

FISCAL 2020 ANNUAL REPORT



Dear Fellow Shareholders,

I write to you as the world rallies to fight a pandemic that has created an unprecedented health and economic crisis. Our Mission: TO HELP OUR CUSTOMERS CREATE A HEALTHIER AND SAFER WORLD has always been a noble calling—a purpose with greater weight and meaning as COVID-19 challenges our resourcefulness, ingenuity and determination.

Before I comment on where we stand as a company today and what we think about our future, I would like to thank the 13,000 people of STERIS—both those working in the field and those behind the scenes in our factories, labs, offices, and increasingly working from their own homes. They are supporting the caregivers of the world with essential healthcare products and services. I am impressed with the way our team has come together during this crisis in support of our Customers and each other.

At STERIS, we have a clear, long-term approach to running the business. Our Customers come first and are followed closely by our people. If we do our jobs well for those two groups, we believe we will deliver above average returns to our shareholders. This philosophy has successfully guided us through several significant challenges over the last decade, and we are confident it will see us through this pandemic as well.

We have chosen a strategy to focus on what we believe to be growth areas in healthcare: procedures, devices, vaccines and biologics. While we are not immune to the downturn in procedures, which we believe to be temporary in nature, we have developed a nice balance of exposure to these areas, along with a good mix of recurring and capital equipment revenue. Finally, we have employed lean techniques and have been insourcing and onshoring to better protect our product and service supply chains for more than a decade - improving quality, delivery reliability, and cost. As a result, we have had a long-term positive run, and so far, our business has fared comparatively well amidst this significant disruption in the global economy.

For fiscal 2020, it is safe to say that we would be celebrating the phenomenal, record-breaking year we just completed, were it not for the pandemic. A few highlights: we broke the \$3 billion revenue mark for the first time; STERIS joined the S&P 500; we had another year of very strong growth rates in revenue and profitability as revenue grew 9% and reported earnings per diluted share reached \$4.76 (constant currency organic revenue grew 10% for fiscal 2020 and adjusted earnings per diluted share reached a new high, growing 15% to \$5.64); and we ended the year with a strong balance sheet. We continued to work toward sustainability. In particular, we made nice progress on our industry-leading efforts to drive sustainable ethylene oxide sterilization. In addition, we are pleased to be able to provide some relief to healthcare providers through our Emergency Use Authorization to decontaminate N95 masks during the COVID-19 crisis. It really was a tremendous year for the people of STERIS, and this stellar performance puts us in an enviable position to face the opportunities and challenges before us.

From a segment perspective, Applied Sterilization Technologies (AST) led the way, growing constant currency organic revenue 15% for the year as a result of increased demand from our core medical device Customers. We expect demand will be strong in the long run, and we plan to continue investing in overall capacity and modality neutrality in this business in fiscal 2021 and beyond.

Healthcare Specialty Services also had another outstanding year, growing constant currency organic revenue 12%. We look forward to ongoing success across our spectrum of offerings, particularly in the United States.

Life Sciences had a better year than anticipated, growing revenue 11% on a constant currency organic basis. Capital equipment sales in this business followed our typical lumpy cadence in fiscal 2020, but our full-year growth exceeded our expectations at 10%. In consumables, we saw high-single- to low-double-digit growth all year, with an upward spike in the fourth quarter at least partially due to COVID-19.

And finally, Healthcare Products had a strong year, with 7% constant currency organic revenue growth driven by new products, particularly in our infection prevention business. Segment growth also benefited from several tuck-in acquisitions this year. Offsetting the acquisitions, Healthcare Products experienced the most immediate revenue loss from the pandemic, particularly in endoscopy products. The positive impact of acquisitions, less the COVID-19 related reductions, combined to essentially offset each other in this segment for the fourth quarter.

Our balance sheet remains on solid ground, as our debt-to-EBITDA leverage ratio continues to fall. Our priorities for capital remain consistent: we plan to continue paying dividends and to invest in our existing businesses to drive anticipated future growth. Our M&A activities have become more selective and deferred due to lack of visibility for near-term expectations, but we continue to evaluate opportunities. And lastly, share buybacks have been discontinued for now, even those to offset dilution for executive compensation.

We have taken some actions to protect our near-term cash flow from a slowdown. These include implementing a hiring freeze for most positions, deferring executive wage increases, canceling significant travel events, restricting routine travel, and deferring capital expenditures except for strategic growth capital.

For the most part, our plants and service operations have been running to support our Customers. We have had limited shutdowns of some facilities and some are operating below normal capacity. We continue to invest in R&D at our normal levels and intend to continue strategic growth capital spending... particularly in AST and in outsourced reprocessing. We firmly believe that these investments will pay off in the intermediate and longer term.

As you know, STERIS is an essential business supporting healthcare. We are very fortunate to be in the business we are in, and to be in a strong financial position. We continue working to enhance the value of your Company as we balance the short-term impacts of the pandemic with our longer-term opportunities.

In closing, we believe the future for STERIS is bright. We appreciate the continued support of our long-term shareholders, the counsel of our Board of Directors, and the dedication of STERIS people working each and every day to make STERIS a great company and help our Customers create a safer and healthier world.

Until next year,

Walt Rosebrough

President and Chief Executive Officer

June 2020

United States Securities and Exchange Commission

Washington, D. C. 20549

		FORM	10-K		
(Maı	k One)				
X	ANNUAL REPORT PURSUANT TO SEC	TION 13 OR 15(d)	OF THE SECURITIES	S EXCHANGE ACT OF 1934	
	For the fiscal year ended March 31, 2020				
			OR		
	TRANSITION REPORT PURSUANT TO	SECTION 13 OR	15(d) OF THE SECUR	ITIES EXCHANGE ACT OF 193	34
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	(Exact		specified in its charter)		
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	Ireland (State or other jurisdiction of			-1455064 Employer	
	incorporation or organization)			fication No.)	
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	(Address of principal executive offices)		(Z	tip code)	
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DOCUMENTS INCORPORATED BY REFERENCE

The number of Ordinary Shares outstanding as of May 22, 2020: 84,918,305

Portions of the Proxy Statement for the 2020 Annual Meeting – Part III

STERIS PLC AND SUBSIDIARIES

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

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PART I

Throughout this Annual Report, references to STERIS plc, "STERIS," "us," or "our," mean STERIS Ireland and its subsidiaries for periods from and after the Redomiciliation and STERIS UK and its subsidiaries for periods prior to the Redomiciliation (as such terms are hereinafter defined), unless otherwise noted. References in this Annual Report to a particular "year," "fiscal," "fiscal year," or "year-end" mean our fiscal year, which ends on March 31. For example, fiscal year 2020 ended on March 31, 2020.

ITEM 1. BUSINESS

INTRODUCTION

STERIS plc is a leading provider of infection prevention and other procedural products and services. Our MISSION IS TO HELP OUR CUSTOMERS CREATE A HEALTHIER AND SAFER WORLD by providing innovative healthcare and life science product and service solutions around the globe. We offer our Customers a unique mix of innovative capital equipment products, such as sterilizers and washers, surgical tables, lights and equipment management systems and connectivity solutions such as operating room integration; consumable products including detergents and gastrointestinal endoscopy accessories and other products and services, including equipment installation and maintenance, microbial reduction of medical devices, instrument and scope repair solutions, laboratory services and outsourced instrument reprocessing.

On March 28, 2019, STERIS plc, a public limited company organized under the laws of England and Wales ("STERIS UK"), completed a redomiciliation from the United Kingdom to Ireland (the "Redomiciliation"). The Redomiciliation was achieved through the insertion of a new Irish public limited holding company ("STERIS Ireland") on top of STERIS UK pursuant to a court-approved scheme of arrangement under English law (the "Scheme"). Following the Scheme effectiveness, STERIS UK was re-registered as a private limited company with the name STERIS Limited, and STERIS Emerald IE Limited, a company established in Ireland and a wholly-owned direct subsidiary of STERIS Ireland, was interposed as the direct parent company of STERIS UK.

STERIS plc's registered office is located in Dublin, Ireland. STERIS plc has approximately 13,000 employees worldwide. Through our field sales and service and a network of dealers and distributors, we serve Customers in more than 100 countries around the world.

We operate and report in four reportable business segments: Healthcare Products, Healthcare Specialty Services, Life Sciences, and Applied Sterilization Technologies. We disclose a measure of segment income that is consistent with the way management operates and views the business. The accounting policies for reportable segments are the same as those for the consolidated Company. In fiscal 2019, we ceased the allocation of certain corporate costs to our segments to align with internal management measures. The fiscal 2018 period operating income measures have been recast for comparability.

The bulk of our revenues are derived from healthcare provider, pharmaceutical and medical device Customers. Much of the growth in these industries is driven by the aging of the population throughout the world, as an increasing number of individuals are entering their prime healthcare consumption years, and are dependent upon advancement in healthcare delivery, acceptance of new technologies, government policies, and general economic conditions. The pharmaceutical industry has been impacted by increased regulatory scrutiny of cleaning and validation processes, mandating that manufacturers improve their processes. Within healthcare, there is concern regarding the level of hospital acquired infections around the world; increased demand for medical procedures, including preventive screenings such as endoscopies and colonoscopies; and a desire by our Customers to operate more efficiently, all of which are driving increased demand for many of our products and services. The COVID-19 pandemic is resulting in the deferral of certain elective medical procedures, which is negatively impacting the demand for some of our products and services.

INFORMATION RELATED TO BUSINESS SEGMENTS

Our chief operating decision maker is our President and Chief Executive Officer ("CEO"). The CEO is responsible for performance assessment and resource allocation. The CEO regularly receives discrete financial information about each reportable segment and uses this information to assess performance and allocate resources. The accounting policies of the reportable segments are the same as those described in Note 1 to the Consolidated Financial Statements titled, "Nature of Operations and Summary of Significant Accounting Policies," of this Annual Report.

HEALTHCARE PRODUCTS SEGMENT

Description of Business. Our Healthcare Products segment provides a broad portfolio of infection prevention, procedural and GI solutions including: consumable products, equipment maintenance and installation services, and capital equipment to acute care hospitals, ambulatory surgery centers and GI clinics. These solutions aid our Customers in improving the safety, quality, productivity, and utility consumption of their surgical, sterile processing, gastrointestinal, and emergency environments.

Products Offered. Our solutions include cleaning chemistries and sterility assurance products, accessories for GI procedures, washers, sterilizers and other pieces of capital equipment essential to the operations of a sterile processing department ("SPD") and equipment used directly in the operating room, including surgical tables, lights, equipment management services, and connectivity solutions.

Services Offered. Our Healthcare Products segment service associates install, maintain, upgrade, repair, and troubleshoot capital equipment throughout the world. We offer various preventive maintenance programs and repair services to support the effective operation of capital equipment over its lifetime.

Customer Concentration. Our Healthcare Products segment sells consumables, services and capital equipment, to Customers in many countries throughout the world. For the year ended March 31, 2020, no Customer represented more than 10% of the Healthcare Product segment's total revenues.

Competition. We compete with a number of large companies that have significant product portfolios and global reach, as well as a number of small companies with very limited product offerings and operations in one or a limited number of countries. On a product basis, competitors include 3M, Belimed, Cantel Medical, Ecolab, Getinge, Hill-Rom, Fortive, Stryker and Skytron.

HEALTHCARE SPECIALTY SERVICES SEGMENT

Description of Business. Our Healthcare Specialty Services segment provides a range of solutions and managed services including: hospital sterilization services and instrument and scope repairs to acute care hospitals and other healthcare settings that aid our Customers in improving the safety, quality and productivity of their operations.

Services Offered. Our Healthcare Specialty Services segment provides comprehensive instrument and endoscope repair and maintenance solutions (on-site or at one of our dedicated facilities), custom process improvement consulting and outsourced instrument sterile processing (on-site at the hospital and in off-site reprocessing centers).

Customer Concentration. Our Healthcare Specialty Services segment offers an array of services to Customers in many countries throughout the world. For the year ended March 31, 2020, no Customer represented more than 10% of the Healthcare Specialty Services segment's total revenues.

Competition. We compete with a number of large companies that have significant product portfolios and global reach, as well as a number of small companies with very limited service offerings and operations in one or a limited number of countries. On a service line basis, competitors include BBraun, Berendsen plc, CleanLease (Clean Lease Fortex), Karl Storz, Mobile, Northfield, Olympus, Owens & Minor, Pentax, Rentex Awé and Rentex Floren and Sterilog Limited.

LIFE SCIENCES SEGMENT

Description of Business. Our Life Sciences segment designs, manufactures and sells consumable products, equipment maintenance, specialty services and capital equipment primarily to pharmaceutical manufacturers around the world.

Products Offered. These solutions include formulated cleaning chemistries, barrier products, sterility assurance products, steam and vaporized hydrogen peroxide sterilizers and washer disinfectors.

Services Offered. Our Life Sciences segment service associates install, maintain, upgrade, repair, and troubleshoot equipment throughout the world. We offer various preventive maintenance programs and repair services to support the effective operation of capital equipment over its lifetime.

Customer Concentration. Our Life Sciences segment sells consumables, services and capital equipment, to Customers in many countries throughout the world. For the year ended March 31, 2020, no Customer represented more than 10% of the Life Sciences segment's total revenues.

Competition. Our Life Sciences segment operates in highly regulated environments where the most intense competition results from technological innovations, product performance, convenience and ease of use, and overall cost-effectiveness. We compete for pharmaceutical Customers with a number of large companies that have significant product portfolios and global reach, as well as a number of small companies with very limited product offerings and operations in one or a limited number of countries. Competitors include Belimed, Ecolab, Fedegari, Getinge, MECO, Stilmas, and Techniplast.

APPLIED STERILIZATION TECHNOLOGIES SEGMENT

Description of Business. Our Applied Sterilization Technologies ("AST") segment provides contract sterilization and testing services for medical device and pharmaceutical manufacturers. As a technology neutral service provider, we offer unbiased technology assessments dependent on the individual requirements of each product. Our Customers are primarily medical device and pharmaceutical manufacturers.

Services Offered. We offer a wide range of sterilization modalities as well as an array of testing services that complements the manufacturing of sterile products. Our locations are in major population centers and core distribution corridors throughout the

Americas, Europe and Asia. Our technical services group supports Customers in all phases of product development, materials testing, and process validation.

Customer Concentration. Our Applied Sterilization Technologies segment's services are offered to Customers throughout the world. For the year ended March 31, 2020, no Customer represented more than 10% of the segment's revenues.

Competition. Applied Sterilization Technologies operates in a highly regulated industry and competes with Sterigenics International, Inc., other smaller contract sterilization companies and manufacturers that sterilize products in-house.

INFORMATION WITH RESPECT TO OUR BUSINESS IN GENERAL

Sources and Availability of Raw Materials. We purchase raw materials, sub-assemblies, components, and other supplies needed in our operations from numerous suppliers in the United States and internationally. The principal raw materials and supplies used in our operations include stainless and carbon steel, organic and inorganic chemicals, fuel, and plastic components. These raw materials and supplies are generally available from several suppliers and in sufficient quantities that we do not currently expect any significant sourcing problems in fiscal 2021. We have long-term supply contracts for certain materials for which there are few suppliers, or those that are single-sourced in certain regions of the world, such as EO and cobalt-60, which are necessary to our AST operations. In addition, we have developed a plan to expand our irradiation processing capacity with accelerator-based technologies, which may reduce the potential supply risk.

Intellectual Property. We protect our technology and products by, among other means, obtaining United States and foreign patents. There can be no assurance, however, that any patent will provide adequate protection for the technology, system, product, service, or process it covers. In addition, the process of obtaining and protecting patents can be long and expensive. We also rely upon trade secrets, technical know-how, and continuing technological innovation to develop and maintain our competitive position.

As of March 31, 2020, we held approximately 410 United States patents and approximately 1,640 in other jurisdictions and had approximately 145 United States patent applications and 360 patent applications pending in other jurisdictions. Patents for individual products extend for varying periods according to the date of filing or grant and legal term of patents in various countries where a patent is obtained. The actual protection a patent provides varies from country to country and depends in part upon the type of patent, the scope of its coverage, and the availability of legal remedies in each country.

Our products are sold around the world under various brand names and trademarks. We consider our brand names and trademarks to be valuable in the marketing of our products. As of March 31, 2020, we had a total of approximately 1,430 trademark registrations worldwide.

Quality Assurance. We manufacture, assemble, and package products in several countries. Each of our production facilities are dedicated to particular processes and products. Our success depends upon Customer confidence in the quality of our production process and the integrity of the data that supports our product safety and effectiveness. We have implemented quality assurance procedures to support the quality and integrity of scientific information and production processes.

Government Regulation. Our business is subject to various degrees of governmental regulation in the countries in which we operate. In the United States, the United States Food and Drug Administration ("FDA"), the United States Environmental Protection Agency ("EPA"), the United States Nuclear Regulatory Commission ("NRC"), and other governmental authorities regulate the development, manufacture, sale, and distribution of our products and services. Our international operations also are subject to a significant amount of government regulation, including country-specific rules and regulations and U.S. regulations applicable to our international operations. Government regulations include detailed inspection of, and controls over, research and development, clinical investigations, product approvals and manufacturing, marketing and promotion, sampling, distribution, record-keeping, storage, and disposal practices.

Compliance with applicable regulations is a significant expense for us. Past, current or future regulations, their interpretation, or their application could have a material adverse impact on our operations. Also, additional governmental regulation may be passed that could prevent, delay, revoke, or result in the rejection of regulatory clearance of our products. We cannot predict the effect on our operations resulting from current or future governmental regulation or the interpretation or application of these regulations.

If we fail to comply with any applicable regulatory requirements, sanctions could be imposed on us. For more information about the risks we face regarding regulatory requirements, see Part I, Item 1A of this Annual Report titled, "Risk Factors". We are subject to extensive regulatory requirements and must receive and maintain regulatory clearance or approval for many products and operations. Failure to receive or maintain, or delays in receiving, clearance or approvals may hurt our revenues, profitability, financial condition, or value.

In the past, we have received warning letters, paid civil penalties, conducted product recalls and field corrections, and been subject to other regulatory sanctions. We believe that we are currently compliant in all material respects with applicable

regulatory requirements. However, there can be no assurance that future or current regulatory, governmental, or private action will not have a material adverse affect on us or on our performance, results, or financial condition.

Environmental Matters. We are subject to various laws and governmental regulations concerning environmental matters and employee safety and health in Ireland, the United States and other countries. We have made, and continue to make, significant investments to comply with these laws and regulations. We cannot predict the future capital expenditures or operating costs required to comply with environmental laws and regulations. We believe that we are currently compliant with applicable environmental, health, and safety requirements in all material respects. However, there can be no assurance that future or current regulatory, governmental, or private action will not have a material adverse affect on our performance, results, or financial condition. Please refer to Note 10 of our consolidated financial statements titled, "Commitments and Contingencies" for further information.

In the future, if a loss contingency related to environmental matters, employee safety, health or conditional asset retirement obligations is significantly greater than the current estimated amount, we would record a liability for the obligation and it may result in a material impact on net income for the annual or interim period during which the liability is recorded. The investigation and remediation of environmental obligations generally occur over an extended period of time, and therefore we do not know if these events would have a material adverse affect on our financial condition, liquidity, or cash flow, nor can there be any assurance that such liabilities would not have a material adverse affect on our performance, results, or financial condition.

Competition. The markets in which we operate are highly competitive and generally highly regulated. Competition is intense in all of our business segments and includes many large and small competitors. Brand, design, quality, safety, ease of use, serviceability, price, product features, warranty, delivery, service, and technical support are important competitive factors to us. We expect to face continued competition in the future as new infection prevention, sterile processing, contamination control, gastrointestinal and surgical support products and services enter the market. We believe many organizations are working with a variety of technologies and sterilizing agents. Also, a number of companies have developed disposable medical instruments and other devices designed to address the risk of contamination.

We believe that our long-term competitive position depends on our success in discovering, developing, and marketing innovative, cost-effective products and services. We devote significant resources to research and development efforts and we believe STERIS is positioned as a global competitor in the search for technological innovations. In addition to research and development, we invest in quality control, Customer programs, distribution systems, technical services, and other information services.

There can be no assurance that we will develop significant new products or services, or that the new products or services we provide or develop in the future will be more commercially successful than those provided or developed by our competitors. In addition, some of our existing or potential competitors may have greater resources than us. Therefore, a competitor may succeed in developing and commercializing products more rapidly than we do. Competition, as it relates to our business segments and product categories, is discussed in more detail in the section above titled, "Information Related to Business Segments."

Employees. As of March 31, 2020, we had approximately 13,000 employees throughout the world including certain locations subject to collective bargaining agreements and works council representation. We believe we generally have good relations with our employees.

Methods of Distribution. Sales and service activities are supported by a staff of regionally based clinical specialists, system planners, corporate account managers, and in-house Customer service and field support departments. We also contract with distributors and dealers in select markets.

Customer training is important to our business. We provide a variety of courses at Customer locations, at our training and education centers, and over the internet. Our training programs help Customers understand the science, technology, and operation of our products and services. Many of our operator training programs are approved by professional certifying organizations and offer continuing education credits to eligible course participants.

Seasonality. Our financial results have been, from time to time, subject to seasonal patterns. We cannot assure you that these patterns will continue.

Backlog. We define backlog as the amount of unfilled capital equipment purchase orders at a point in time. At March 31, 2020, we had a backlog of \$242.5 million. Of this amount, \$170.1 million and \$72.4 million related to our Healthcare Products and Life Sciences segments, respectively. At March 31, 2019, we had backlog orders of \$215.2 million. Of this amount, \$154.5 million and \$60.7 million related to our Healthcare Products and Life Sciences segments, respectively.

Availability of Securities and Exchange Commission Filings. We make available free of charge on or through our website our Annual Reports on Form 10-K, Quarterly Reports on Form 10-Q, and Current Reports on Form 8-K, and amendments to

these reports, as soon as reasonably practicable after we electronically file such material with, or furnish such material to, the Securities and Exchange Commission ("SEC"). You may access these documents, as well as other SEC filings related to the Company, on the Investor Relations page of our website at http://www.steris-ir.com. You may also obtain copies of these documents by accessing the SEC's website at http://www.sec.gov. The content on or accessible through any website referred to in this Annual Report on Form 10-K is not incorporated by reference into this Form 10-K unless expressly noted.

We also make available free of charge on our website our Corporate Governance Guidelines, our Director Code of Ethics, and our Code of Business Conduct, as well as the Charters of the Audit Committee, the Compensation and Organization Development Committee, the Nominating and Governance Committee, and the Compliance Committee of the Company's Board of Directors.

INFORMATION ABOUT OUR EXECUTIVE OFFICERS

The following table presents certain information regarding our executive officers at March 31, 2020. All executive officers serve at the pleasure of the Board of Directors.

Name	Age	Position
Karen L. Burton	52	Vice President, Controller and Chief Accounting Officer
Daniel A. Carestio	47	Senior Vice President and Chief Operating Officer
Cary L. Majors	45	Senior Vice President, North America Commercial Operations
Walter M Rosebrough, Jr.	66	President and Chief Executive Officer
Renato G. Tamaro	51	Vice President and Corporate Treasurer
Michael J. Tokich	51	Senior Vice President and Chief Financial Officer
J. Adam Zangerle	53	Senior Vice President, General Counsel, and Secretary

The following discussion provides a summary of each executive officer's recent business experience through March 31, 2020:

Karen L. Burton serves as Vice President, Controller and Chief Accounting Officer. She assumed this role in January 2017. She served as Vice President, Corporate Controller from May 2008 to January 2017.

Daniel A. Carestio serves as Senior Vice President and Chief Operating Officer. He assumed this role in August 2018. From February 2018 to August 2018 he served as Senior Vice President, Sterilization and Disinfection. From August 2015 to February 2018, he served as Senior Vice President, STERIS Applied Sterilization Technologies and Life Sciences. From 2011 to August 2015, he served as Vice President, Sales and Marketing for Isomedix Services and General Manager of Life Sciences.

Cary L. Majors serves as Senior Vice President, North America Commercial Operations. He assumed this role in August 2019. From April 2014 to August 2019 he served as Vice President, North America Commercial Operations.

Walter M Rosebrough, Jr. serves as President and Chief Executive Officer. He assumed this role when he joined STERIS in October 2007. Mr. Rosebrough is also a Director of STERIS plc and Varex Imaging Corporation.

Renato G. Tamaro serves as Vice President and Corporate Treasurer. He assumed this role in August 2017. From March 2006 to July 2017, he served as Assistant Treasurer.

Michael J. Tokich serves as Senior Vice President and Chief Financial Officer. He assumed this role in August 2017. From February 2014 to July 2017, he served as the Senior Vice President, Chief Financial Officer and Treasurer.

J. Adam Zangerle serves as Senior Vice President, General Counsel, and Secretary. He assumed this role in July 2018. From July 2013 to July 2018 he served as Vice President, General Counsel, and Secretary.

ITEM 1A. RISK FACTORS

This section describes certain risk factors that could affect our business, financial condition and results of operations. You should consider these risk factors when evaluating the forward-looking statements contained in this Annual Report on Form 10-K, because our actual results and financial condition might differ materially from those projected in the forward-looking statements should these risks occur. We face other risks besides those highlighted below. These other risks include additional uncertainties not presently known to us or that we currently believe are immaterial, but may ultimately have a significant impact. In addition, the impact of the COVID-19 pandemic may also exacerbate any of these risks, which could have a material effect on us. Should any of these risks, described below or otherwise, actually occur, our business, financial condition, performance, prospects, value, or results of operations could be negatively affected.

MARKET RISKS

Risk or uncertainty	Discussion
Doing business internationally	
We conduct manufacturing, sales and distribution operations on a worldwide basis and are subject to a variety of risks associated with doing business internationally. Implementation and achievement of international growth objectives also may be impeded by political, social, and economic uncertainties or unrest in countries in which we conduct operations or market or distribute our products.	We maintain significant international operations, including operations in the U.S., Canada, Mexico, Europe, Asia Pacific and Latin America. As a result, we are subject to a number of risks and complications associated with international manufacturing, sales, services, and other operations. These include: risks associated with currency exchange rate fluctuations; difficulties in enforcing agreements and collecting receivables through some foreign legal systems; enhanced credit risks in certain European countries as well as emerging market regions; Customers with longer payment cycles than Customers in the United States; significant variations in tax rates among the countries in which we do business, and tax withholding obligations in respect of our earnings; tax laws that restrict our ability to use tax credits, offset gains, or repatriate funds; tariffs, exchange controls or other trade restrictions including transfer pricing restrictions when products produced in one country are sold to an affiliated entity in another country; the impact of the COVID-19 pandemic on our supply chain and the industries in which we operate; general economic and political conditions in countries where we operate or where end users of our products are situated, including the potential implications of the COVID-19 pandemic, the U.K. "Brexit", for the U.K. and/or regional or global economies, or the withdrawal from the EU of other member countries; difficulties associated with managing a large organization spread throughout various countries; difficulties in enforcing intellectual property rights or weaker intellectual property right protections in some countries and difficulties associated with compliance with a variety of laws and regulations governing international trade, including the U.S. Foreign Corrupt Practices Act and the U.K. Bribery Act and laws and regulations dealing with trade with persons in sanctioned countries.
Compliance with multiple, and potentially conflicting, international laws and regulations, import and export limitations, anti-corruption laws, and exchange controls may be difficult, burdensome or expensive.	We are subject to compliance with various laws and regulations, including the U.S. Foreign Corrupt Practices Act, the U.K. Bribery Act, and similar anti-bribery laws, which generally prohibit companies and their intermediaries from making improper payments to officials for the purpose of obtaining or retaining business. We are also subject to limitations on trade with persons in sanctioned countries. While our employees and agents are required to comply with these laws, we cannot assure you that our internal policies and procedures will always protect us from violations of these laws, despite our commitment to legal compliance and corporate ethics.

Economic conditions and financial market access

The COVID-19 pandemic has disrupted our operations and could have a material adverse effect on our business and financial condition.

The COVID-19 pandemic, along with the response to the pandemic by governmental and other actors, has disrupted our operations. We have experienced temporary mandatory and voluntary facility closures in certain jurisdictions in which we operate. Furthermore, we have experienced less demand for some of our products and services as a result of official prohibitions or voluntary deferrals of certain medical procedures, and other factors, which we believe has been exacerbated by the impact of stay-at-home orders. Additionally, the COVID-19 outbreak has, caused temporary disruptions in our supply chain.

Long-term facility closures or other restrictions could materially adversely affect our ability to adequately staff, supply or otherwise maintain our operations. Such restrictions also may have a substantial impact on our Customers and our sales cycles. The COVID-19 pandemic may put pressure on overall spending for our products and services, and may cause our Customers to modify spending priorities or delay or abandon purchasing decisions. Moreover, because a large number of our employees have transitioned to working from home, we may be subject to increased vulnerability to cyber and other information technology risks. We have modified, and may further modify, our business practices in response to the risks and negative impacts associated with the COVID-19 pandemic. However, there can be no assurance that these measures will be temporary or successful.

The impact of the COVID-19 pandemic continues to evolve and its ultimate duration, severity and disruption to our business, Customers and supply chain, and the related financial impact to us, cannot be accurately forecasted at this time. Should such disruption continue for an extended period, the adverse effect on our business, results of operations and financial condition could be more severe. Additionally, continued weak economic conditions generally could result in extended weak demand for our products and services. Furthermore, future public health crises are possible and could involve some or all of the risks discussed above.

Changes in economic climate may adversely affect us.

Adverse economic cycles or conditions, and Customer, regulatory or government response to those cycles or conditions, have affected and could further affect our results of operations. The onset of these cycles or conditions may not be foreseeable and there can be no assurance when they will begin to improve after they occur. There also can be no assurance as to the strength or length of any recovery from a business downturn or recession. Credit and liquidity problems may make it difficult for some businesses to access credit markets and obtain financing and may cause some businesses to curtail spending to conserve cash in anticipation of persistent business slowdowns and liquidity needs. If our Customers have difficulty financing their purchases due to tight credit markets or related factors or because of other operational or utilization problems they may be experiencing or otherwise decide to curtail their purchases, our business could be adversely affected. Our exposure to bad debt losses could also increase if Customers are unable to pay for products previously ordered and delivered.

Many of our Customers are governmental entities or other entities that rely on government healthcare systems or government funding. If government funding for healthcare becomes limited or restricted in countries in which we operate, including as a result of the impacts of the COVID-19 pandemic, our Customers may be unable to pay their obligations on a timely basis or to make payment in full and it may become necessary to increase reserves. In addition, there can be no assurance that there will not be an increase in collection difficulties. Prospectively, additional adverse effects resulting from these conditions may include decreased healthcare utilization, further pricing pressure on our products and services, and/or weaker overall demand for our products and services, particularly capital products.

Our acquisition activity and ability to grow organically may be adversely affected if we are unable to continue to access the financial markets.

Our recent acquisitions have been financed largely through cash on hand and borrowings under our bank credit facilities. Future acquisitions or other capital requirements will necessitate additional cash. To the extent our existing sources of cash are insufficient to fund these or other future activities, we may need to raise additional funds through new or expanded borrowing arrangements or equity. There can be no assurance that we will be able to obtain additional funds beyond those available under existing bank credit facilities on terms favorable to us, or at all, or that such facilities can be replaced when they terminate.

LEGAL, REGULATORY AND TAX RISKS

Risk or uncertainty Discussion Healthcare laws and

Changes in healthcare laws or government and other third-party payor reimbursement levels to healthcare providers, or failure to meet healthcare reimbursement or other requirements, might negatively impact our business.

reimbursement

We sell many of our products and services to hospitals and other healthcare providers and pharmaceutical manufacturers. Many of these Customers are subject to or supported by government programs or receive reimbursement for services from third-party payors, such as government programs, including Medicare and Medicaid in the U.S., private insurance plans, and managed care programs. Reimbursement systems vary significantly by country. Government-managed healthcare systems control reimbursement for healthcare services in many countries. Public budgetary constraints may significantly impact the ability of hospitals, pharmaceutical manufacturers, and other Customers supported by such systems to purchase our products. Government or other third-party payors may deny or change coverage, reduce their current levels of reimbursement for healthcare services, or otherwise implement measures to regulate pricing or contain costs. In addition, our costs may increase more rapidly than reimbursement levels or permissible pricing increases or we may not satisfy the standards or requirements for reimbursement.

Among other provisions, the U.S. Patient Protection and Affordable Care Act, as amended by the Health Care and Education Affordability Reconciliation Act, imposed an excise tax on medical devices manufactured or offered for sale in the United States. Late in 2019, U.S. Congress enacted legislation that repealed the excise tax, which had been suspended during calendar years 2016 through 2019. In addition, we have been required to commit significant resources to "Sunshine Act" compliance. Various additional health care reform proposals have emerged at the federal and state level, and we are unable to predict which, if any, of those proposals will be enacted.

Risk or uncertainty

Discussion

Product and service related regulations and claims

We are subject to extensive regulatory requirements and must receive and maintain regulatory clearance or approval for many products and operations. Failure to receive or maintain, or delays in receiving, clearance or approvals may hurt our revenues, profitability, financial condition, or value.

Our operations are subject to extensive regulation in the countries where we do business. In the United States, our products and services are regulated by the FDA and other regulatory authorities. In many foreign countries, sales of our products and services are subject to extensive regulations that may or may not be comparable to those of the FDA. In Europe, our products are regulated primarily by country and community regulations of those countries within the European Economic Area and must conform to the requirements of those authorities.

Government regulation applies to nearly all aspects of testing, manufacturing, safety, labeling, storing, recordkeeping, reporting, promoting, distributing, and importing or exporting of medical devices, products, and services. In general, unless an exemption applies, a sterilization, decontamination or medical device or product or service must receive regulatory approval or clearance before it can be marketed or sold. Modifications to existing products or the marketing of new uses for existing products also may require regulatory approvals, approval supplements or clearances. If we are unable to obtain any required approvals, approval supplements or clearances for any modification to a previously cleared or approved device, we may be required to cease manufacturing and sale, or recall or restrict the use of such modified device, pay fines, or take other action until such time as appropriate clearance or approval is obtained.

Regulatory agencies may refuse to grant approval or clearance, or review and disagree with our interpretation of approvals or clearances, or with our decision that regulatory approval is not required or has been maintained. Regulatory submissions may require the provision of additional data and may be time consuming and costly, and their outcome is uncertain. Regulatory agencies may also change policies, adopt additional regulations, or revise existing regulations, each of which could prevent or delay approval or clearance of devices, or could impact our ability to market a previously cleared, approved, or unregulated device. Our failure to comply with the regulatory requirements of the FDA or other applicable regulatory requirements in the United States or elsewhere might subject us to administratively or judicially imposed sanctions. These sanctions include, among others, warning letters, fines, civil penalties, criminal penalties, injunctions, debarment, product seizure or detention, product recalls and total or partial suspension of production, sale and/or promotion.

The COVID-19 pandemic may disrupt the operations of regulatory bodies with responsibility for oversight of healthcare and health and medical products. Such disruptions could result in the focus and prioritization of regulatory resources on emergent matters, which could divert regulatory resources away from more routine regulatory matters that are not COVID-19 related but that have the potential to impact our business. For example, there could be delays in FDA review of applications for marketing authorization, including those which may be necessary for or in connection with proposed changes to our products or the changes to the processes by which they are manufactured. It is unknown how long these disruptions could continue, were they to occur. Any elongation or de-prioritization or delay in regulatory review resulting from such disruptions could materially affect our ongoing device design, development, and commercialization plans.

Our products are subject to recalls and restrictions, even after receiving United States or foreign regulatory clearance or approval. Ongoing medical device reporting regulations require that we report to appropriate governmental authorities in the United States and/or other countries when our products cause or contribute to a death or serious injury or malfunction in a way that would be reasonably likely to contribute to a death or serious injury if the malfunction were to recur. Governmental authorities can require product recalls or impose restrictions for product design, manufacturing, labeling, clearance, or other issues. For the same reasons, we may voluntarily elect to recall or restrict the use of a product. Any recall or restriction could divert managerial and financial resources and might harm our reputation among our Customers and other healthcare professionals who use or recommend our products and services.

We may be adversely affected by product liability claims or other legal actions or regulatory or compliance matters. We face an inherent business risk of exposure to product liability claims and other legal and regulatory actions. A significant increase in the number, severity, amount, or scope of these claims and actions may, as described above with respect to recalls and restrictions, result in substantial costs and harm our reputation or otherwise adversely affect product sales and our business. Product liability claims and other legal and regulatory actions may also distract management from other business responsibilities.

We are also subject to a variety of other types of claims, proceedings, investigations, and litigation initiated by government agencies or third parties and other potential risks and liabilities. These include compliance matters, product regulation or safety, taxes, employee benefit plans, employment discrimination, health and safety, environmental, antitrust, customs, import/export, government contract compliance, financial controls or reporting, intellectual property, allegations of misrepresentation, false claims or false statements, commercial claims, claims regarding promotion of our products and services, or other similar or different matters. Any such claims, proceedings, investigations or litigation, regardless of the merits, might result in substantial costs, restrictions on product use or sales, or otherwise injure our business.

Administratively or judicially imposed or agreed sanctions might include warning letters, fines, civil penalties, criminal penalties, loss of tax benefits, injunctions, product seizure, recalls, suspensions or restrictions, re-labeling, detention, and/or debarment. We also might be required to take actions such as payment of substantial amounts, or revision of financial statements, or to take, or be subject to, the following types of actions with respect to our products, services, or business: redesign, re-label, restrict, or recall products; cease manufacturing and selling products; seizure of product inventory; comply with a court injunction restricting or prohibiting further marketing and sale of products or services; comply with a consent decree, which could result in further regulatory constraints; dedication of significant internal and external resources and costs to respond to and comply with legal and regulatory issues and constraints; respond to claims, litigation, and other proceedings brought by Customers, users, governmental agencies, and others; disruption of product improvements and product launches; discontinuation of certain product lines or services; or other restrictions or limitations on product sales, use or operation, or other activities or business practices.

Some product replacements or substitutions may not be possible or may be prohibitively costly or time consuming. The impact of any legal, regulatory, or compliance claims, proceeding, investigation, or litigation, is difficult to predict.

We maintain product liability and other insurance with coverages believed to be adequate. However, product liability or other claims may exceed insurance coverage limits, fines, penalties and regulatory sanctions may not be covered by insurance, or insurance may not continue to be available or available on commercially reasonable terms. Additionally, our insurers might deny claim coverage for valid or other reasons or may become insolvent.

Our business and financial condition could be adversely affected by difficulties in acquiring or maintaining a proprietary intellectual ownership position.

To maintain our competitive position for our products, we need to obtain patent or other proprietary rights for new and improved products and to maintain and enforce our existing patents and other proprietary rights. We typically apply for patents in the United States and in strategic other countries. We may also acquire patents through acquisitions. We may encounter difficulties in obtaining or protecting patents.

We rely on a combination of patents, trademarks, trade secrets, know-how, and confidentiality agreements to protect the proprietary aspects of our technology. These measures afford only limited protection, and competitors may gain access to our intellectual property and proprietary information. Litigation may be necessary to enforce or defend our intellectual property rights, to protect our trade secrets, and to determine the validity and scope of our proprietary rights. Litigation may also be brought against us claiming that we have violated the intellectual property rights of others. Litigation may be costly and may divert management's attention from other matters. Additionally, in some foreign countries with weaker intellectual property rights, it may be difficult to maintain and enforce patents and other proprietary rights or defend against claims of infringement.

Tax and trade risks

Current economic and political conditions make tax rules in any jurisdiction subject to significant change.

The U.S. Tax Cuts and Jobs Act ("TCJA") was signed into law on December 22, 2017. Guidance continues to be issued clarifying the application of this new legislation. We cannot predict the overall impact that the additional guidance may have on our business. It is reasonable to expect that global taxing authorities will be reviewing current legislation for potential modifications in reaction to the implementation of the TCJA. In addition, further changes in the tax laws of other jurisdictions could arise, including as a result of the base erosion and profit shifting (BEPS) project undertaken by the Organization for Economic Cooperation and Development (OECD). The OECD, which represents a coalition of member countries, has issued recommendations that, in some cases, would make substantial changes to numerous long-standing tax positions and principles. These contemplated changes, to the extent adopted by OECD members and/or other countries, could increase tax uncertainty and may adversely impact our provision for income taxes.

Our tax rate is uncertain and may vary from expectations, which could have a material impact on our results of operations and earnings per share. There can be no assurance that we will be able to maintain any particular worldwide effective corporate tax rate. We cannot give any assurance as to what our effective tax rate will be in the future because of, among other things, uncertainty regarding the tax policies of the jurisdictions in which we and our affiliates operate. Our actual effective tax rate may vary from our expectations, and such variance may be material. Additionally, tax laws or their implementation and applicable tax authority practices in any particular jurisdiction could change in the future, possibly on a retroactive basis, and any such change could have a material adverse impact on us and our affiliates.

Changes in tax treaties and trade agreements could negatively impact our costs, results of operations and earnings per share. Legislative and regulatory action may be taken in the U.S. which, if ultimately adopted, could override or otherwise adversely impact tax treaties upon which we rely or broaden the circumstances under which STERIS plc would be considered a U.S. resident, each of which could materially and adversely affect our tax obligations. We cannot predict the outcome of any specific legislative or regulatory proposals. However, if proposals were adopted that had the effect of disregarding our organization in Ireland or limiting our ability as an Irish company to take advantage of tax treaties with the U.S., we could be subject to increased taxation and/or potentially significant expense.

Existing free trade laws and regulations provide certain beneficial duties and tariffs for qualifying imports and exports, subject to compliance with the applicable classification and other requirements. Changes in laws and regulations or policies governing the terms of foreign trade, and in particular, increased trade restrictions, including as a result of the COVID-19 pandemic, tariffs or taxes on imports from countries where we manufacture products could have a material adverse impact on our business and financial results.

Proposed legislation relating to	Various U.S. federal and state legislative proposals that would deny governmental
the denial of U.S. federal or state	contracts to redomiciled companies may adversely affect us if adopted into law. We are
governmental contracts to U.S.	unable to predict the likelihood that any such proposed legislation might become law, the
companies that redomicile abroad	nature of regulations that may be promulgated under any future legislative enactments, or
could adversely affect our	the effect such enactments or increased regulatory scrutiny could have on our business.
business.	
The U.S. Internal Revenue	Although we are organized under the laws of Ireland and are a tax resident in Ireland for
Service (the "IRS") may not agree	Irish tax purposes, the IRS may assert that we should be treated as a U.S. corporation
that we are a foreign corporation	(and, therefore, a U.S. tax resident) for U.S. federal tax purposes pursuant to
for U.S. federal tax purposes.	Section 7874 of the Internal Revenue Code of 1986, as amended (the "Code" and such
	Section, "Section 7874"). For U.S. federal tax purposes, a company generally is

Irish tax purposes, the IRS may assert that we should be treated as a U.S. corporation (and, therefore, a U.S. tax resident) for U.S. federal tax purposes pursuant to Section 7874 of the Internal Revenue Code of 1986, as amended (the "Code" and such Section, "Section 7874"). For U.S. federal tax purposes, a company generally is considered to be a tax resident in the jurisdiction of its organization. Because we are organized under the laws of Ireland, we would generally be classified as a non-U.S. corporation (and, therefore, a non-U.S. tax resident) under these rules. Section 7874, however, provides an exception to this general rule under which a non-U.S. organized entity may be treated as a U.S. corporation for U.S. federal tax purposes.

If we were to be treated as a U.S. corporation for U.S. federal tax purposes, we could be subject to substantial additional U.S. tax liability. Additionally, if we were treated as a U.S. corporation for U.S. federal tax purposes, non-U.S. holders of our ordinary shares would be subject to U.S. withholding tax on the gross amount of any dividends we paid to such shareholders. For Irish tax purposes, we are expected, regardless of any application of Section 7874, to be treated as an Ireland tax resident. Consequently, if we are treated as a U.S. corporation for U.S. federal tax purposes under Section 7874, we could be liable for both U.S. and Ireland taxes, which could have a material adverse effect on our financial condition and results of operations.

BUSINESS AND OPERATIONAL RISKS

Risk or uncertainty Discussion				
Competition				
Our businesses are highly competitive, and if we fail to compete successfully, our revenues and results of operations may be hurt.	We operate in a highly competitive global environment. Our businesses compete with other broad-line manufacturers, as well as many smaller businesses specializing in particular products or services, primarily on the basis of brand, design, quality, safety, ease of use, serviceability, price, product features, warranty, delivery, service, and technical support. We face increased competition from new infection prevention, sterile processing, contamination control, surgical support, cleaning consumables, gastrointestinal endoscopy accessories, contract sterilization, and other products and services entering the market. Competitors and potential competitors also are attempting to develop alternate technologies and sterilizing agents, as well as disposable medical instruments and other devices designed to address the risk of contamination.			
Consolidations among our healthcare and pharmaceutical Customers may result in a loss of Customers or more significant pricing pressures.	A number of our Customers have consolidated. These consolidations are due in part to healthcare cost reduction measures initiated by competitive pressures as well as legislators, regulators and third-party payors. This may result in greater pricing pressures on us and in some cases loss of Customers. Additional consolidations could result in a loss of Customers or more significant pricing pressures.			

Decreased availability or increased costs of raw materials or energy supplies or other supplies might increase our production costs or limit our production capabilities or curtail our operations.

We purchase raw materials, fabricated and other components, and energy supplies from a variety of suppliers. Key materials include stainless steel, organic and inorganic chemicals, fuel, cobalt-60, EO, and plastic components. The availability and prices of raw materials and energy supplies are subject to volatility and are influenced by worldwide economic conditions, speculative action, world supply and demand balances, inventory levels, availability of substitute materials, currency exchange rates, anticipated or perceived shortages, and other factors. Also, certain of our key materials and components have a limited number of suppliers. Some are single-sourced in certain regions of the world, such as cobalt-60 and EO, which are necessary to our AST operations. Changes in regulatory requirements regarding the use of, the unavailability or short supply of these products might disrupt or cause shutdowns of portions of our AST operations or have other adverse consequences. We have developed a plan to expand our irradiation processing capacity with accelerator-based technologies which may reduce the potential supply risk. Shortages in supply, increased regulatory or security requirements, or increases in the price of raw materials, components and energy supplies may adversely affect us.

Our operations, and those of our suppliers, are subject to a variety of business continuity hazards and risks, any of which could interrupt production or operations or otherwise adversely affect our performance, results, or value. Business continuity hazards and other risks include: explosions, fires, earthquakes, public health crises, inclement weather, and other disasters; utility or other mechanical failures; unscheduled downtime; labor difficulties; inability to obtain or maintain any required licenses or permits; disruption of communications; data security, preservation and redundancy disruptions; inability to hire or retain key management or employees; disruption of supply or distribution; and regulation of the safety, security or other aspects of our operations.

The occurrence of these types of events has disrupted and may in the future disrupt or shut down operations, or otherwise adversely impact the production or profitability of a particular facility, or our operations as a whole. Certain casualties also might cause personal injury and loss of life, or severe damage to or destruction of property and equipment, and for casualties occurring at our facilities, result in liability claims against us. Although we maintain property and casualty insurance and liability and similar insurance of the types and in the amounts that we believe are customary for our industries, our insurance coverages have limits and we are not fully insured against all potential hazards and risks incident to our business.

Our operations are subject to regulations and permitting, which may be changed or amended by the relevant authorities, and which may limit or eliminate our current operations or increase the complexity, burden, or expense of compliance and regulated materials or processes that we use in our operations may become the focus of litigation.

Our Applied Sterilization Technologies ("AST") segment is a technology-neutral contract sterilization service that offers our Customers a wide range of sterilization modalities through a worldwide network of over 50 contract sterilization and laboratory facilities. One of the modalities offered by our AST operations is Ethylene Oxide ("EO") sterilization. In the United States, several regulators, including the U.S. Environmental Protection Agency ("EPA"), U.S. Food and Drug Administration ("FDA"), and agencies at the state and local level, play a role in regulating the use of EO sterilization. In 2016, the EPA changed the cancer risk basis for EO and determined that EO is carcinogenic to humans. Recent announcements of the temporary or permanent closure of EO sterilization facilities operated by others have been associated with state and/or local regulatory or other legal action related to EO emissions at those facilities. Our AST operations have taken and will continue to take measures to comply with all applicable emissions regulations and to reduce emissions. However, no assurance can be given that current or future legislative or regulatory action, or current or future litigation to which we are or may become a party, will not significantly increase the costs of conducting our EO contract sterilization operations or curtail or eliminate the use of EO in our contract sterilization operations. A significant reduction in our EO contract sterilization activities may have a material adverse effect on our financial condition and results of operations. Further, we could be liable for damages and fines as a result of legislative or regulatory action or litigation, and any liability could exceed our insurance and indemnification coverage, if any, and have a material adverse effect on our financial condition. Additionally, for many medical devices, EO sterilization may be the only current method of sterilization that effectively sterilizes and does not damage the device during the sterilization process. In the event of regulatory, legislative, or legal action that curtails or eliminates EO sterilization, there could be a shortage of medical devices and consequently a decline in surgical procedures. A decline in surgical procedures could result in a decline in demand for the products and services provided by our Healthcare Products and Healthcare Specialty Services businesses, which may have a material adverse effect on our financial condition and results of operations.

We engage in acquisitions and affiliations, divestitures, and other business arrangements. Our growth may be adversely affected if we are unable to successfully identify, price, and integrate strategic business candidates or otherwise optimize our business portfolio.

Our success depends, in part, on strategic acquisitions and joint ventures, which are intended to complement or expand our businesses, divestiture of non-strategic businesses, and other actions intended to optimize our portfolio of businesses. This strategy depends upon our ability to identify, appropriately price, and complete these types of business development transactions or arrangements and to obtain any necessary financing. In the last several fiscal years we have made a number of acquisitions. We also completed several divestitures of non-strategic businesses or product lines during the last several years

Our success with respect to these recent and future acquisitions will depend on our ability to integrate the businesses acquired, retain key personnel, realize identified cost synergies and otherwise execute our strategies. Our success will also depend on our ability to develop satisfactory working arrangements with our strategic partners in joint ventures or other affiliations, or to divest or realign businesses. Competition for strategic business candidates may result in increases in costs and price for acquisition candidates and market valuation issues may reduce the value available for divestiture of non-strategic businesses. These types of transactions are also subject to a number of other risks and uncertainties, including: delays in realizing or failure to realize anticipated benefits of the transactions; diversion of management's time and attention from other business concerns; difficulties in retaining key employees, Customers, or suppliers of the acquired or divested businesses; difficulties in maintaining uniform standards, controls, procedures and policies, or other integration or divestiture difficulties; adverse effects on existing business relationships with suppliers or Customers; other events contributing to difficulties in generating future cash flows; risks associated with the assumption of contingent or other liabilities of acquisition targets or retention of liabilities for divested businesses and difficulties in obtaining financing.

If our continuing efforts to create a lean business and in-source production to reduce costs are not successful, our profitability may be hurt or our business otherwise might be adversely affected. We have undertaken various activities to create a lean business, including in-sourcing. We continue to look for opportunities to in-source production that is currently provided by third parties. These activities may not produce the full efficiencies and cost reduction benefits that we expect or efficiencies and benefits might be delayed. Implementation costs also might exceed expectations.

The COVID-19 pandemic or similar public health crises could have a material adverse impact on ability to staff our operations.

As supplier to Healthcare and Life Sciences Customers, we fall within a "critical infrastructure" sector, and are also considered an essential business and therefore exempt under various stay at home/shelter in place orders. Accordingly, our employees continue to work because of the importance of our operations to the health and well-being of citizens in the countries in which we operate. We have implemented telework policies wherever possible for appropriate categories of employees. However, our employees that are unable to telework continue to work at our facilities and those of our Customers, and we have implemented appropriate safety measures, such as social distancing and increased cleaning protocols. While we believe that we have taken appropriate measures to ensure the health and well-being of our employees, there can be no assurances that our measures will be sufficient to protect our employees in our workplace or that they may not otherwise be exposed to COVID-19 outside of our workplace. If a number of our essential employees become ill, incapacitated or are otherwise unable or unwilling to continue working during the current or any future health crises, our operations may be adversely impacted.

Our business and results of operations may be adversely affected if we are unable to recruit and retain qualified management and other personnel or other compliance matters adversely impact our personnel. Our continued success depends, in large part, on our ability to hire and retain highly qualified people and if we are unable to do so, our business and operations may be impaired or disrupted. Competition for highly qualified people is intense and there is no assurance that we will be successful in attracting or retaining replacements to fill vacant positions, successors to fill retirements or employees moving to new positions, or other highly qualified personnel. In addition, legal, regulatory or compliance matters create significant distraction or diversion of significant or unanticipated resources or attention that could have a material adverse effect on the responsibilities and retention of qualified employees.

We could experience a failure of a key information technology system, process or site or a breach of information security, including a cybersecurity breach or failure of one or more key information technology systems, networks, processes, associated sites or service providers. We rely extensively on information technology (IT) systems to conduct business. In addition, we rely on networks and services, including internet sites, data hosting and processing facilities and tools and other hardware, software and technical applications and platforms, some of which are managed, hosted, provided and/or used by third-parties or their vendors, to assist in conducting our business. Numerous and evolving cybersecurity threats pose potential risks to the security of our IT systems, networks and services, as well as the confidentiality, availability and integrity of our data. While we have made investments seeking to address these threats, including monitoring of networks and systems, hiring of experts, employee training and security policies for employees and third-party providers, the techniques used in these attacks change frequently and may be difficult to detect for periods of time and we may face difficulties in anticipating and implementing adequate preventative measures. If our IT systems are damaged or cease to function properly, the networks or service providers we rely upon fail to function properly, or we or one of our third-party providers suffer a loss or disclosure of our business or stakeholder information due to any number of causes ranging from catastrophic events or power outages to improper data handling or security breaches and our business continuity plans do not effectively address these failures on a timely basis, we may be exposed to reputational, competitive and business harm as well as litigation and regulatory action. In addition, the COVID-19 pandemic may increase the risk of such vulnerability and attacks, including unauthorized access or attacks exploiting the fact that a large number of employees are working remotely during government shutdowns and closures. Enforcement of the General Data Protection Regulation ("GDPR") was effective as of May 2018. The GDPR is focused on the protection of personal data not merely the privacy of personal data. The GDPR creates a range of new compliance obligations and will significantly increase financial penalties for noncompliance (including possible fines of up to 4% of global annual revenues for the preceding financial year or €20 million (whichever is higher) for the most serious infringements).

ITEM 1B. UNRESOLVED STAFF COMMENTS

None.

ITEM 2. PROPERTIES

The following discussion sets forth materially important properties of the Company and its subsidiaries as of March 31, 2020. The Company believes that its facilities are adequate for operations and are maintained in good condition. The Company is confident that, if needed, it will be able to acquire additional facilities at commercially reasonable rates. In the following discussion "International" is defined as all countries other than Ireland and the United States.

The Company's principal executive office is located in Dublin, Ireland and its primary administrative offices are located in Mentor, OH (U.S.).

The Company owns 44 and leases 11 contact sterilization locations, utilized in the Applied Sterilization Technologies Segment that are located in major population centers and core distribution corridors throughout the Americas, Europe and Asia.

The Company operates over 75 locations representing sales, administrative and operational locations in the U.S. and 19 other countries, the majority of which are leased and support one or multiple business segments. Operational locations are primarily comprised of service centers and distribution warehouses. Our locations are geographically spread to be in close proximity to our Customers to ensure timely delivery of products and services.

The Company owns and leases several material manufacturing locations that support one or more of our Healthcare Products, Healthcare Specialty Services and Life Sciences segments, which are disclosed in the following table:

Location	U.S./INTL*	Leased/Owned
Montgomery, AL	U.S.	Owned/Leased
St. Louis, MO	U.S.	Owned/Leased
Mentor, OH	U.S.	Owned/Leased
Sharon Hill, PA	U.S.	Owned
Franklin Park, IL	U.S.	Leased
Point Richmond, CA	U.S.	Leased
Quebec City, Canada	INTL	Owned
Tuusula, Finland	INTL	Owned/Leased
Bordeaux, France	INTL	Owned
Leicester, England	INTL	Owned/Leased
Shanghai, China	INTL	Leased
Guadalupe, Mexico	INTL	Leased
Bishop Stortford, England	INTL	Leased

^{*} International includes all countries other than Ireland and the U.S.

ITEM 3. LEGAL PROCEEDINGS

Information regarding our legal proceedings is included in Item 7 of Part II, Management's Discussion and Analysis ("MD&A"), and Note 10 of our consolidated financial statements titled, "Commitments and Contingencies," and is incorporated herein by reference thereto.

ITEM 4. MINE SAFETY DISCLOSURES

None.

PART II

ITEM 5. MARKET FOR REGISTRANT'S ORDINARY EQUITY, RELATED SHAREHOLDER MATTERS AND ISSUER PURCHASES OF EQUITY SECURITIES

Market Information. Our ordinary shares are traded on the New York Stock Exchange under the symbol "STE."

Holders. As of March 31, 2020, there were approximately 1,040 holders of record of our ordinary shares.

Dividend Policy. The Company's Board of Directors decides the timing and amount of any dividends we may pay. The Board expects to be able to continue to pay cash dividends for the foreseeable future.

Purchases of Equity Securities by the Issuer and Affiliated Purchasers. On August 9, 2016, STERIS UK announced that its Board of Directors had authorized the purchase of up to \$300.0 million (net of taxes, fees and commissions) of our ordinary shares. As a result of the Redomiciliation, this authorization terminated.

On May 7, 2019, our Board of Directors authorized the continuation of the foregoing share repurchase program by STERIS plc, resulting in a share repurchase authorization of \$79.0 million (net of taxes, fees and commissions). On July 30, 2019, our Board of Directors approved an increase in the May 7, 2019 authorization of an additional amount of \$300.0 million (net of taxes, fees and commissions).

As of March 31, 2020, there was approximately \$339.0 million (net of taxes, fees and commissions) of remaining availability under the authorizations.

Under the authorizations, shares may be repurchased from time to time through open market transactions, including 10b5-1 plans. Any repurchase program may be activated, suspended or discontinued at any time.

We purchased 273,259 of our ordinary shares during fiscal 2020 for the aggregate amount of \$40.0 million, pursuant to the 2019 authorizations.

The following table presents information with respect to purchases STERIS made of its ordinary shares during the fourth quarter of fiscal year 2020:

	(a) Total Number of Shares Purchased	(b) Average Price Paid Per Share	(c) Total Number of Shares Purchased as Part of Publicly Announced Plans	of SI Be Pi Pla	(d) Maximum Dollar Value of Shares that May Yet Be Purchased Under the Plans at Period End (dollars in thousands)		
January 1-31	_	\$ -	_	\$	348,979		
February 1-29	22,500	166.2	0 22,500		345,239		
March 1-31	45,700	136.5	5 45,700		338,979		
Total	68,200 (1)	\$ 146.0	8 (1) 68,200	\$	338,979		

⁽¹⁾ Does not include 8 shares purchased during the quarter at an average price of \$151.44 per share by the STERIS Corporation 401(k) Plan on behalf of an executive officer of the Company who may be deemed to be an affiliated purchaser.

ITEM 6. SELECTED FINANCIAL DATA

	Years Ended March 31,									
in thousands, except per share data)		2020 (1)		2019 (1) (2)		2018 (2)		2017 (2)		2016 ⁽²⁾
Statements of Income Data:										
Revenues	\$	3,030,895	\$	2,782,170	\$	2,619,996	\$	2,612,756	\$	2,238,764
Gross profit		1,319,923		1,175,427		1,092,746		1,026,213		895,348
Restructuring expenses		673		30,987		103		215		(820)
Income from continuing operations		536,973		411,465		399,883		226,206		237,576
Income taxes		90,876		64,394		63,360		74,015		60,299
Net income attributable to shareholders Basic income per ordinary share:		407,605		304,051		290,915		109,965		110,763
Net income	\$	4.81	\$	3.59	\$	3.42	\$	1.29	\$	1.57
Shares used in computing net income per ordinary share – basic Diluted income per ordinary share:		84,778		84,577		85,028		85,473		70,698
Net income	\$	4.76	\$	3.56	\$	3.39	\$	1.28	\$	1.56
Shares used in computing net income per ordinary share – diluted		85,641		85,468		85,713		86,094		71,184
Dividends per ordinary share	\$	1.45	\$	1.33	\$	1.21	\$	1.09	\$	0.98
Balance Sheets Data:										
Working capital	\$	705,144	\$	588,539	\$	591,195	\$	636,219	\$	571,919
Total assets		5,425,582		5,073,071		5,200,334		4,924,555		5,346,416
Long-term indebtedness		1,150,521		1,183,227		1,316,001		1,478,361		1,567,796
Total liabilities		2,018,858		1,887,273		1,983,034		2,114,422		2,307,524
Total shareholders' equity	\$	3,393,876	\$	3,177,810	\$	3,205,960	\$	2,798,602	\$	3,023,034

⁽¹⁾ See "Management's Discussion and Analysis of Financial Condition and Results of Operations."
(2) As a result of our adoption of ASU 2017-07, prior year amounts on our Consolidated Statements of Income have been reclassified to retroactively apply the components of the net periodic benefit cost of our defined benefit pension plans and our other post-retirements benefit plan.

ITEM 7. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

INTRODUCTION

In Management's Discussion and Analysis ("MD&A"), we explain the general financial condition and the results of operations for STERIS and its subsidiaries including:

- what factors affect our business:
- what our earnings and costs were;
- why those earnings and costs were different from the year before;
- where our earnings came from;
- how this affects our overall financial condition;
- what our expenditures for capital projects were; and
- where cash will come from to fund future debt principal repayments, growth outside of core operations, repurchase ordinary shares, pay cash dividends and fund future working capital needs.

The MD&A also analyzes and explains the annual changes in the specific line items in the Consolidated Statements of Income. As you read the MD&A, it may be helpful to refer to information in Item 1, "Business," Item 6, "Selected Financial Data," and our consolidated financial statements, which present the results of our operations for fiscal 2020, 2019 and 2018 as well as Part I, Item 1A, "Risk Factors" and Note 10 of our consolidated financial statements titled, "Commitments and Contingencies" for a discussion of some of the matters that can adversely affect our business and results of operations. This information, discussion, and disclosure may be important to you in making decisions about your investments in STERIS.

Information on our financial condition and results of our operations for our 2018 fiscal year period can be found in Item 7 titled, "Management's Discussion and Analysis of Financial Condition and Results of Operations", of our Annual Report on Form 10-K for the fiscal year ended March 31, 2019, filed with the SEC on May 30, 2019.

FINANCIAL MEASURES

In the following sections of the MD&A, we may, at times, refer to financial measures that are not required to be presented in the consolidated financial statements under U.S. GAAP. We sometimes use the following financial measures in the context of this report: backlog; debt-to-total capital; and days sales outstanding. We define these financial measures as follows:

- <u>Backlog</u> We define backlog as the amount of unfilled capital equipment purchase orders at a point in time. We use this figure as a measure to assist in the projection of short-term financial results and inventory requirements.
- <u>Debt-to-total capital</u> We define debt-to-total capital as total debt divided by the sum of total debt and shareholders' equity. We use this figure as a financial liquidity measure to gauge our ability to borrow and fund growth.
- <u>Days sales outstanding ("DSO")</u> We define DSO as the average collection period for accounts receivable. It is calculated as net accounts receivable divided by the trailing four quarters' revenues, multiplied by 365 days. We use this figure to help gauge the quality of accounts receivable and expected time to collect.

We, at times, may also refer to financial measures which are considered to be "non-GAAP financial measures" under SEC rules. We have presented these financial measures because we believe that meaningful analysis of our financial performance is enhanced by an understanding of certain additional factors underlying that performance. These financial measures should not be considered an alternative to measures required by accounting principles generally accepted in the United States. Our calculations of these measures may differ from calculations of similar measures used by other companies and you should be careful when comparing these financial measures to those of other companies. Additional information regarding these financial measures, including reconciliations of each non- GAAP financial measure, is available in the subsection of MD&A titled, "Non-GAAP Financial Measures."

REVENUES- DEFINED

As required by Regulation S-X, we separately present revenues generated as either product revenues or service revenues on our Consolidated Statements of Income for each period presented. When we discuss revenues, we may, at times, refer to revenues summarized differently than the Regulation S-X requirements. The terminology, definitions, and applications of terms that we use to describe revenues may be different from terms used by other companies. We use the following terms to describe revenues:

- Revenues Our revenues are presented net of sales returns and allowances.
- <u>Product Revenues</u> We define product revenues as revenues generated from sales of consumable and capital equipment products.
- <u>Service Revenues</u> We define service revenues as revenues generated from parts and labor associated with the maintenance, repair, and installation of our capital equipment. Service revenues also include hospital sterilization services, instrument and scope repairs, and linen management as well as revenues generated from contract sterilization and laboratory services offered through our Applied Sterilization Technologies segment.
- <u>Capital Equipment Revenues</u> We define capital equipment revenues as revenues generated from sales of capital equipment, which includes steam sterilizers, low temperature liquid chemical sterilant processing systems, including SYSTEM 1 and 1E, washing systems, VHP[®] technology, water stills, and pure steam generators; surgical lights and tables; and integrated OR.
- <u>Consumable Revenues</u> We define consumable revenues as revenues generated from sales of the consumable family
 of products, which includes SYSTEM 1 and 1E consumables, V-PRO consumables, gastrointestinal endoscopy
 accessories, sterility assurance products, skin care products, cleaning consumables, barrier product solutions and
 surgical instruments.
- <u>Recurring Revenues</u> We define recurring revenues as revenues generated from sales of consumable products and service revenues.

GENERAL OVERVIEW AND EXECUTIVE SUMMARY

STERIS plc is a leading provider of infection prevention and other procedural products and services. Our MISSION IS TO HELP OUR CUSTOMERS CREATE A HEALTHIER AND SAFER WORLD by providing innovative healthcare and life science product and service solutions around the globe. We offer our Customers a unique mix of innovative consumable products, such as detergents, gastrointestinal ("GI") endoscopy accessories, barrier product solutions, and other products and services, including: equipment installation and maintenance, microbial reduction of medical devices, instrument and scope repair solutions, laboratory testing services, on-site and off-site reprocessing, and capital equipment products, such as sterilizers and surgical tables, and connectivity solutions such as operating room ("OR") integration.

On March 28, 2019, STERIS plc, a public limited company organized under the laws of England and Wales ("STERIS UK"), completed a redomiciliation from the United Kingdom to Ireland (the "Redomiciliation"). The Redomiciliation was achieved through the insertion of a new Irish public limited holding company ("STERIS Ireland") on top of STERIS UK pursuant to a court-approved scheme of arrangement under English law (the "Scheme"). Following the Scheme effectiveness, STERIS UK was re-registered as a private limited company with the name STERIS Limited, and STERIS Emerald IE Limited, a company established in Ireland and a wholly-owned direct subsidiary of STERIS Ireland, was interposed as the direct parent company of STERIS UK.

We operate and report in four reportable business segments: Healthcare Products, Healthcare Specialty Services, Life Sciences, and Applied Sterilization Technologies. We describe our business segments in Note 11 to our consolidated financial statements, titled "Business Segment Information."

The bulk of our revenues are derived from the healthcare, medical device and pharmaceutical industries. Much of the growth in these industries is driven by the aging of the population throughout the world, as an increasing number of individuals are entering their prime healthcare consumption years, and is dependent upon advancement in healthcare delivery, acceptance of new technologies, government policies, and general economic conditions. The pharmaceutical industry has been impacted by increased regulatory scrutiny of cleaning and validation processes, mandating that manufacturers improve their processes. Within healthcare, there is increased concern regarding the level of hospital acquired infections around the world; increased demand for medical procedures, including preventive screenings such as endoscopies and colonoscopies; and a desire by our Customers to operate more efficiently, all which are driving increased demand for many of our products and services. The COVID-19 pandemic is resulting in the deferral of certain elective medical procedures, which is negatively impacting the demand for some of our products and services.

We completed several tuck in acquisitions and asset purchases in fiscal 2020 and 2019 that expanded our product and service offerings to our Customers.

During fiscal 2020, we sold the operations of our Healthcare Specialty Services business that were located in China with annual revenues of approximately \$5.0 million.

We continue to invest in manufacturing in-sourcing projects and lean process improvements for the purpose of improving quality, cost and delivery of our products to our Customers.

U.S. Tax Reform. On December 22, 2017, the U.S. government enacted comprehensive tax legislation commonly referred to as the Tax Cuts and Jobs Act (the "TCJA"). The TCJA made broad and complex changes to the U.S. tax code including, but not limited to, (1) reduction of the U.S. federal corporate income tax rate; (2) elimination of the corporate alternative minimum tax ("AMT"); (3) the creation of the base erosion anti-abuse tax ("BEAT"), a new minimum tax; (4) a general elimination of U.S. federal income taxes on dividends from non-U.S. subsidiaries; (5) a new provision designed to tax global intangible low-taxed income ("GILTI"), which allows for the possibility of using foreign tax credits ("FTCs") and a deduction of up to 50 percent to offset the income tax liability (subject to some limitations); (6) a new limitation on deductible interest expense; (7) the repeal of the domestic production activity deduction; (8) limitations on the deductibility of certain executive compensation; (9) limitations on the use of FTCs to reduce the U.S. income tax liability; and (10) limitations on net operating losses ("NOLs") generated after December 31, 2017, to 80.0 percent of taxable income.

Fiscal 2019 Restructuring Plan. During the third quarter of fiscal year 2019, we adopted and announced a targeted restructuring plan (the "Fiscal 2019 Restructuring Plan"), which included the closure of two manufacturing facilities, one in Brazil and one in England, as well as other actions including, the rationalization of certain products. Fewer than 200 positions were eliminated. The Company has relocated the production of certain impacted products to other existing manufacturing operations during fiscal 2020. These restructuring actions were designed to enhance profitability and improve efficiency. For additional information on restructuring see the subsection titled "Restructuring Expenses", located in the Results of Operations section of this MD&A, or Note 2 of our Consolidated Financial Statements, titled "Restructuring".

Highlights. Revenues increased \$248.7 million, or 8.9%, to \$3,030.9 million for the year ended March 31, 2020, as compared to \$2,782.2 million for the year ended March 31, 2019. This increase reflects organic growth in all business segments, which was partially offset by unfavorable fluctuations in currencies.

Fiscal 2020 operating income increased 30.5% to \$537.0 million over fiscal 2019 operating income of \$411.5 million. The increase is primarily attributable to lower restructuring expenses, increased revenue volumes and higher gross margin attainment in fiscal 2020 over fiscal 2019.

Net cash flows from operations were \$590.6 million and free cash flow was \$380.2 million in fiscal 2020 compared to net cash flows from operations of \$539.5 million and free cash flow of \$355.4 million in fiscal 2019 (see subsection of MD&A titled, "Non-GAAP Financial Measures" for additional information and related reconciliation of non-GAAP financial measures to the most comparable GAAP measures). The increase in free cash flow is primarily due to the improvement in cash from operations.

Our debt-to-total capital ratio was 25.3% at March 31, 2020. During the year, we increased our quarterly dividend for the fourteenth consecutive year to \$0.37 per share per quarter.

Outlook. In fiscal 2021 and beyond, we expect to continue to manage our costs, grow our business with internal product and service development, invest in greater capacity, and augment these value creating methods with potential acquisitions of additional products and services.

However, the COVID-19 pandemic began to impact our business late in fiscal 2020. The coronavirus pandemic and related public health recommendations and mandated precautions to mitigate the spread of COVID-19, including deferral of medical procedures and treatments and shelter-in-place orders or similar measures, is negatively affecting, and is expected to continue to affect some of our operations which would impact our financial position and cash flows in fiscal 2021. We have experienced and expect to continue to experience unpredictable fluctuations in demand for certain of our products and services, including some products and services that are experiencing increased demand.

We cannot predict the ultimate impact that the COVID-19 pandemic and related actions will have on our Customers' operations, financial position and cash flows and therefore, on the demand for our products and services.

Further, the broader economic impact of the COVID-19 pandemic response could cause interest rate variability and generate unanticipated fluctuations in currency rates that impact our revenues and costs outside of the United States, creating variability in our results.

As a result, we are unable to estimate the ultimate impact of the COVID-19 pandemic to our consolidated results of operations, financial position and cash flows for fiscal 2021 and beyond.

NON-GAAP FINANCIAL MEASURES

We, at times, refer to financial measures which are considered to be "non-GAAP financial measures" under SEC rules. We, at times, also refer to our results of operations excluding certain transactions or amounts that are non-recurring or are not indicative of future results, in order to provide meaningful comparisons between the periods presented.

These non-GAAP financial measures are not intended to be, and should not be, considered separately from or as an alternative to the most directly comparable GAAP financial measures.

These non-GAAP financial measures are presented with the intent of providing greater transparency to supplemental financial information used by management and the Board of Directors in their financial analysis and operational decision-making. These amounts are disclosed so that the reader has the same financial data that management uses with the belief that it will assist investors and other readers in making comparisons to our historical operating results and analyzing the underlying performance of our operations for the periods presented.

We believe that the presentation of these non-GAAP financial measures, when considered along with our GAAP financial measures and the reconciliation to the corresponding GAAP financial measures, provide the reader with a more complete understanding of the factors and trends affecting our business than could be obtained absent this disclosure. It is important for the reader to note that the non-GAAP financial measure used may be calculated differently from, and therefore may not be comparable to, a similarly titled measure used by other companies.

We define free cash flow as net cash provided by operating activities as presented in the Consolidated Statements of Cash Flows less purchases of property, plant, equipment, and intangibles plus proceeds from the sale of property, plant, equipment, and intangibles, which are also presented within investing activities in the Consolidated Statements of Cash Flows. We use this as a measure to gauge our ability to pay cash dividends, fund growth outside of core operations, fund future debt principal repayments, and repurchase shares. The following table summarizes the calculation of our free cash flow for the years ended March 31, 2020 and 2019:

Vegre Ended March 31

	rears Ended March				
(dollars in thousands)		2020		2019	
Net cash flows provided by operating activities	\$	590,559	\$	539,505	
Purchases of property, plant, equipment and intangibles, net		(214,516)		(189,715)	
Proceeds from the sale of property, plant, equipment and intangibles		4,156		5,567	
Free cash flow	\$	380,199	\$	355,357	

RESULTS OF OPERATIONS

The COVID-19 pandemic began to impact our business late in fiscal 2020 and therefore did not have a material impact on our fiscal 2020 results of operations.

In the following subsections, we discuss our earnings and the factors affecting them. We begin with a general overview of our operating results and then separately discuss earnings for our operating segments.

FISCAL 2020 AS COMPARED TO FISCAL 2019

Revenues. The following table compares our revenues, in total and by type and geography, for the year ended March 31, 2020 to the year ended March 31, 2019:

	Years Ende	d Ma	rch 31,			Percent	
(dollars in thousands)	2020		2019		Change	Change	
Total revenues \$	3,030,895	\$	2,782,170	\$	248,725	8.9%	
Revenues by type:							
Service revenues	1,628,107		1,486,145		141,962	9.6%	
Consumable revenues	672,329		605,631		66,698	11.0%	
Capital equipment revenues	730,459		690,394		40,065	5.8%	
Revenues by geography:							
Ireland revenues	63,821		56,784		7,037	12.4%	
United States revenues	2,211,722		1,976,814		234,908	11.9%	
Other foreign revenues	755,352		748,572		6,780	0.9%	

Revenues increased \$248.7 million, or 8.9%, to \$3,030.9 million for the year ended March 31, 2020, as compared to \$2,782.2 million for the year ended March 31, 2019. This increase reflects organic growth in all business segments and favorable pricing, which was partially offset by unfavorable fluctuations in currencies.

Service revenues for fiscal 2020 increased \$142.0 million, or 9.6% over fiscal 2019, reflecting growth in all business segments. Consumable revenues for fiscal 2020 increased \$66.7 million, or 11.0%, over fiscal 2019, reflecting growth in the Healthcare Products and Life Sciences segments. Capital equipment revenues for fiscal 2020 increased by \$40.1 million, or 5.8%, over fiscal 2019, reflecting strong shipment volumes in the Healthcare Products and Life Science business segments.

Ireland revenues for fiscal 2020 were \$63.8 million, representing an increase of \$7.0 million, or 12.4%, over fiscal 2019 revenues of \$56.8 million, reflecting strong growth in service revenues.

United States revenues for fiscal 2020 were \$2,211.7 million, representing an increase of \$234.9 million, or 11.9%, over fiscal 2019 revenues of \$1,976.8 million, reflecting double digit growth in service, consumable and capital equipment revenues.

Revenues from other foreign locations for fiscal 2020 were \$755.4 million, representing an increase of 0.9% over the fiscal 2019 revenues of \$748.6 million, reflecting strength in Canada and the Latin America region. The Europe, Middle East and Africa ("EMEA") region slightly declined primarily due to actions taken in conjunction with the 2019 Restructuring Plan.

Gross Profit. The following table compares our gross profit for the year ended March 31, 2020 to the year ended March 31, 2019:

		Years Ende			Percent		
(dollars in thousands)		2020		2019		Change	Change
Gross profit:							
Product	\$	652,586	\$	593,730	\$	58,856	9.9%
Service		667,337		581,697		85,640	14.7%
Total gross profit	\$	1,319,923	\$	1,175,427	\$	144,496	12.3%
Gross profit percentage:							
Product		46.5%)	45.8%	, 0		
Service		41.0%)	39.1%	, 0		
Total gross profit percentage	_	43.5%		42.2%	ó		
	_				_		

Our gross profit is affected by the volume, pricing and mix of sales of our products and services, as well as the costs associated with the products and services that are sold. Our gross profit increased \$144.5 million and gross profit percentage increased 130 basis points to 43.5% for fiscal 2020 as compared to 42.2% for fiscal 2019. The increase in gross margin percentage is primarily due to the favorable impact of pricing (50 basis points), lower current period expenses related to the Fiscal 2019 Restructuring Plan (20 basis points), our recent divestitures (10 basis points) and mix and other adjustments (50 basis points). Productivity enhancements fully offset material, labor and facility cost increases.

Operating Expenses. The following table compares our operating expenses for the year ended March 31, 2020 to the year ended March 31, 2019:

		Years Ende		Percent			
(dollars in thousands)	2020 2019				Change	Change	
Operating expenses:							
Selling, general, and administrative	\$	716,731	\$	669,937	\$ 46,794	7.0%	
Research and development		65,546		63,038	2,508	4.0%	
Restructuring expenses		673		30,987	(30,314)	NM	
Total operating expenses	\$	782,950	\$	763,962	\$ 18,988	2.5%	

NM - Not meaningful

Selling, General, and Administrative Expenses. Significant components of total selling, general, and administrative expenses ("SG&A") are compensation and benefit costs, fees for professional services, travel and entertainment, facilities costs, gains or losses from divestitures, and other general and administrative expenses. SG&A increased 7.0% in fiscal 2020 over fiscal 2019. Volume sensitive costs like commissions and third party purchasing organization fees increased 15% in fiscal 2020 over fiscal 2019, but continue to be approximately 3% of revenues. Higher compensation costs related to our annual employee bonus and additional operating expenses from our newly acquired businesses also contributed to the fiscal 2020 increase.

Research and Development. Research and development expenses increased \$2.5 million during fiscal 2020, as compared to fiscal 2019, due primarily to increased spending within the Healthcare Products segment. Research and development expenses are influenced by the number and timing of in-process projects and labor hours and other costs associated with these projects. Our research and development initiatives continue to emphasize new product development, product improvements, and the development of new technological platform innovations. During fiscal 2020, our investments in research and development continued to be focused on, but were not limited to, enhancing capabilities of sterile processing combination technologies, procedural products and accessories, and devices and support accessories used in gastrointestinal endoscopy procedures.

Restructuring Expenses. During the third quarter of fiscal 2019, we adopted and announced a targeted restructuring plan (the "Fiscal 2019 Restructuring Plan"), which included the closure of two manufacturing facilities, one in Brazil and one in England, as well as other actions including the rationalization of certain products. Fewer than 200 positions were eliminated. The Company has relocated the production of certain impacted products to other existing manufacturing operations during fiscal 2020. These restructuring actions were designed to enhance profitability and improve efficiency.

We have incurred pre-tax expenses totaling \$43.9 million related to these restructuring actions, of which \$31.7 million was recorded as restructuring expenses and \$12.2 million was recorded in cost of revenues, with a total of \$31.2 million, \$2.5 million, \$0.7 million, and \$7.8 million related to the Healthcare Products, Healthcare Specialty Services, Life Sciences, and Applied Sterilization Technologies segments, respectively. Corporate related restructuring charges were \$1.7 million. Additional restructuring expenses related to this plan are not expected to be material to our results of operations.

The following table summarizes our total pre-tax restructuring expenses for fiscal 2020 and 2019:

Fiscal 2019 Restructuring Plan	Ma	ar Ended arch 31, 2020	Year Ended March 31, 2019
(dollars in thousands)			
Severance and other compensation related costs	\$	1,554 \$	5,651
Accelerated depreciation and amortization		_	16,194
(Gain) on disposal of asset		(1,164)	
Asset impairment		_	4,312
Lease termination costs and other		283	4,830
Product rationalization (1)		2,470	9,721
Total restructuring expenses	\$	3,143 \$	40,708

⁽¹⁾ Recorded in cost of revenues on the Consolidated Statements of Income.

Non-Operating Expenses, Net. Non-operating expense (income), net consists of interest expense on debt, offset by interest earned on cash, cash equivalents, short-term investment balances, and other miscellaneous expense. The following table

compares our non-operating expense (income), net for the year ended March 31, 2020 to the year ended March 31, 2019:

(dollars in thousands)		2020	2019	Change
Non-operating expenses, net:				
Interest expense	\$	40,279 \$	45,015	\$ (4,736)
Interest income and miscellaneous expense		(1,987)	(3,020)	1,033
Non-operating expenses, net	\$	38,292 \$	41,995	\$ (3,703)

Interest expense decreased \$4.7 million during fiscal 2020, as compared to fiscal 2019, primarily due to lower outstanding debt levels in the fiscal 2020 period as compared to the same prior year period (refer to our Note 6 to our consolidated financial statements, titled "Debt", for more information). Interest income and miscellaneous expense is not material.

Additional information regarding our outstanding debt is included in Note 6 to our consolidated financial statements titled, "Debt," and in the subsection of this MD&A titled, "Liquidity and Capital Resources."

Income Tax Expense. The following table compares our income tax expense and effective income tax rates for the years ended March 31, 2020 and March 31, 2019:

	Years Ende		Percent		
(dollars in thousands)	 2020	2019	Change	Change	
Income tax expense	\$ 90,876	\$ 64,394	\$ 26,482	41.1%	-
Effective income tax rate	18.2%	17.4%			

The effective income tax rate for fiscal 2020 was 18.2% as compared to 17.4% for fiscal 2019. The fiscal 2020 effective tax rate increased when compared to fiscal 2019 primarily due to an increased percentage of profits earned and taxed in jurisdictions with a higher tax rate.

Business Segment Results of Operations. We operate and report in four reportable business segments: Healthcare Products, Healthcare Specialty Services, Life Sciences, and Applied Sterilization Technologies. Non-allocated operating costs that support the entire Company and items not indicative of operating trends are excluded from segment operating income.

Our Healthcare Products segment offers infection prevention and procedural solutions for healthcare providers worldwide, including consumable products, equipment maintenance and installation services, and capital equipment.

Our Healthcare Specialty Services segment provides a range of specialty services for healthcare providers including hospital sterilization services and instrument and scope repairs.

Our Life Sciences segment offers consumable products, equipment maintenance, specialty services and capital equipment primarily for pharmaceutical manufacturers.

Our Applied Sterilization Technologies ("AST") segment provides contract sterilization and testing services for medical device and pharmaceutical manufacturers.

We disclose a measure of segment income that is consistent with the way management operates and views the business. The accounting policies for reportable segments are the same as those for the consolidated Company.

For more information regarding our segments please refer to Note 11 to our consolidated financial statements titled "Business Segment Information," and Item 1, "Business".

The following table compares business segment and Corporate and other revenues and operating income for the year ended March 31, 2020 to the year ended March 31, 2019:

		Years ende		Percent	
(dollars in thousands)		2020	2019	Change	Change
Revenues:				_	
Healthcare Products	\$	1,423,198	\$ 1,338,428	\$ 84,770	6.3 %
Healthcare Specialty Services		563,611	510,057	53,554	10.5 %
Life Sciences		416,939	378,558	38,381	10.1 %
Applied Sterilization Technologies		627,147	555,127	72,020	13.0 %
Total revenues	\$	3,030,895	\$ 2,782,170	\$ 248,725	8.9 %
Operating income (loss):					
Healthcare Products		356,419	323,684	32,735	10.1 %
Healthcare Specialty Services		64,217	64,222	(5)	— %
Life Sciences		144,088	132,129	11,959	9.1 %
Applied Sterilization Technologies		270,917	221,828	49,089	22.1 %
Corporate		(207,015)	(184,900)	(22,115)	12.0 %
Total operating income before adjustments	\$	628,626	\$ 556,963	\$ 71,663	12.9 %
Less: Adjustments					
Amortization of acquired intangible assets (1)		71,675	86,878		
Acquisition and integration related charges (2)		8,225	8,901		
Redomiciliation and tax restructuring costs (3)		3,699	8,783		
(Gain) on fair value adjustment of acquisition related contingent consideration (1)		_	(842)		
Net loss (gain) on divestiture of businesses (1)		1,770	(1,370)		
Amortization of property "step up" to fair value (1)		2,392	2,440		
Restructuring charges (4)		3,143	40,708		
COVID-19 incremental costs (5)	_	749	 		
Total operating income	\$	536,973	\$ 411,465		

⁽¹⁾ For more information regarding our recent acquisitions and divestitures see Note 18 titled, "Business Acquisitions and Divestitures". Amortization of purchased intangible assets fiscal 2019 total includes an impairment charge of \$16.2 million, see Note 3 titled, "Goodwill and Intangible Assets", for more information.

Healthcare Products revenues increased 6.3% in fiscal 2020, as compared to fiscal 2019, reflecting growth in consumable, service and capital equipment revenues of 9.5%, 6.1% and 4.2%, respectively. The increase reflects organic growth, which was partially offset by unfavorable fluctuations in currencies. At March 31, 2020, the Healthcare Products segment's backlog amounted to \$170.1 million, increasing \$15.6 million, or 10.1%, as compared to the backlog of \$154.5 million at March 31, 2019.

Healthcare Specialty Services revenues increased 10.5% in fiscal 2020, as compared to fiscal 2019. The increase reflects organic growth, which was partially offset by the divestiture of our Healthcare Specialty Services business in China and unfavorable fluctuations in currencies.

Life Sciences revenues increased 10.1% in fiscal 2020, as compared to fiscal 2019, reflecting growth in consumable, capital equipment and service revenues of 14.9%, 9.8% and 3.7%, respectively. The increase reflects organic growth and favorable pricing, which were partially offset by unfavorable fluctuations in currencies. Life Sciences backlog at March 31, 2020 amounted to \$72.4 million, increasing \$11.7 million, or 19.3%, as compared to backlog of \$60.7 million at March 31, 2019.

Applied Sterilization Technologies revenues increased 13.0% in fiscal 2020, as compared to fiscal 2019. The increase reflects organic growth, which was primarily attributable to increased demand from medical device Customers, which was partially offset by unfavorable fluctuations in currencies.

⁽²⁾ Acquisition and integration related charges include transaction costs and integration expenses associated with acquisitions.

⁽³⁾ Costs incurred in connection with the Redomiciliation and subsequent tax restructuring.

⁽⁴⁾ For more information regarding our restructuring activities see Note 2 titled, "Restructuring".

⁽⁵⁾ COVID-19 incremental costs includes the additional costs attributable to COVID-19 such as enhanced cleaning protocols, personal protective equipment for our employees, event cancellation fees, and payroll costs associated with our response to COVID-19, net of any government subsidies available.

The Healthcare Products segment's operating income increased \$32.7 million to \$356.4 million in fiscal year 2020, as compared to \$323.7 million in fiscal year 2019. The segment's operating margins were 25.0% for fiscal year 2020 and 24.2% for fiscal year 2019. The increases in the fiscal 2020 period were primarily due to increased volumes and favorable product mix.

The Healthcare Specialty Services segment's operating income was flat at \$64.2 million in fiscal years 2020 and 2019. The segment's operating margins were 11.4% for fiscal year 2020 and 12.6% for fiscal year 2019. The fiscal 2020 operating margin benefited from increased volumes, which were more than offset by investments being made to add capacity in anticipation of continuing demand.

The Life Sciences business segment's operating income increased \$12.0 million to \$144.1 million in fiscal year 2020, as compared to \$132.1 million in fiscal year 2019, primarily due to increased volumes. The segment's operating margins were 34.6% for fiscal year 2020 and 34.9% for fiscal year 2019. The decline in the fiscal 2020 operating margin was primarily due to unfavorable product mix.

The Applied Sterilization Technologies segment's operating income increased \$49.1 million to \$270.9 million in fiscal year 2020, as compared to \$221.8 million in fiscal year 2019. The Applied Sterilization Technologies segment's operating margins were 43.2% for fiscal year 2020 and 40.0% for fiscal year 2019. The increases in the fiscal 2020 period were primarily due to increased volumes.

Effective April 1, 2020, and consistent with the way management will operate and view the business, the current Healthcare Products and Healthcare Specialty Services segments will be combined and reported as one segment, simply called Healthcare. Going forward we will operate and report in three business segments: Healthcare, Life Sciences and Applied Sterilization Technologies. Corporate will continue to be presented separately and contain the costs that are associated with being a publicly traded company and certain other corporate costs.

LIQUIDITY AND CAPITAL RESOURCES

The following table summarizes significant components of our cash flows for the years ended March 31, 2020 and 2019:

	Years Ende					
(dollars in thousands)	 2020		2019			
Net cash provided by operating activities	\$ 590,559	\$	539,505			
Net cash used in investing activities	(319,735)		(213,224)			
Net cash used in financing activities	(163,146)		(294,792)			
Debt-to-total capital ratio	25.3%)	27.1%			
Free cash flow	\$ 380,199	\$	355,357			

Net Cash Provided By Operating Activities – The net cash provided by our operating activities was \$590.6 million for the year ended March 31, 2020 compared to \$539.5 million for the year ended March 31, 2019. The following discussion summarizes the significant changes in our operating cash flows for the years ended March 31, 2020 and 2019:

Net cash provided by operating activities increased in fiscal 2020 by 9.5%, as compared to fiscal 2019, primarily due to
higher net income attainment in the fiscal 2020 period, which was partially offset by higher cash requirements to fund
operating assets and liabilities.

Net Cash Used In Investing Activities – The net cash used in our investing activities was \$319.7 million for the year ended March 31, 2020, compared to \$213.2 million for the year ended March 31, 2019. The following discussion summarizes the significant changes in our investing cash flows for the years ended March 31, 2020 and 2019:

- <u>Purchases of property, plant, equipment, and intangibles, net</u> Capital expenditures totaled \$214.5 million and \$189.7 million for fiscal 2020 and 2019, respectively. The fiscal 2020 increase was primarily due to our previously announced expansion projects in the Applied Sterilization Technologies and Healthcare Specialty Services segments.
- Proceeds from the sale of property, plant, equipment and intangibles During fiscal 2020 and 2019 we received \$4.2 million and \$5.6 million respectively, for proceeds from the sale of property, plant, equipment and intangibles. The majority of the fiscal 2020 and fiscal 2019 proceeds were related to the sale of Healthcare Products facilities located in the U.K.
- <u>Proceeds from the sale of business</u> During fiscal 2020 and 2019 we received \$0.4 million and \$2.5 million, respectively, for proceeds from the sale of certain non-core businesses. For more information, refer to our Note 18 to our consolidated financial statements, titled "Business Acquisitions and Divestitures".

- Purchases of investments During fiscal 2019, we completed an equity investment for approximately \$5.0 million.
- <u>Investments in business, net of cash acquired</u> During fiscal 2020 and 2019, we used \$109.8 million and \$13.3 million, respectively, for acquisitions. For more information on these acquisitions refer to Note 18 to our consolidated financial statements titled, "Business Acquisitions and Divestitures".
- Other During fiscal 2019 we provided approximately \$13.4 million under borrowing agreements. For more information on these agreements refer to our Note 18 to our consolidated financial statements, titled "Business Acquisitions and Divestitures".

Net Cash Used In Financing Activities – Net cash used in financing activities was \$163.1 million for the year ended March 31, 2020, compared to net cash used in financing activities of \$294.8 million for the year ended March 31, 2019. The following discussion summarizes the significant changes in our financing cash flows for the years ended March 31, 2020 and 2019:

- Payments on long-term obligations During fiscal 2019 we repaid \$85.0 million in private placement notes that matured on August 15, 2018. For more information on our debt refer to Note 6 to our consolidated financial statements titled, "Debt".
- (Payments) proceeds under credit facilities, net At the end of fiscal 2020, \$275.4 million of debt was outstanding under our bank credit facility, compared to \$301.8 million of debt outstanding under this facility at the end of fiscal 2019. We provide additional information about our bank credit facility in Note 6 to our consolidated financial statements titled, "Debt".
- Repurchases of shares During fiscal 2020, we purchased 273,259 of our ordinary shares in the aggregate amount of \$40.0 million. We also obtained 122,884 of our ordinary shares in connection with our stock-based compensation award programs in the amount of \$11.2 million. During fiscal 2019, we purchased 659,393 of our ordinary shares in the aggregate amount of \$73.2 million, which included \$0.4 million of taxes and commissions. We also obtained 112,356 of our ordinary shares in connection with our stock-based compensation award programs in the amount \$8.3 million. We provide additional information about our share repurchases in Note 13 to our consolidated financial statements titled, "Repurchases of Ordinary Shares."
- <u>Deferred financing fees and debt issuance costs</u> We paid \$1.3 million and \$0.5 million in fiscal 2020 and 2019 respectively, for financing fees and debt issuance costs related to our Credit Agreement and Private Placement debt. For more information on our debt refer to Note 6 to our consolidated financial statements titled, "Debt".
- <u>Cash dividends paid to ordinary shareholders</u> During fiscal 2020, we paid cash dividends totaling \$123.0 million or \$1.45 per outstanding share. During fiscal 2019, we paid cash dividends totaling \$112.5 million or \$1.33 per outstanding share.
- Stock option and other equity transactions, net We generally receive cash for issuing shares upon the exercise of options under our employee stock option program. During fiscal 2020 and fiscal 2019, we received cash proceeds totaling \$34.7 million and \$13.3 million, respectively, under these programs. During fiscal 2020, we received contributions from noncontrolling interest holders of \$6.1 million and paid \$1.2 million in distributions to noncontrolling interest holders. During fiscal 2019 we paid \$0.3 million in distributions to noncontrolling interest holders.

Cash Flow Measures. Free cash flow was \$380.2 million in fiscal 2020 compared to \$355.4 million in fiscal 2019. The increase in free cash flow is primarily due to the improvement in cash from operations.

Our debt-to-total capital ratio was 25.3% at March 31, 2020 and 27.1% at March 31, 2019.

Cash Requirements. We intend to use our existing cash and cash equivalent balances and cash generated from operations to fund capital expenditures and meet our other liquidity needs. Our capital requirements depend on many uncertain factors, including our rate of sales growth, our Customers' acceptance of our products and services, the costs of obtaining adequate manufacturing capacities, the timing and extent of our research and development projects, changes in our operating expenses and other factors. To the extent that existing and anticipated sources of cash are not sufficient to fund our future activities, we may need to raise additional funds through additional borrowings or the sale of equity securities. There can be no assurance that our financing arrangements will provide us with sufficient funds or that we will be able to obtain any additional funds on terms favorable to us or at all.

Sources of Credit. Our sources of credit as of March 31, 2020 are summarized in the following table:

(dollars in thousands)	Maximum Amounts Available	Reductions in Available Credit acility for Other Financial Instruments	N	March 31, 2020 Amounts Outstanding	N	March 31, 2020 Amounts Available
Sources of Credit						
Private placement	\$ 878,409	\$ _	\$	878,409	\$	_
Credit Agreement (1)	1,000,000	6,768		275,449		717,783
Total Sources of Credit	\$ 1,878,409	\$ 6,768	\$	1,153,858	\$	717,783

⁽¹⁾ At March 31, 2020, there was \$6.8 million of letters of credit outstanding under the Credit Agreement.

Our sources of funding from credit as of March 31, 2020 are summarized below:

- On March 23, 2018, STERIS UK and certain of its subsidiaries entered into a Credit Agreement (the "Credit Agreement") with various financial institutions as lenders, and JPMorgan Chase Bank, N.A., as Administrative Agent. STERIS Ireland subsequently became a borrower and guarantor under the Credit Agreement. The Credit Agreement replaced a bank credit facility dated March 31, 2015. The Credit Agreement provides up to \$1.0 billion of credit, in the form of a revolver facility, which may be utilized for revolving credit borrowings, swing line borrowings and letters of credit, with sublimits for swing line borrowings and letters of credit. The revolver facility may be increased in specified circumstances by up to \$500.0 million. The Credit Agreement will mature on March 23, 2023, and all unpaid borrowings, together with accrued and unpaid interest thereon, are repayable on that date. The Credit Agreement contains leverage and interest coverage covenants. Borrowings may be taken in U.S. dollars, euros, and pounds sterling and certain other specified currencies and bear interest at our option based upon either the Base Rate or the Eurocurrency Rate, plus the Applicable Margin in effect from time to time under the Credit Agreement. The Applicable Margin is determined based on the ratio of Consolidated Total Debt to Consolidated EBITDA (as such terms are defined in the Credit Agreement). Interest on Base Rate Advances is payable quarterly in arrears and interest on Eurocurrency Rate Advances is payable at the end of the relevant interest period therefor, but in no event less frequently than every three months. Borrowings at closing were used to repay outstanding balances of debt outstanding under the former bank credit facility dated March 31, 2015 that was scheduled to mature on March 31, 2020 and for other general corporate purposes.
- The Credit Agreement was amended in March 2019, in connection with the Redomiciliation to permit the Redomiciliation.
 The amendments did not effect any material changes in the terms of the Credit Agreement regarding borrowings or the issuance of letters of credit.

Our outstanding Senior Notes at March 31, 2020 were as follows:

(dollars in thousands)	Applicable Note Purchase Agreement	Maturity Date	U.S. Dollar Value at March 31, 2020
\$35,000 Senior notes at 6.43%	2008 Private Placement	August 2020	35,000
\$91,000 Senior notes at 3.20%	2012 Private Placement	December 2022	91,000
\$80,000 Senior notes at 3.35%	2012 Private Placement	December 2024	80,000
\$25,000 Senior notes at 3.55%	2012 Private Placement	December 2027	25,000
\$125,000 Senior notes at 3.45%	2015 Private Placement	May 2025	125,000
\$125,000 Senior notes at 3.55%	2015 Private Placement	May 2027	125,000
\$100,000 Senior notes at 3.70%	2015 Private Placement	May 2030	100,000
\$50,000 Senior notes at 3.93%	2017 Private Placement	February 2027	50,000
€60,000 Senior notes at 1.86%	2017 Private Placement	February 2027	66,342
\$45,000 Senior notes at 4.03%	2017 Private Placement	February 2029	45,000
€20,000 Senior notes at 2.04%	2017 Private Placement	February 2029	22,114
£45,000 Senior notes at 3.04%	2017 Private Placement	February 2029	55,767
€19,000 Senior notes at 2.30%	2017 Private Placement	February 2032	21,008
£30,000 Senior notes at 3.17%	2017 Private Placement	February 2032	37,178
Total Senior Notes			\$ 878,409

- On February 27, 2017, STERIS UK issued and sold an aggregate principal amount of \$95.0 million, €99.0 million, and £75.0 million, of senior notes in a private placement to certain institutional investors in an offering that was exempt from the registration requirements of the Securities Act of 1933. These notes have maturities of between 10 and 15 years from the issue date. The agreement governing these notes contains leverage and interest coverage covenants.
- On May 15, 2015, STERIS Corporation issued and sold \$350.0 million of senior notes, in a private placement to certain
 institutional investors in an offering that was exempt from the registration requirements of the Securities Act of 1933.
 These notes have maturities of 10 to 15 years from the issue date. The agreement governing these notes contains leverage
 and interest coverage covenants.
- The agreements governing certain senior notes issued and sold in February 2013, December 2012, and August 2008, were amended and restated in their entirety on March 31, 2015. All of these notes were issued and sold in private placements to certain institutional investors in offerings that were exempt from the registration requirements of the Securities Act of 1933. The amended and restated agreements, which have been consolidated into a single agreement for the 2013 and 2012 notes, and a separate single agreement for the 2008 notes, contain leverage and interest coverage covenants.
- All of the note agreements were amended in March 2019, in connection with the Redomiciliation. The amendments waived
 certain repurchase rights of the note holders and increased the size of certain baskets to more closely align with Credit
 Agreement baskets.

As of March 31, 2020, a total of \$275.4 million was outstanding under the Credit Agreement, based on currency exchange rates as of March 31, 2020. At March 31, 2020, we had \$717.8 million of unused funding available under the Credit Agreement. The Credit Agreement includes a sub-limit that reduces the maximum amount available to us by letters of credit outstanding. At March 31, 2020, there was \$6.8 million in letters of credit outstanding under the Credit Agreement.

At March 31, 2020, we were in compliance with all financial covenants associated with our indebtedness. We provide additional information regarding our debt structure and payment obligations in the section of the MD&A titled, "Liquidity and Capital Resources" in the subsection titled, "Contractual and Commercial Commitments" and in Note 6 to our consolidated financial statements titled, "Debt."

CAPITAL EXPENDITURES

Our capital expenditure program is a component of our long-term strategy. This program includes, among other things, investments in new and existing facilities, business expansion projects, radioisotope (cobalt-60), and information technology enhancements and research and development advances. During fiscal 2020, our capital expenditures amounted to \$214.5 million. We use cash provided by operating activities and our cash and cash equivalent balances to fund capital expenditures. In fiscal 2021, we expect to continue to invest in facility expansions, particularly within the Applied Sterilization Technologies segment and in ongoing maintenance for existing facilities. The outbreak of COVID-19 has become a global pandemic. We may choose to temporarily defer planned capital expenditures due to fluctuations in demand for our products and services resulting from the COVID-19 pandemic and our Customers' needs.

CONTRACTUAL AND COMMERCIAL COMMITMENTS

At March 31, 2020, we had commitments under non-cancelable operating leases totaling \$173.9 million.

Our contractual obligations and commercial commitments as of March 31, 2020 are presented in the following tables. Commercial commitments include standby letters of credit, letters of credit required as security under our self-insured risk retention policies, and other potential cash outflows resulting from events that require us to fulfill commitments.

	Payments due by March 31,											
(dollars in thousands)	2021		2022		2023		2024		2025 and thereafter		Total	
Contractual Obligations:				-							_	
Debt	\$	35,000	\$		\$	366,449	\$		\$	752,409	\$ 1,153,858	
Operating leases		25,302		21,064		17,271		14,045		96,249	173,931	
Purchase obligations		67,866		75,968		10,297				_	154,131	
Benefit payments under defined benefit plans		5,872		6,025		6,600		6,336		41,810	66,643	
Trust assets available for benefit payments under defined benefit plans		(5,872)		(6,025)		(6,600)		(6,336)		(41,810)	(66,643)	
Benefit payments under other post- retirement benefits plans		1,510		1,392		1,252		1,115		4,733	10,002	
Expected contributions to defined benefit plans		3,839		3,954		1,991		_		_	9,784	
Total Contractual Obligations	\$	133,517	\$	102,378	\$	397,260	\$	15,160	\$	853,391	\$ 1,501,706	

The table above includes only the principal amounts of our contractual obligations. We provide information about the interest component of our long-term debt in the subsection of MD&A titled, "Liquidity and Capital Resources," and in Note 6 to our consolidated financial statements titled, "Debt."

Purchase obligations shown in the table above relate to minimum purchase commitments with suppliers for materials purchases and long term construction contracts.

The table above excludes contributions we make to our defined contribution plans. Our future contributions to the defined contribution plans depend on uncertain factors, such as the amount and timing of employee contributions and discretionary employer contributions. We provide additional information about our defined benefit pension plans, defined contribution plan, and other post-retirement benefits plan in Note 9 to our consolidated financial statements titled, "Benefit Plans."

	Amount of Commitment Expiring March 31,									
(dollars in thousands)	2021		2022		2023		2024		2025 and hereafter	Totals
Commercial Commitments:										
Letters of credit and surety bonds	\$ 56,899	\$	7,062	\$	1,118	\$	353	\$	2,324	\$ 67,756
Letters of credit as security for self- insured risk retention policies	 12,474		_		_		_		_	12,474
Total Commercial Commitments	\$ 69,373	\$	7,062	\$	1,118	\$	353	\$	2,324	\$ 80,230

CRITICAL ACCOUNTING POLICIES, ESTIMATES, AND ASSUMPTIONS

The following subsections describe our most critical accounting policies, estimates, and assumptions. Our accounting policies are more fully described in Note 1 to our consolidated financial statements titled, "Nature of Operations and Summary of Significant Accounting Policies."

Estimates and Assumptions. Our discussion and analysis of financial condition and results of operations is based on our consolidated financial statements that were prepared in accordance with United States generally accepted accounting principles. We make certain estimates and assumptions that we believe to be reasonable when preparing these financial statements. These estimates and assumptions involve judgments with respect to numerous factors that are difficult to predict and are beyond management's control. As a result, actual amounts could be materially different from these estimates. We periodically review these critical accounting policies, estimates, assumptions, and the related disclosures with the Audit Committee of the Company's Board of Directors.

Revenue Recognition. Revenue is recognized when obligations under the terms of the contract are satisfied and control of the promised products or services has transferred to the Customer. Revenues are measured at the amount of consideration that we expect to be paid in exchange for the products or services. Product revenue is recognized when control passes to the Customer, which is generally based on contract or shipping terms. Service revenue is recognized when the Customer benefits from the service, which occurs either upon completion of the service or as it is provided to the Customer. Our Customers include end users as well as dealers and distributors who market and sell our products. Our revenue is not contingent upon resale by the dealer or distributor, and we have no further obligations related to bringing about resale. Our standard return and restocking fee policies are applied to sales of products. Shipping and handling costs charged to Customers are included in Product revenues. The associated expenses are treated as fulfillment costs and are included in Cost of revenues. Revenues are reported net of sales and value-added taxes collected from Customers.

We have individual Customer contracts that offer discounted pricing. Dealers and distributors may be offered sales incentives in the form of rebates. We reduce revenue for discounts and estimated returns, rebates, and other similar allowances in the same period the related revenues are recorded. The reduction in revenue for these items is estimated based on historical experience and trend analysis to the extent that it is probable that a significant reversal of revenue will not occur. Estimated returns are recorded gross on the Consolidated Balance Sheets.

In transactions that contain multiple performance obligations, such as when products, maintenance services, and other services are combined, we recognize revenue as each product is delivered or service is provided to the Customer. We allocate the total arrangement consideration to each performance obligation based on its relative standalone selling price, which is the price for the product or service when it is sold separately.

Payment terms vary by the type and location of the Customer and the products or services offered. Generally, the time between when revenue is recognized and when payment is due is not significant. We do not evaluate whether the selling price contains a financing component for contracts that have a duration of less than one year.

We do not capitalize sales commissions as substantially all of our sales commission programs have an amortization period of one year or less.

Certain costs to fulfill a contract are capitalized and amortized over the term of the contract if they are recoverable, directly related to a contract and generate resources that we will use to fulfill the contract in the future. At March 31, 2020 assets related to costs to fulfill a contract were not material to our Consolidated Financial Statements.

Allowance for Doubtful Accounts Receivable. We maintain an allowance for uncollectible accounts receivable for estimated losses in the collection of amounts owed by Customers. We estimate the allowance based on analyzing a number of factors, including amounts written off historically, Customer payment practices, and general economic conditions. We also analyze significant Customer accounts on a regular basis and record a specific allowance when we become aware of a specific Customer's inability to pay. As a result, the related accounts receivable are reduced to an amount that we reasonably believe is collectible. These analyses require judgment. If the financial condition of our Customers worsens, or economic conditions change, we may be required to make changes to our allowance for doubtful accounts receivable.

Allowance for Sales Returns. We maintain an allowance for sales returns based upon known returns and estimated returns for both capital equipment and consumables. We estimate returns of capital equipment and consumables based upon historical experience.

Inventories and Reserves. Inventories are stated at the lower of their cost or market value. We determine cost based upon a combination of the last-in, first-out ("LIFO") and first-in, first-out ("FIFO") cost methods. We determine the LIFO inventory value at the end of the year based on inventory levels and costs at that time. For inventories valued using the LIFO method, we believe that the use of the LIFO method results in a matching of current costs and revenues. Inventories valued using the LIFO method represented approximately 25.3% and 25.2% of total inventories at March 31, 2020 and 2019, respectively. Inventory costs include material, labor, and overhead. If we had used only the FIFO method of inventory costing, inventories would have been \$16.9 million and \$16.8 million higher than those reported at March 31, 2020 and 2019, respectively.

We review inventory on an ongoing basis, considering factors such as deterioration and obsolescence. We record an allowance for estimated losses when the facts and circumstances indicate that particular inventories will not be usable. If future market conditions vary from those projected, and our estimates prove to be inaccurate, we may be required to write-down inventory values and record an adjustment to cost of revenues.

Asset Impairment Losses. Property, plant, equipment, and identifiable intangible assets are reviewed for impairment when events and circumstances indicate that the carrying value of such assets may not be recoverable. Impaired assets are recorded at the lower of carrying value or estimated fair value. We conduct this review on an ongoing basis and, if impairment exists, we record the loss in the Consolidated Statements of Income during that period.

When we evaluate assets for impairment, we make certain judgments and estimates, including interpreting current economic indicators and market valuations, evaluating our strategic plans with regards to operations, historical and anticipated performance of operations, and other factors. If we incorrectly anticipate these factors, or unexpected events occur, our operating results could be materially affected.

Asset Retirement Obligations. We incur retirement obligations for certain assets. We record an initial liability for the asset retirement obligations (ARO) at fair value. Accounting for the ARO at inception and in subsequent periods includes the determination of the present value of a liability and offsetting asset, the subsequent accretion of that liability and depletion of the asset, and a periodic review of the ARO liability estimates and discount rates used in the analysis. We provide additional information about our asset retirement obligations in Note 5 to our consolidated financial statements titled, "Property, Plant and Equipment."

Restructuring. We record specific accruals in connection with plans for restructuring elements of our business. These accruals include estimates principally related to employee separation costs, the closure and/or consolidation of facilities, and contractual obligations. Actual amounts could differ from the original estimates. We review our restructuring-related accruals on a quarterly basis and changes to plans are appropriately recognized in the Consolidated Statements of Income in the period the change is identified.

Purchase Accounting and Goodwill. Assets and liabilities of the business acquired are accounted for at their estimated fair values as of the acquisition date. Any excess of the cost of the acquisition over the fair value of the net tangible and intangible assets acquired is recorded as goodwill. We supplement management expertise with valuation specialists in performing appraisals to assist us in determining the fair values of assets acquired and liabilities assumed. These valuations require us to make estimates and assumptions, especially with respect to intangible assets. We generally amortize our intangible assets over their useful lives with the exception of indefinite lived intangible assets. We do not amortize goodwill, but we evaluate it annually for impairment. Therefore, the allocation of the purchase price to intangible assets and goodwill has a significant impact on future operating results.

We evaluate the recoverability of recorded goodwill amounts annually, or when evidence of potential impairment exists. We may consider qualitative indicators of the fair value of a reporting unit when it is unlikely that a reporting unit has impaired goodwill. We may also utilize a discounted cash flow analysis that requires certain assumptions and estimates be made regarding market conditions and our future profitability. In those circumstances, we test goodwill for impairment by reviewing the book value compared to the fair value at the reporting unit level. We calculate the fair value of our reporting units based on the present value of estimated future cash flows. Management's judgment is necessary to evaluate the impact of operating and macroeconomic changes and to estimate future cash flows to measure fair value. Assumptions used in our impairment evaluations, such as forecasted growth rates and cost of capital, are consistent with internal projections and operating plans. We believe such assumptions and estimates are also comparable to those that would be used by other marketplace participants.

As a result of our annual impairment review for goodwill and other indefinite lived intangible assets for fiscal year 2020 no indicators of impairment were identified.

We evaluate indefinite lived intangible assets annually, or when evidence of potential impairment exists. We evaluate several qualitative indicators and assumptions, and trends that influence the valuation of the assets to determine if any evidence of potential impairment exists. During the third quarter of fiscal 2019, management adopted a branding strategy that included phasing out the usage of a tradename associated with certain products in the Healthcare Products business segment. As a result, management recorded an impairment charge of \$16.2 million, which is included within the Selling, general, and administrative line of the Consolidated Statements of Income. The remaining fair value of the asset was calculated using an income approach (the relief from royalty method). The remaining fair value was not material and was amortized over the asset's remaining useful life. Fair value calculated using this approach is classified within Level 3 of the fair value hierarchy and requires several assumptions.

Income Taxes. Our provision for income taxes is based on our current period income, changes in deferred income tax assets and liabilities, income tax rates, changes in uncertain tax benefits, and tax planning opportunities available to us in the various jurisdictions in which we operate. Tax laws are complex and subject to different interpretations by the taxpayer and the respective governmental taxing authorities. We use judgment in determining our annual effective income tax rate and evaluating our tax positions. We prepare and file tax returns based on our interpretation of tax laws and regulations, and we record estimates based on these judgments and interpretations. We cannot be sure that the tax authorities will agree with all of the tax positions taken by us. The actual income tax liability for each jurisdiction in any year can, in some instances, ultimately be determined be several years after the tax return is filed and the financial statements are published.

We evaluate our tax positions using the recognition threshold and measurement attribute in accordance with current accounting guidance. We determine whether it is more-likely-than-not that a tax position will be sustained upon examination, including resolution of related appeals or litigation processes, based on the technical merits of the position. In evaluating whether a tax position has met the more-likely-than-not recognition threshold, we presume that the position will be examined by the appropriate taxing authority and that the taxing authority will have full knowledge of all relevant information. A tax position that meets the more-likely-than-not recognition threshold is measured at the largest amount of benefit that is greater than 50 percent likely of being realized upon ultimate settlement. The appropriate unit of account for determining what constitutes an individual tax position, and whether the more-likely-than-not recognition threshold is met for a tax position, is a matter of judgment based on the individual facts and circumstances of that position evaluated in light of all available evidence. We review and adjust our tax estimates periodically because of ongoing examinations by and settlements with the various taxing authorities, as well as changes in tax laws, regulations and precedent.

We recognize deferred tax assets and liabilities based on the differences between the financial statement carrying amounts and the tax basis of assets and liabilities. We regularly review our deferred tax assets for recoverability and establish a valuation allowance based on historical taxable income, projected future taxable income, the expected timing of the reversals of existing temporary differences, and the implementation of tax planning strategies. If we are unable to generate sufficient future taxable income in certain tax jurisdictions, or if there is a material change in the effective income tax rates or time period within which the underlying temporary differences become taxable or deductible, we could be required to increase our valuation allowance, which would increase our effective income tax rate and could result in an adverse impact on our consolidated financial position, results of operations, or cash flows.

We believe that adequate accruals have been made for income taxes. Differences between the estimated and actual amounts determined upon ultimate resolution, individually or in the aggregate, are not expected to have a material adverse effect on our consolidated financial position, but could possibly be material to our consolidated results of operations or cash flows for any one period.

Additional information regarding income taxes is included in Note 8 to our consolidated financial statements titled, "Income Taxes."

Self-Insurance Liabilities. We record a liability for self-insured risks that we retain for general and product liabilities, workers' compensation, and automobile liabilities based on actuarial calculations. We use our historical loss experience and actuarial methods to calculate the estimated liability. This liability includes estimated amounts for both losses and incurred but not reported claims. We review the assumptions used to calculate the estimated liability at least annually to evaluate the adequacy of the amount recorded. We maintain insurance policies to cover losses greater than our estimated liability, which are subject to the terms and conditions of those policies. The obligation covered by insurance contracts will remain on the balance sheet as we remain liable to the extent insurance carriers do not meet their obligation. Estimated amounts receivable under the contracts are included in the "Prepaid expenses and other current assets" line, and the "Other assets" line of our consolidated balance sheets. Our accrual for self-insured risk retention as of March 31, 2020 and 2019 was \$23.2 million and \$19.7 million, respectively.

We are also self-insured for employee medical claims. We estimate a liability for incurred but not reported claims based upon recent claims experience. Our self-insured liabilities contain uncertainties because management must make assumptions and apply judgments to estimate the ultimate cost to settle reported claims and claims incurred but not reported as of the balance sheet date. If actual results are not consistent with these assumptions and judgments, we could be exposed to additional costs in subsequent periods.

Contingencies. We are, and will likely continue to be, involved in a number of legal proceedings, government investigations, and claims, which we believe generally arise in the course of our business, given our size, history, complexity, and the nature of our business, products, Customers, regulatory environment, and industries in which we participate. These legal proceedings, investigations and claims generally involve a variety of legal theories and allegations, including, without limitation, personal injury (e.g., slip and falls, burns, vehicle accidents), product liability or regulation (e.g., based on product operation or claimed malfunction, failure to warn, failure to meet specification, or failure to comply with regulatory requirements), product exposure (e.g., claimed exposure to chemicals, asbestos, contaminants, radiation), property damage (e.g., claimed damage due to leaking equipment, fire, vehicles, chemicals), commercial claims (e.g., breach of contract, economic loss, warranty, misrepresentation), financial (e.g., taxes, reporting), employment (e.g., wrongful termination, discrimination, benefits matters), and other claims for damage and relief.

We record a liability for such contingencies to the extent we conclude that their occurrence is both probable and estimable. We consider many factors in making these assessments, including the professional judgment of experienced members of management and our legal counsel. We have made estimates as to the likelihood of unfavorable outcomes and the amounts of such potential losses. In our opinion, the ultimate outcome of these proceedings and claims is not anticipated to have a material adverse affect on our consolidated financial position, results of operations, or cash flows. However, the ultimate outcome of

proceedings, government investigations, and claims is unpredictable and actual results could be materially different from our estimates. We record expected recoveries under applicable insurance contracts when we are assured of recovery. Refer to Note 10 of our consolidated financial statements titled, "Commitments and Contingencies" for additional information.

We are subject to taxation from federal, state and local, and foreign jurisdictions. Tax positions are settled primarily through the completion of audits within each individual tax jurisdiction or the closing of a statute of limitation. Changes in applicable tax law or other events may also require us to revise past estimates. The IRS of the United States routinely conducts audits of our federal income tax returns.

Additional information regarding our commitments and contingencies is included in Note 10 to our consolidated financial statements titled, "Commitments and Contingencies."

Benefit Plans. We provide defined benefit pension plans for certain employees and retirees. In addition, we sponsor an unfunded post-retirement benefits plan for two groups of United States retirees. Benefits under this plan include retiree life insurance and retiree medical insurance, including prescription drug coverage.

Employee pension and post-retirement benefits plans are a cost of conducting business and represent obligations that will be settled in the future and therefore, require us to use estimates and make certain assumptions to calculate the expense and liabilities related to the plans. Changes to these estimates and assumptions can result in different expense and liability amounts. Future actual experience may be significantly different from our current expectations. We believe that the most critical assumptions used to determine net periodic benefit costs and projected benefit obligations are the expected long-term rate of return on plan assets and the discount rate. A summary of significant assumptions used to determine the March 31, 2020 projected benefit obligations and the fiscal 2020 net periodic benefit costs is as follows:

	Synergy Health plc	Isotron BV	Synergy Health Daniken AG	Synergy Health Radeberg	Synergy Health Allershausen	Harwell Dosimeters Ltd	U.S. Post- Retirement Benefits Plan
Funding Status	Funded	Funded	Funded	Unfunded	Unfunded	Funded	Unfunded
Assumptions used to determine March 31, 2020							
Benefit obligations:							
Discount rate	2.40%	1.60%	0.20%	1.60%	0.50%	2.45%	3.00%
Assumptions used to determine fiscal 2020							
Net periodic benefit costs:							
Discount rate	2.50%	1.20%	0.20%	1.60%	1.75%	2.45%	3.50%
Expected return on plan assets	4.80%	1.20%	0.65%	n/a	n/a	n/a	n/a
371 37 . 11 11							

NA – Not applicable.

We develop our expected long-term rate of return on plan assets assumptions by evaluating input from third-party professional advisors, taking into consideration the asset allocation of the portfolios, and the long-term asset class return expectations. Generally, net periodic benefit costs increase as the expected long-term rate of return on plan assets assumption decreases. Holding all other assumptions constant, lowering the expected long-term rate of return on plan assets assumption for our funded defined benefit pension plans by 50 basis points would have increased the fiscal 2020 benefit costs by less than \$0.1 million.

We develop our discount rate assumptions by evaluating input from third-party professional advisers, taking into consideration the current yield on country specific investment grade long-term bonds which provide for similar cash flow streams as our projected benefit obligations. Generally, the projected benefit obligations and the net periodic benefit costs both increase as the discount rate assumption decreases. Holding all other assumptions constant, lowering the discount rate assumption for our defined benefit pension plans and for the other post-retirement benefits plan by 50 basis points would have decreased the fiscal 2020 net periodic benefit costs by less than \$0.1 million and would have increased the projected benefit obligations by approximately \$10.5 million at March 31, 2020.

We have made assumptions regarding healthcare costs in computing our other post-retirement benefit obligation. The assumed rates of increase generally decline ratably over a five year-period from the assumed current year healthcare cost trend rate of 6.8% to the assumed long-term healthcare cost trend rate. A 100 basis point change in the assumed healthcare cost trend rate (including medical, prescription drug, and long-term rates) would have had the following effect at March 31, 2020:

	100 E	Basis F	oint
(dollars in thousands)	Increase		Decrease
Effect on total service and interest cost components	\$ -	- \$	_
Effect on postretirement benefit obligation		7	(6)

We recognize an asset for the overfunded status or a liability for the underfunded status of defined benefit pension and post-retirement benefit plans in our balance sheets. This amount is measured as the difference between the fair value of plan assets and the benefit obligation (the projected benefit obligation for pension plans and the accumulated post-retirement benefit obligation for other post-retirement benefit plans). Changes in the funded status of the plans are recorded in other comprehensive income in the year they occur. We measure plan assets and obligations as of the balance sheet date. Note 9 to our consolidated financial statements titled, "Benefit Plans," contains additional information about our pension and other post-retirement welfare benefits plans.

Share-Based Compensation. We measure the estimated fair value for share-based compensation awards, including grants of employee stock options, at the grant date and recognize the related compensation expense over the period in which the share-based compensation vests. We selected the Black-Scholes-Merton option pricing model as the most appropriate method for determining the estimated fair value of our share-based stock option compensation awards. This model involves assumptions that are judgmental and affect share-based compensation expense.

Share-based compensation expense was \$23.8 million in fiscal 2020, \$24.0 million in fiscal 2019 and \$22.2 million in fiscal 2018. Note 14 to our consolidated financial statements titled, "Share-Based Compensation," contains additional information about our share-based compensation plans.

RECENTLY ISSUED ACCOUNTING STANDARDS IMPACTING THE COMPANY

Recently issued accounting standards that are relevant to us are presented in Note 1 to our consolidated financial statements titled, "Nature of Operations and Summary of Significant Accounting Policies."

INFLATION

Our business has not been significantly impacted by the overall effects of inflation. We monitor the prices we charge for our products and services on an ongoing basis and plan to adjust those prices to take into account future changes in the rate of inflation. However, we may not be able to completely offset the impact of inflation.

FORWARD-LOOKING STATEMENTS

This Form 10-K may contain statements concerning certain trends, expectations, forecasts, estimates, or other forward-looking information affecting or relating to STERIS or its industry, products or activities that are intended to qualify for the protections afforded "forward-looking statements" under the Private Securities Litigation Reform Act of 1995 and other laws and regulations. Forward-looking statements speak only as to the date the statement is made and may be identified by the use of forward-looking terms such as "may," "will," "expects," "believes," "anticipates," "plans," "estimates," "projects," "targets," "forecasts," "outlook," "impact," "potential," "confidence," "improve," "optimistic," "deliver," "orders," "backlog," "comfortable," "trend", and "seeks," or the negative of such terms or other variations on such terms or comparable terminology. Many important factors could cause actual results to differ materially from those in the forward-looking statements including. without limitation, disruption of production or supplies, changes in market conditions, political events, pending or future claims or litigation, competitive factors, technology advances, actions of regulatory agencies, and changes in laws, government regulations, labeling or product approvals or the application or interpretation thereof. Other risk factors are described herein and in STERIS's other securities filings, including Item 1A of this Annual Report on Form 10-K. Many of these important factors are outside of STERIS's control. No assurances can be provided as to any result or the timing of any outcome regarding matters described in STERIS's securities filings or otherwise with respect to any regulatory action, administrative proceedings, government investigations, litigation, warning letters, cost reductions, business strategies, earnings or revenue trends or future financial results. References to products are summaries only and should not be considered the specific terms of the product clearance or literature. Unless legally required, STERIS does not undertake to update or revise any forward-looking statements even if events make clear that any projected results, express or implied, will not be realized. Other potential risks and uncertainties that could cause actual results to differ materially from those in the forward-looking statements include, without limitation, (a) the impact of the COVID-19 pandemic on STERIS's operations, performance, results, prospects, or value, (b)

STERIS's ability to achieve the expected benefits regarding the accounting and tax treatments of the Redomiciliation transaction, (c) operating costs, Customer loss and business disruption (including, without limitation, difficulties in maintaining relationships with employees, Customers, clients or suppliers) being greater than expected following the Redomiciliation, (d) STERIS's ability to meet expectations regarding the accounting and tax treatment of the Tax Cuts and Jobs Act ("TCJA") or the possibility that anticipated benefits resulting from the TCJA will be less than estimated, (e) changes in tax laws or interpretations that could increase our consolidated tax liabilities, including changes in tax laws that would result in STERIS being treated as a domestic corporation for United States federal tax purposes, (f) the potential for increased pressure on pricing or costs that leads to erosion of profit margins, (g) the possibility that market demand will not develop for new technologies, products or applications or services, or business initiatives will take longer, cost more or produce lower benefits than anticipated, (h) the possibility that application of or compliance with laws, court rulings, certifications, regulations, regulatory actions, including without limitation those relating to FDA warning notices or letters, government investigations, the outcome of any pending FDA requests, inspections or submissions, or other requirements or standards may delay, limit or prevent new product introductions, affect the production and marketing of existing products or services or otherwise affect STERIS's performance, results, prospects or value, (i) the potential of international unrest, economic downturn or effects of currencies, tax assessments, tariffs and/or other trade barriers, adjustments or anticipated rates, raw material costs or availability, benefit or retirement plan costs, or other regulatory compliance costs, (i) the possibility of reduced demand, or reductions in the rate of growth in demand, for STERIS's products and services, (k) the possibility of delays in receipt of orders, order cancellations, or delays in the manufacture or shipment of ordered products or in the provision of services, (1) the possibility that anticipated growth, cost savings, new product acceptance, performance or approvals, or other results may not be achieved, or that transition, labor, competition, timing, execution, regulatory, governmental, or other issues or risks associated with STERIS's businesses, industry or initiatives including, without limitation, those matters described in this Form 10-K and other securities filings, may adversely impact STERIS's performance, results, prospects or value, (m) the impact on STERIS and its operations, or tax liabilities, of Brexit or the exit of other member countries from the EU, and the Company's ability to respond to such impacts, (n) the impact on STERIS and its operations of any legislation, regulations or orders, including but not limited to any new trade or tax legislation, regulations or orders, that may be implemented by the U.S. administration or Congress, or of any responses thereto, (o) the possibility that anticipated financial results or benefits of recent acquisitions, or of STERIS's restructuring efforts, or of recent divestitures, or of the targeted restructuring plan will not be realized or will be other than anticipated, and (p) the effects of contractions in credit availability, as well as the ability of STERIS's Customers and suppliers to adequately access the credit markets when needed.

ITEM 7A. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK

In the ordinary course of business, we are exposed to various risks, including, but not limited to, interest rate, foreign currency, and commodity risks. These risks are described in the sections that follow.

INTEREST RATE RISK

As of March 31, 2020, we had \$878.4 million in fixed rate senior notes outstanding. As of March 31, 2020, we had \$275.4 million in outstanding borrowings under our Credit Agreement which are exposed to changes in interest rates. We monitor our interest rate risk, but do not engage in any hedging activities using derivative financial instruments. For additional information regarding our debt structure, refer to Note 6 to our Consolidated Financial Statements titled, "Debt."

FOREIGN CURRENCY RISK

We are exposed to the impact of foreign currency exchange fluctuations. This foreign currency exchange risk arises when we conduct business in a currency other than the U.S. dollar. For most operations, local currencies have been determined to be the functional currencies. The financial statements of subsidiaries are translated to their U.S. dollar equivalents at end-of-period exchange rates for assets and liabilities and at average currency exchange rates for revenues and expenses. Translation adjustments for subsidiaries whose local currency is their functional currency are recorded as a component of accumulated other comprehensive income (loss) within equity. Note 19 to our consolidated financial statements titled, "Reclassifications out of Accumulated Other Comprehensive Income (Loss)," contains additional information about the impact of translation on accumulated other comprehensive income (loss) and equity. Transaction gains and losses arising from fluctuations in currency exchange rates on transactions denominated in currencies other than the functional currency are recognized in the Consolidated Statements of Income. Since we operate internationally and approximately 30% of our revenues and 40% of our cost of revenues are generated outside the United States, foreign currency exchange rate fluctuations can significantly impact our financial position, results of operations, and competitive position.

We enter into foreign currency forward contracts to hedge monetary assets and liabilities denominated in foreign currencies, including inter-company transactions. We do not use derivative financial instruments for speculative purposes. At March 31, 2020, we held a foreign currency forward contract to buy 6.0 million Canadian dollars.

COMMODITY RISK

We are dependent on basic raw materials, sub-assemblies, components, and other supplies used in our operations. Our financial results could be affected by the availability and changes in prices of these materials. Some of these materials are sourced from a limited number of suppliers or only a single supplier. These materials are also key source materials for our competitors. Therefore, if demand for these materials rises, we may experience increased costs and/or limited or unavailable supplies. As a result, we may not be able to acquire key production materials on a timely basis, which could impact our ability to produce products and satisfy incoming sales orders on a timely basis. In addition, the costs of these materials can rise suddenly and result in significantly higher costs of production. We believe that we have adequate sources of supply for many of our key materials and energy sources. Where appropriate, we enter into long-term supply contracts as a basis to guarantee a reliable supply. We may also enter into commodity swap contracts to hedge price changes in a certain commodity that impacts raw materials included in our cost of revenues. At March 31, 2020, we held commodity swap contracts to buy 715,200 pounds of nickel.

ITEM 8. FINANCIAL STATEMENTS AND SUPPLEMENTARY DATA

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Report of Independent Registered Public Accounting Firm

To the Shareholders and the Board of Directors of STERIS plc

Opinion on the Financial Statements

We have audited the accompanying consolidated balance sheets of STERIS plc and subsidiaries (the Company) as of March 31, 2020 and 2019, the related consolidated statements of income, comprehensive income, shareholders' equity and cash flows for each of the three years in the period ended March 31, 2020, and the related notes and the financial statement schedule listed in the Index at Item 15(a) (collectively referred to as the "consolidated financial statements"). In our opinion, the consolidated financial statements present fairly, in all material respects, the financial position of the Company at March 31, 2020 and 2019, and the results of its operations and its cash flows for each of the three years in the period ended March 31, 2020, in conformity with U.S. generally accepted accounting principles.

We also have audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States) (PCAOB), the Company's internal control over financial reporting as of March 31, 2020, based on criteria established in Internal Control-Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission (2013 framework), and our report dated May 29, 2020 expressed an unqualified opinion thereon.

Basis for Opinion

These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on the Company's financial statements based on our audits. We are a public accounting firm registered with the PCAOB and are required to be independent with respect to the Company in accordance with the U.S. federal securities laws and the applicable rules and regulations of the Securities and Exchange Commission and the PCAOB.

We conducted our audits in accordance with the standards of the PCAOB. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement, whether due to error or fraud. Our audits included performing procedures to assess the risks of material misstatement of the financial statements, whether due to error or fraud, and performing procedures that respond to those risks. Such procedures included examining, on a test basis, evidence regarding the amounts and disclosures in the financial statements. Our audits also included evaluating the accounting principles used and significant estimates made by management, as well as evaluating the overall presentation of the financial statements. We believe that our audits provide a reasonable basis for our opinion.

Critical Audit Matter

The critical audit matter communicated below is a matter arising from the current period audit of the financial statements that was communicated or required to be communicated to the audit committee and that: (1) relates to accounts or disclosures that are material to the financial statements and (2) involved our especially challenging, subjective, or complex judgments. The communication of the critical audit matter does not alter in any way our opinion on the consolidated financial statements, taken as a whole, and we are not, by communicating the critical audit matter below, providing a separate opinion on the critical audit matter or on the accounts or disclosure to which it relates.

Description of the Matter

Uncertain Tax Positions

As discussed in Note 8 to the consolidated financial statements, the Company received three notices of proposed tax adjustments from the U.S. Internal Revenue Service (the "IRS") regarding the deductibility of interest paid on certain intercompany debt for the fiscal years 2016, 2017 and 2018. The IRS adjustments would result in a cumulative tax liability of approximately \$40 million. The Company believes it is more-likely-than-not that they will be able to sustain the interest deductions taken in the U.S. and has not recorded a liability for an uncertain tax position related to this matter.

Auditing management's analysis of tax positions related to interest paid on certain intercompany debt was challenging as the analysis is highly judgmental due to complex interpretations of tax laws and legal rulings. This tax position must be evaluated, and there may be uncertainties around initial recognition and de-recognition of tax positions, including regulatory changes, litigation and examination activity.

How We Addressed the Matter in Our Audit

We obtained an understanding, evaluated the design and tested the operating effectiveness of controls over the Company's accounting process for uncertain tax positions. For example, we tested controls over management's identification of uncertain tax positions and its application of the recognition and measurement principles, including management's review of the facts and circumstances and the corresponding tax laws relied upon to conclude that it is currently more-likely-than-not that they will realize the benefit recorded.

Our audit procedures included, among others, involving income tax professionals to assess the technical merits of the Company's tax positions related to certain intercompany debt and cross border transactions. We assessed the Company's correspondence with the relevant tax authorities and evaluated income tax opinions and other third-party advice obtained by the Company. We analyzed the Company's assumptions and data used to determine the amount of tax benefit to recognize and we tested the accuracy of the calculations performed. We also evaluated the adequacy of the Company's disclosures included in Note 8 to the consolidated financial statements in relation to these matters.

We have served as the Company's auditor since 1989.

/s/ Ernst & Young LLP Cleveland, Ohio May 29, 2020

STERIS PLC AND SUBSIDIARIES CONSOLIDATED BALANCE SHEETS (in thousands)

March 31,	2020			2019
Assets				
Current assets:				
Cash and cash equivalents	\$	319,581	\$	220,633
Accounts receivable (net of allowances of \$12,051 and \$9,645, respectively)		586,481		564,830
Inventories, net		248,259		208,243
Prepaid expenses and other current assets		54,430		60,029
Total current assets		1,208,751		1,053,735
Property, plant, and equipment, net		1,111,855		1,031,582
Lease right-of-use assets, net		131,837		_
Goodwill		2,356,085		2,322,928
Intangibles, net		565,473		604,614
Other assets		51,581		60,212
Total assets	\$	5,425,582	\$	5,073,071
Liabilities and equity				
Current liabilities:				
Accounts payable	\$	149,341	\$	152,913
Accrued income taxes		14,013		15,460
Accrued payroll and other related liabilities		128,261		109,058
Lease obligations due within one year		19,809		_
Accrued expenses and other		192,183		187,765
Total current liabilities		503,607		465,196
Long-term indebtedness		1,150,521		1,183,227
Deferred income taxes, net		160,270		151,038
Long-term lease obligations		114,114		_
Other liabilities		90,346		87,812
Total liabilities	\$	2,018,858	\$	1,887,273
Commitments and contingencies (see Note 10)				
Ordinary shares, with \$0.001 and \$75.00 par value, respectively; 500,000 shares authorized; 84,924 and 84,517 ordinary shares issued and outstanding, respectively		1,982,164		1,998,564
Retained earnings		1,647,175		1,339,024
Accumulated other comprehensive (loss)		(235,463)		(159,778)
Total shareholders' equity		3,393,876		3,177,810
Noncontrolling interests		12,848		7,988
Total equity		3,406,724		3,185,798
Total liabilities and equity	\$	5,425,582	\$	5,073,071

CONSOLIDATED STATEMENTS OF INCOME

(in thousands, except per share amounts)

Years Ended March 31,	2020	2019	2018
Revenues:			
Product	\$ 1,402,788	\$ 1,296,025	\$ 1,220,633
Service	 1,628,107	1,486,145	1,399,363
Total revenues	3,030,895	2,782,170	2,619,996
Cost of revenues:			
Product	750,202	702,295	646,177
Service	960,770	904,448	881,073
Total cost of revenues	1,710,972	1,606,743	1,527,250
Gross profit	1,319,923	1,175,427	1,092,746
Operating expenses:			
Selling, general, and administrative	716,731	669,937	631,978
Research and development	65,546	63,038	60,782
Restructuring expenses	673	30,987	103
Total operating expenses	782,950	763,962	692,863
Income from operations	536,973	411,465	399,883
Non-operating expenses, net:			
Interest expense	40,279	45,015	50,629
Interest income and miscellaneous expense	 (1,987)	(3,020)	(5,728)
Total non-operating expenses, net	38,292	41,995	44,901
Income before income tax expense	498,681	369,470	354,982
Income tax expense	90,876	64,394	63,360
Net income	407,805	305,076	291,622
Less: Net income attributable to noncontrolling interests	200	1,025	707
Net income attributable to shareholders	\$ 407,605	\$ 304,051	\$ 290,915
Net income per share attributable to shareholders:			
Basic	\$ 4.81	\$ 3.59	3.42
Diluted	\$ 4.76	\$ 3.56	3.39
Cash dividends declared per ordinary share outstanding	\$ 1.45	\$ 1.33	\$ 1.21

STERIS PLC AND SUBSIDIARIES CONSOLIDATED STATEMENTS OF COMPREHENSIVE INCOME (in thousands)

Years Ended March 31,		2020	2019	2018
Net income	\$	407,805	\$ 305,076	\$ 291,622
Less: Net income attributable to noncontrolling interests		200	1,025	707
Net income attributable to shareholders	\$	407,605	\$ 304,051	\$ 290,915
Other comprehensive (loss) income				
Unrealized gain on available for sale securities, (net of taxes of \$0, \$0 and \$516, respectively)		_	_	1,792
Pension and postretirement benefit plan changes (net of taxes of \$295, (\$423), and \$1,860, respectively)	(2,609)		2,538	(4,387)
Change in cumulative foreign currency translation adjustment		(73,076)	(172,031)	254,982
Total other comprehensive (loss) income attributable to shareholders		(75,685)	(169,493)	252,387
Comprehensive income attributable to shareholders	\$	331,920	\$ 134,558	\$ 543,302

STERIS PLC AND SUBSIDIARIES CONSOLIDATED STATEMENTS OF CASH FLOWS

(in thousands)

Years Ended March 31,	2020		2019	2018
Operating activities:				
Net income	\$ 407,80)5 \$	305,076	\$ 291,622
Adjustments to reconcile net income to net cash provided by operating activities:				
Depreciation, depletion, and amortization	197,2	35	225,921	178,332
Deferred income taxes	9,42	23	(6,511)	(24,722)
Share-based compensation expense	23,8	1	23,965	22,187
Loss (gain) on the disposal of property, plant, equipment, and intangibles, net	(1'	74)	924	2,582
Loss (gain) on sale of businesses	1,7	70	(1,370)	14,547
Other items	42	26	(18,397)	32,229
Changes in operating assets and liabilities, net of effects of acquisitions:				
Accounts receivable, net	(17,80	66)	(48,486)	(37,731)
Inventories, net	(39,0	57)	(14,617)	(5,178)
Other current assets	3,78	34	(7,371)	(1,244)
Accounts payable	(2,7'	79)	21,244	563
Accruals and other, net	6,19)1	59,127	(15,555)
Net cash provided by operating activities	590,5	59	539,505	457,632
Investing activities:				
Purchases of property, plant, equipment, and intangibles, net	(214,5)	(6)	(189,715)	(165,457)
Proceeds from the sale of property, plant, equipment, and intangibles	4,1	56	5,567	2,094
Proceeds from the sale of businesses	4,	39	2,478	8,888
Purchases of investments	-	_	(4,955)	_
Acquisition of business, net of cash acquired	(109,8	(4)	(13,313)	(46,271)
Other	-	_	(13,286)	(3,083)
Net cash used in investing activities	(319,7)	35)	(213,224)	(203,829)
Financing activities:				
Payments on long-term obligations	-	_	(85,000)	(222,500)
(Payments) proceeds under credit facilities, net	(26,5)) 0)	(27,087)	29,065
Deferred financing fees and debt issuance costs	(1,28	31)	(488)	(2,029)
Acquisition related deferred or contingent consideration	(62	26)	(1,327)	(2,064)
Repurchases of shares	(51,24	11)	(81,494)	(65,485)
Cash dividends paid to common shareholders	(123,0)	34)	(112,503)	(102,929)
Contributions from noncontrolling interest	6,0	50	_	_
Distributions to noncontrolling interest	(1,2)	15)	(255)	(1,400)
Stock option and other equity transactions, net	34,7	31	13,362	11,158
Net cash used in financing activities	(163,14	16)	(294,792)	(356,184)
Effect of exchange rate changes on cash and cash equivalents	(8,73	30)	(12,390)	20,997
Increase (decrease) in cash and cash equivalents	98,9	18	19,099	(81,384)
Cash and cash equivalents at beginning of period	220,63	33	201,534	282,918
Cash and cash equivalents at end of period	\$ 319,58	81 \$	220,633	\$ 201,534

CONSOLIDATED STATEMENTS OF SHAREHOLDERS' EQUITY

(in thousands, except per share amounts)

Equity compensation programs 592 32,470 — — — — — 32 Distributions to noncontrolling interest — — — — — — (1,400) (1 Cash dividends – \$1.21 per ordinary share — — — — (102,929) — — — (102 Change in noncontrolling interest — — — — — 602 Balance at March 31, 2018 84,747 \$ 2,048,037 100 \$ 15 \$ 1,146,223 \$ 11,685 \$ 11,340 \$ 3,217 Comprehensive income:	
Comprehensive income: Net income — — — — 290,915 — 707 291 Other comprehensive loss — — — — — 252,387 — 252 Repurchases of ordinary shares (793) (69,567) — — 4,082 — — — (65 Equity compensation programs 592 32,470 — — — — — 32 Distributions to noncontrolling interest — — — — — — — (1,400) (1 Cash dividends – \$1.21 per ordinary share — — — — — — — — 602 Change in noncontrolling interest — — — — — — 602 Balance at March 31, 2018 84,747 \$ 2,048,037 100 \$ 15 \$ 1,146,223 \$ 11,685 \$ 11,340 \$ 3,217	0.022
Net income),033
Other comprehensive loss — — — — — 252,387 — 252 Repurchases of ordinary shares (793) (69,567) — — 4,082 — — — (65 Equity compensation programs 592 32,470 — — — — — — 32 Distributions to noncontrolling interest — — — — — — (1,400) (1 Cash dividends – \$1.21 per ordinary share — — — — — — — — (102,929) — — — 602 Balance at March 31, 2018 84,747 \$ 2,048,037 100 \$ 15 \$ 1,146,223 \$ 11,685 \$ 11,340 \$ 3,217 Comprehensive income:	1 (22
Repurchases of ordinary shares (793) (69,567) — — 4,082 — — — (65,567) Equity compensation programs 592 32,470 — — — — — 32 Distributions to noncontrolling interest — — — — — — (1,400) (1 Cash dividends – \$1.21 per ordinary share — — — — — — — (102,929) — — — (102 Change in noncontrolling interest — — — — — — 602 Balance at March 31, 2018 84,747 \$ 2,048,037 100 \$ 15 \$ 1,146,223 \$ 11,685 \$ 11,340 \$ 3,217 Comprehensive income: — <	
Equity compensation programs 592 32,470 — — — — — 32 Distributions to noncontrolling interest — — — — — — (1,400) (1 Cash dividends – \$1.21 per ordinary share — — — — (102,929) — — — (102 Change in noncontrolling interest — — — — — 602 Balance at March 31, 2018 84,747 \$ 2,048,037 100 \$ 15 \$ 1,146,223 \$ 11,685 \$ 11,340 \$ 3,217 Comprehensive income:	
Distributions to noncontrolling interest — — — — — — — — — — — — — — — — — — —	5,485)
Cash dividends - \$1.21 per ordinary share - - - (102,929) - - - (102 Change in noncontrolling interest - - - - - - 602 Balance at March 31, 2018 84,747 \$ 2,048,037 100 \$ 15 \$1,146,223 \$ 11,685 \$ 11,340 \$ 3,217 Comprehensive income:	2,470
Change in noncontrolling interest — — — — — 602 Balance at March 31, 2018 84,747 \$ 2,048,037 100 \$ 15 \$1,146,223 \$ 11,685 \$ 11,340 \$ 3,217 Comprehensive income:	1,400)
Balance at March 31, 2018 84,747 \$ 2,048,037 100 \$ 15 \$1,146,223 \$ 11,685 \$ 11,340 \$ 3,217 Comprehensive income:	2,929)
Comprehensive income:	602
•	7,300
Net income — — — — 304,051 — 1,025 305	
	5,076
Other comprehensive loss — — — — — — — — — — — (169,493) — — (169	9,493)
Repurchases of ordinary shares (763) (86,414) — — 4,920 — — (81	1,494)
	6,941
Retirement of shares resulting from Redomiciliation (84,514) (10,592,117) (100) (15) — — — (10,592)	2,132)
Issuance of shares resulting from Redomiciliation 84,514 10,592,117 — — — — — 10,592	2,117
Adoption of Accounting Standards (note 1) — — — — (3,667) (1,970) — (5	5,637)
Cash dividends – \$1.33 per ordinary share – – – (112,503) – – (112	2,503)
Distributions to noncontrolling interest — — — — — — — (255)	(255)
Other changes in noncontrolling interest	4,122)
Balance at March 31, 2019 84,517 \$ 1,998,564 — \$ — \$1,339,024 \$ (159,778) \$ 7,988 \$ 3,185	5,798
Comprehensive income:	
Net income — — — 407,605 — 200 407	7,805
Other comprehensive income — — — — — — (75,685) — (75	5,685)
Repurchases of ordinary shares (396) (74,821) — — 23,580 — — (51	1,241)
Equity compensation programs and other 803 58,421 — — — — 58	8,421
Cash dividends – \$1.45 per ordinary share — — — — — — — — — — — — — — — — — — —	3,034)
Distributions to noncontrolling interest — — — — — — — — (1,245)	1,245)
Contributions from noncontrolling interest — — — — — — 6,050 6	6,050
Other changes in noncontrolling interest	(145)
Balance at March 31, 2020 84,924 \$ 1,982,164 — \$ — \$1,647,175 \$ (235,463) \$ 12,848 \$ 3,406	6,724

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

1. NATURE OF OPERATIONS AND SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

Nature of Operations. On March 28, 2019, STERIS plc, a public limited company organized under the laws of England and Wales ("STERIS UK"), completed a redomiciliation from the United Kingdom to Ireland (the "Redomiciliation"). The Redomiciliation was achieved through the insertion of a new Irish public limited holding company ("STERIS Ireland") on top of STERIS UK pursuant to a court-approved scheme of arrangement under English law (the "Scheme"). Following the Scheme effectiveness, STERIS UK was re-registered as a private limited company with the name STERIS Limited, and STERIS Emerald IE Limited, a company established in Ireland and a wholly-owned direct subsidiary of STERIS Ireland, was interposed as the direct parent company of STERIS UK.

STERIS plc is a leading provider of infection prevention and other procedural products and services. We offer our Customers a unique mix of innovative consumable products, such as detergents, gastrointestinal ("GI") endoscopy accessories, barrier product solutions, and other products and services, including: equipment installation and maintenance, microbial reduction of medical devices, instrument and scope repair solutions, laboratory testing services, on-site and off-site reprocessing, and capital equipment products, such as sterilizers and surgical tables, and connectivity solutions such as operating room ("OR") integration.

We operate and report in four reportable business segments: Healthcare Products, Healthcare Specialty Services, Life Sciences, and Applied Sterilization Technologies. We describe our business segments in Note 11 to our consolidated financial statements titled, "Business Segment Information."

Our fiscal year ends on March 31. References in this Annual Report to a particular "year," "fiscal," "fiscal year," or "year-end" mean our fiscal year. The significant accounting policies applied in preparing the accompanying consolidated financial statements of the Company are summarized below.

Principles of Consolidation. We use the consolidation method to report our investment in our subsidiaries. Therefore, the accompanying consolidated financial statements include the accounts of the Company and its wholly-owned and majority-owned subsidiaries. We eliminate inter-company accounts and transactions when we consolidate these accounts. Investments in equity of unconsolidated affiliates, over which the Company has significant influence, but not control, over the financial and operating polices, are accounted for primarily using the equity method. These investments are immaterial to the Company's Consolidated Financial Statements.

Use of Estimates. We make certain estimates and assumptions when preparing financial statements according to U.S. GAAP that affect the reported amounts of assets and liabilities at the financial statement dates and the reported amounts of revenues and expenses during the periods presented. These estimates and assumptions involve judgments with respect to many factors that are difficult to predict and are beyond our control. Actual results could be materially different from these estimates. We revise the estimates and assumptions as new information becomes available.

Cash Equivalents and Supplemental Cash Flow Information. Cash equivalents are all highly liquid investments with a maturity of three months or less when purchased. We invest our excess cash in short-term instruments including money market funds and time deposits with major banks and financial institutions. We select investments in accordance with the criteria established in our investment policy. Our investment policy specifies, among other things, maturity, credit quality and concentration restrictions with the objective of preserving capital and maintaining adequate liquidity.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Information supplementing our Consolidated Statements of Cash Flows is as follows:

Years Ended March 31,		2020	2019	2018
Cash paid during the year for:	,			_
Interest	\$	38,021	\$ 44,118	\$ 48,663
Income taxes		92,462	64,668	85,629
Cash received during the year for income tax refunds		4,378	2,189	7,747

Revenue Recognition and Associated Liabilities. We adopted Accounting Standards Update ("ASU") 2014-09 "Revenue from Contracts with Customers" and the subsequently issued amendments on April 1, 2018 using the modified retrospective approach to contracts that were not completed as of April 1, 2018. Under this standard, certain capital equipment contracts are comprised of a single performance obligation, resulting in the deferral of the corresponding capital equipment revenue and cost of revenues until installation is complete. Previously, these capital equipment revenues and cost of revenues were recognized based upon shipping terms. We recorded a cumulative effect adjustment in the beginning of fiscal 2019 to Retained earnings of \$5,637, based on the terms and conditions for certain open capital equipment contracts as of March 31, 2018.

Revenue is recognized when obligations under the terms of the contract are satisfied and control of the promised products or services have transferred to the Customer. Revenues are measured at the amount of consideration that we expect to be paid in exchange for the products or services. Product revenue is recognized when control passes to the Customer, which is generally based on contract or shipping terms. Service revenue is recognized when the Customer benefits from the service, which occurs either upon completion of the service or as it is provided to the Customer. Our Customers include end users as well as dealers and distributors who market and sell our products. Our revenue is not contingent upon resale by the dealer or distributor, and we have no further obligations related to bringing about resale. Our standard return and restocking fee policies are applied to sales of products. Shipping and handling costs charged to Customers are included in Product revenues. The associated expenses are treated as fulfillment costs and are included in Cost of revenues. Revenues are reported net of sales and value-added taxes collected from Customers.

We have individual Customer contracts that offer discounted pricing. Dealers and distributors may be offered sales incentives in the form of rebates. We reduce revenue for discounts and estimated returns, rebates, and other similar allowances in the same period the related revenues are recorded. The reduction in revenue for these items is estimated based on historical experience and trend analysis to the extent that it is probable that a significant reversal of revenue will not occur. Estimated returns are recorded gross on the Consolidated Balance Sheets.

In transactions that contain multiple performance obligations, such as when products, maintenance services, and other services are combined, we recognize revenue as each product is delivered or service is provided to the Customer. We allocate the total arrangement consideration to each performance obligation based on its relative standalone selling price, which is the price for the product or service when it is sold separately.

Payment terms vary by the type and location of the Customer and the products or services offered. Generally, the time between when revenue is recognized and when payment is due is not significant. We do not evaluate whether the selling price contains a financing component for contracts that have a duration of less than one year.

We do not capitalize sales commissions as substantially all of our sales commission programs have an amortization period of one year or less.

Certain costs to fulfill a contract are capitalized and amortized over the term of the contract if they are recoverable, directly related to a contract and generate resources that we will use to fulfill the contract in the future. At March 31, 2020, assets related to costs to fulfill a contract were not material to our Consolidated Financial Statements.

Refer to Note 11, titled "Business Segment Information" for disaggregation of revenue.

Product Revenue

Product revenues consist of revenues generated from sales of consumables and capital equipment. These contracts are primarily based on a Customer's purchase order and may include a Distributor, Dealer or Group Purchasing Organization (GPO) agreement. We recognize revenue for sales of product when control passes to the Customer, which generally occurs either when the products are shipped or when they are received by the Customer. Revenue related to certain capital equipment products is deferred until installation is complete as the capital equipment and installation are highly integrated and form a single performance obligation.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Service Revenue

Within our Healthcare Products and Life Sciences segments, service revenues consist of revenue generated from parts and labor associated with the maintenance, repair and installation of capital equipment. These contracts are primarily based on a Customer's purchase order and may include a Distributor, Dealer, or GPO agreement. For maintenance, repair and installation of capital equipment, revenue is recognized upon completion of the service.

We also offer preventive maintenance and separately priced extended warranty agreements to our Customers, which require us to maintain and repair our products over the duration of the contract. Generally, these contract terms are cancelable without penalty and range from one to five years. Amounts received under these Customer contracts are initially recorded as a service liability and are recognized as service revenue ratably over the contract term using a time-based input measure.

Within our Healthcare Specialty Services segment, revenues relate primarily to outsourced reprocessing services and instrument repairs. Contracts for outsourced reprocessing services are primarily based on an agreement with a Customer, ranging in length from several months to 15 years. Outsourced reprocessing services revenue is recognized ratably over the contract term using a time-based input measure, adjusted for volume and other performance metrics, to the extent that it is probable that a significant reversal of revenue will not occur. Contracts for instrument repairs are primarily based on a Customer's purchase order, and the associated revenue is recognized upon completion of the repair.

Within our Applied Sterilization Technologies segment, service revenues include contract sterilization and laboratory services. Sales contracts for contract sterilization and laboratory services are primarily based on a Customer's purchase order and associated Customer agreement and revenues are generally recognized upon completion of the service.

Contract Liabilities

Payments received from Customers are based on invoices or billing schedules as established in contracts with Customers. Deferred revenue is recorded when payment is received in advance of performance under the contract. Deferred revenue is recognized as revenue upon completion of the performance obligation, which generally occurs within one year. During fiscal 2020, we recognized revenue of \$48,602 that was included in our contract liability balance at the beginning of the period. During fiscal 2019, we recognized revenue of \$30,169 that was included in our contract liability balance at the beginning of the period.

Refer to Note 7, titled "Additional Consolidated Balance Sheet Information" for Deferred revenue balances.

Service Liabilities

Payments received in advance of performance for cancelable preventative maintenance and separately priced extended warranty contracts are recorded as service liabilities. Service liabilities are recognized as revenue as performance is rendered under the contract. Prior to the adoption of Accounting Standards Codification ("ASC") 606, these amounts were included in Deferred revenues.

Refer to Note 7, titled "Additional Consolidated Balance Sheet Information" for Service liability balances.

Remaining Performance Obligations

Remaining performance obligations reflect only the performance obligations related to agreements for which we have a firm commitment from a Customer to purchase, and exclude variable consideration related to unsatisfied performance obligations. With regard to products, these remaining performance obligations include capital equipment and consumable orders which have not shipped. With regard to service, these remaining performance obligations primarily include installation, certification, and outsourced reprocessing services. As of March 31, 2020, the transaction price allocated to remaining performance obligations was approximately \$940,000. We expect to recognize approximately 49% of the transaction price within one year and approximately 45% beyond one year. The remainder has yet to be scheduled for delivery.

Accounts Receivable. Accounts receivable are presented at their face amount, less allowances for sales returns and uncollectible accounts. Accounts receivable consist of amounts billed and currently due from Customers and amounts earned but unbilled. We generally obtain and perfect security interest in products sold in the United States when we have a concern with the Customer's risk profile.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

We maintain an allowance for uncollectible accounts receivable for estimated losses in the collection of amounts owed by Customers. We estimate the allowance based on analyzing a number of factors, including amounts written off historically, Customer payment practices, and general economic conditions. We also analyze significant Customer accounts on a regular basis and record a specific allowance when we become aware of a specific Customer's inability to pay. As a result, the related accounts receivable are reduced to an amount that we reasonably believe is collectible.

We maintain an allowance for sales returns based upon known returns and estimated returns for both capital equipment and consumables. We estimate returns of capital equipment and consumables based upon recent historical experience.

Inventories, net. Inventories are stated at the lower of their cost or market value. We determine cost based upon a combination of the last-in, first-out ("LIFO") and first-in, first-out ("FIFO") cost methods. For inventories valued using the LIFO method, we believe that the use of the LIFO method results in a matching of current costs and revenues. Inventories valued using the LIFO method represented approximately 25.3% and 25.2% of total inventories at March 31, 2020 and 2019, respectively. Inventory costs include material, labor, and overhead. If we had used only the FIFO method of inventory costing, inventories would have been \$16,937 and \$16,757 higher than those reported at March 31, 2020 and 2019, respectively.

We review inventory on an ongoing basis, considering factors such as deterioration, obsolescence, and other items. We record an allowance for estimated losses when the facts and circumstances indicate that particular inventories will not be usable. If future market conditions vary from those projected, and our estimates prove to be inaccurate, we may be required to write-down inventory values and record an adjustment to cost of revenues.

Property, Plant, and Equipment. Our property, plant, and equipment consists of land and land improvements, buildings and leasehold improvements, machinery and equipment, information systems, radioisotope (cobalt-60), and construction in progress. Property, plant, and equipment are presented at cost less accumulated depreciation and depletion. We capitalize additions and improvements. Repairs and maintenance are charged to expense as they are incurred.

Land is not depreciated and construction in progress is not depreciated until placed in service. Depreciation of most assets is computed on the cost less the estimated salvage value by using the straight-line method over the estimated remaining useful lives. Depletion of radioisotope is computed using the annual decay factor of the material, which is similar to the sum-of-the-years-digits method.

We generally depreciate or deplete property, plant, and equipment over the useful lives presented in the following table:

Asset Type	Useful Life
	(years)
Land improvements	3-40
Buildings and leasehold improvements	2-50
Machinery and equipment	2-20
Information Systems	2-20
Radioisotope (cobalt-60)	20

When we sell, retire, or dispose of property, plant, and equipment, we remove the asset's cost and accumulated depreciation from our Consolidated Balance Sheet. We recognize the net gain or loss on the sale or disposition in the Consolidated Statements of Income in the period when the transaction occurs.

Interest. We capitalize interest costs incurred during the construction of long-lived assets. We capitalized interest costs of \$428 and \$495 for the years ended March 31, 2020 and 2019, respectively. Total interest expense for the years ended March 31, 2020, 2019, and 2018 was \$40,279, \$45,015, and \$50,629, respectively.

Identifiable Intangible Assets. Our identifiable intangible assets include product technology rights, trademarks, licenses, and Customer and vendor relationships. We record these assets at cost, or when acquired as part of a business acquisition, at estimated fair value. We generally amortize identifiable intangible assets over periods ranging from 5 to 20 years using the straight-line method. Our intangible assets also include indefinite lived assets including certain trademarks and tradenames that were acquired in connection with business combinations. These assets are tested at least annually for impairment.

Investments. Investments in marketable securities are stated at fair value and are included in "Other assets" on the Consolidated Balance Sheets. Following the fiscal 2019 adoption of ASU 2016-01, "Financial Instruments - Overall - Recognition and Measurement of Financial Assets and Liabilities, changes in the fair value of these investments are recorded in the "Interest income and miscellaneous expense line" of the Consolidated Statement of Income.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Asset Impairment Losses. Property, plant, equipment, and identifiable intangible assets are reviewed for impairment when indicators of impairment exist and circumstances indicate that the carrying value of such assets may not be recoverable. Impaired assets are recorded at the lower of carrying value or estimated fair value. We monitor for such indicators on an ongoing basis and if an impairment exists, we record the loss in the Consolidated Statements of Income during that period.

Asset Retirement Obligations. We incur retirement obligations for certain assets. We record initial liabilities for the asset retirement obligations ("ARO") at fair value. Recognition of ARO includes: estimating the present value of a liability and offsetting asset, the subsequent accretion of that liability and depletion of the asset, and a periodic review of the ARO liability estimates and discount rates used in the analysis. We provide additional information about our asset retirement obligations in Note 5 to our consolidated financial statements titled, "Property, Plant and Equipment."

Acquisitions of Business. Assets acquired and liabilities assumed in a business combination are accounted for at fair value on the date of acquisition. Costs related to the acquisition are expensed as incurred.

Goodwill. We perform our annual impairment test for goodwill in the third quarter of each year. We may consider qualitative indicators of the fair value of a reporting unit when it is unlikely that a reporting unit has impaired goodwill. We may also utilize a discounted cash flow analysis that requires certain assumptions and estimates be made regarding market conditions and our future profitability. We review the book value compared to the fair value at the reporting unit level. We calculate the fair value of our reporting units based on the present value of estimated future cash flows. Management's judgment is necessary to evaluate the impact of operating and macroeconomic changes and to estimate future cash flows to measure fair value. Assumptions used in our impairment evaluations, such as forecasted growth rates and cost of capital, are consistent with internal projections, strategic plans, and operating plans. We believe such assumptions and estimates are also comparable to those that would be used by other market place participants.

Self-Insurance Liabilities. We record a liability for self-insured risks that we retain for general and product liabilities, workers' compensation, and automobile liabilities based on actuarial calculations. We use our historical loss experience and actuarial methods to calculate the liability. This liability includes estimates for both losses and incurred but not reported claims. We review the assumptions used to calculate the estimated liability at least annually to evaluate the adequacy of the amount recorded. We maintain insurance policies to cover losses greater than our estimated liability, which are subject to the terms and conditions of those policies. We are also self-insured for certain employee medical claims. We estimate a liability for incurred but not reported claims based upon recent claims experience.

Benefit Plans. We sponsor defined benefit pension plans. We also sponsor a post-retirement benefits plan for certain former employees. We determine our costs and obligations related to these plans by evaluating input from third-party professional advisers. These costs and obligations are affected by assumptions including the discount rate, expected long-term rate of return on plan assets, the annual rate of change in compensation for eligible employees, estimated changes in costs of healthcare benefits, and other factors. We review the assumptions used on an annual basis.

We recognize an asset for the overfunded status or a liability for the underfunded status of defined benefit pension and post-retirement benefits plans in our consolidated balance sheets. This amount is measured as the difference between the fair value of plan assets and the benefit obligation (the projected benefit obligation for pension plans and the accumulated post-retirement benefit obligation for other post-retirement benefit plans). Changes in the funded status of the plans are recorded in other comprehensive income in the year they occur. We measure plan assets and obligations as of the balance sheet date. We provide additional information about our pension and other post-retirement benefits plans in Note 9 to our consolidated financial statements titled, "Benefit Plans."

Fair Value of Financial Instruments. Except for long-term debt, our financial instruments are highly liquid or have short-term maturities. We provide additional information about the fair value of our financial instruments in Note 17 titled, "Fair Value Measurements."

Foreign Currency Translation. Most of our operations use their local currency as their functional currency. Financial statements of subsidiaries are translated into U.S. dollars using the exchange rate at each balance sheet date for assets and liabilities and a weighted average exchange rate for each period for revenues, expenses, gains and losses. Translation adjustments for subsidiaries whose local currency is their functional currency are recorded as a component of accumulated other comprehensive income (loss) within equity. Transaction gains and losses resulting from fluctuations in currency exchange rates on transactions denominated in currencies other than the functional currency are recognized as incurred in the accompanying Consolidated Statement of Income, except for certain inter-company balances designated as long-term in nature.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Forward and Swap Contracts. We enter into foreign currency forward contracts to hedge assets and liabilities denominated in foreign currencies, including inter-company transactions. We may also enter into commodity swap contracts to hedge price changes in nickel that impact raw materials included in our cost of revenues. We do not use derivative financial instruments for speculative purposes. These contracts are marked to market, with gains and losses recognized within "Selling, general, and administrative expenses" or "Cost of revenues" in the accompanying Consolidated Statements of Income.

Warranty. Warranties are provided on the sale of certain of our products and services and an accrual for estimated future claims is recorded at the time revenue is recognized. We estimate warranty expense based primarily on historical warranty claim experience.

Shipping and Handling. We record shipping and handling costs in costs of revenues. Shipping and handling costs charged to Customers are recorded as revenues in the period the product revenues are recognized.

Advertising Expenses. Costs incurred for communicating, advertising and promoting our products are generally expensed when incurred as a component of Selling, General and Administrative Expense. We incurred \$12,652, \$10,691, and \$10,886 of advertising costs during the years ended March 31, 2020, 2019, and 2018, respectively.

Research and Development. We incur research and development costs associated with commercial products and expense these costs as incurred. If a Customer reimburses us for research and development costs, the costs are charged to the related contracts as costs of revenues.

Income Taxes. We defer income taxes for all temporary differences between pre-tax financial and taxable income and between the book and tax basis of assets and liabilities. We record valuation allowances to reduce net deferred tax assets to an amount that we expect will more-likely-than-not be realized. In making such a determination, we consider all available information, including scheduled reversals of deferred tax liabilities, projected future taxable income, tax planning strategies, and if applicable, any carryback claims that can be filed. In the event we were to determine that we would be able to realize our deferred income tax assets in the future in excess of their net recorded amount, we would make an adjustment to the valuation allowance which would reduce the provision for income taxes and the effective tax rate.

We evaluate uncertain tax positions in accordance with a two-step process. The first step is recognition: The determination of whether or not it is more-likely-than-not that a tax position will be sustained upon examination, including resolution of any related appeals or litigation processes, based on the technical merits of the position. In evaluating whether a tax position has met the more-likely-than-not recognition threshold, we presume that the position will be examined by the appropriate tax authority and that the tax authority will have full knowledge of all relevant information. The second step is measurement: A tax position that meets the more-likely-than-not threshold is measured to determine the amount of benefit to recognize in the financial statements. The measurement process requires the determination of the range of possible settlement amounts and the probability of achieving each of the possible settlements. The tax position is measured at the largest amount of benefit that is greater than fifty percent likely of being realized upon ultimate settlement. No tax benefits are recognized for positions that do not meet the more-likely-than-not threshold. Tax positions that previously failed to meet the more-likely-than-not threshold are recognized in the first subsequent financial reporting period in which that threshold is met. Previously recognized tax positions that no longer meet the more-likely-than-not recognition threshold are derecognized in the first subsequent financial reporting period in which the threshold is no longer met. We describe income taxes further in Note 8 to our consolidated financial statements titled, "Income Taxes."

Share-Based Compensation. We describe share-based compensation in Note 14 to our consolidated financial statements titled, "Share-Based Compensation." We measure the cost of employee services received in exchange for an award of equity instruments based on the grant-date fair value of the award. We record liability awards at fair value each reporting period and the change in fair value is reflected as share-based compensation expense in our Consolidated Statements of Income. The expense is classified as cost of goods sold, selling, general and administrative expenses or research and development expenses in a manner consistent with the employee's compensation and benefits. These costs are recognized in the Consolidated Statement of Income over the period during which an employee is required to provide service in exchange for the award.

Restructuring. We recognize restructuring expenses as incurred. Asset impairment and accelerated depreciation expenses primarily relate to inventory write-downs for rationalized products and adjustments in the carrying value of the related facilities and machinery and equipment to their estimated fair value. In addition, the remaining useful lives of other property, plant, and equipment associated with the related operations are reevaluated based on the respective restructuring plan, which may result in the acceleration of depreciation and amortization of certain assets.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Recently Issued Accounting Standards Impacting the Company

Recently Issued Accounting Standards Impacting the Company are presented in the following table:

Standard	Date of Issuance	Description	Date of Adoption	Effect on the financial statements or other significant matters
Standards that hav	e recently beer	n adopted		
ASU 2016-02, "Leases" (Topic 842)	February 2016	The standard requires lessees to record all leases, whether finance or operating, on the balance sheet. An asset will be recorded to represent the right to use the leased asset, and a liability will be recorded to represent the lease obligation. The standard is effective for annual periods beginning after December 15, 2018 and interim periods within that period. Early adoption is permitted.	First Quarter Fiscal 2020	We adopted this standard, and related amendments, effective April 1, 2019 using the modified retrospective transition method and have not restated prior periods. We elected to use the package of practical expedients permitted under the transition guidance, which allows the carry forward of historical lease classification of existing leases. We also elected the practical expedient related to land easements, allowing us to carry forward our accounting treatment for land easements on existing or expired agreements. We made an accounting policy election to not recognize lease assets or liabilities for leases with a term of 12 months or less and elected to not separate non-lease components from lease components from lease components to which they relate for all asset classes. We recorded lease right-of-use assets and lease liabilities for operating leases totaling \$120,562. The adoption of the standard did not have a material impact to the Consolidated Statements of Income or Cash Flows. Additional information is disclosed in Note 10 under the heading "Leases".

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

ASU 2017-12 "Targeted Improvements to Accounting for Hedging Activities" (Topic 815)	August 2017	The standard provides targeted improvements to accounting for hedging activities by expanding an entity's ability to hedge nonfinancial and financial risk components and reduce complexity in fair value hedges of interest rate risk. The guidance eliminates the requirement to separately measure and report hedge ineffectiveness and generally requires the entire change in the fair value of a hedging instrument to be presented in the same income statement line as the hedged item. The guidance also eases certain documentation and assessment requirements and modifies the accounting for components excluded from the assessment of hedge effectiveness. The standard is effective for fiscal years, and interim periods within those years, beginning after December 15, 2018. Early adoption is permitted in any interim period after issuance of the standard.	First Quarter Fiscal 2020	We adopted this standard effective April 1, 2019 with no material impact to our Consolidated Balance Sheets. The impact to our Consolidated Statements of Income will depend on the value of future hedging activities.
ASU 2018-02 "Income Statement - Reporting Comprehensive Income" (Topic 220)	February 2018	The standard allows a reclassification from accumulated other comprehensive income to retained earnings for stranded tax effects resulting from the Tax Cuts and Jobs Act ("TCJA") and requires certain disclosures about stranded tax effects. The underlying guidance requiring that the effect of a change in tax laws or rates be included in income from continuing operations is not affected. This standard is effective for fiscal years beginning after December 15, 2018 and interim periods within those years. Early adoption is permitted.	First Quarter Fiscal 2020	We have elected not to reclassify the income tax effects of the TCJA from Accumulated Other Comprehensive Income ("AOCI") to retained earnings. Our policy is to release income tax effects from AOCI when individual units of account are sold or terminated.

ASU 2016-13, "Measurement of Credit Losses on Financial Instruments"	June 2016	The standard requires a financial asset (or group of financial assets) measured at amortized cost to be presented at the net amount expected to be collected. The allowance for credit losses is a valuation account that is deducted from the amortized cost basis of the financial asset(s) to present the net carrying value at the amount expected to be collected on the financial asset. Credit losses relating to available-for-sale debt securities should be recorded through an allowance for credit losses. The standard is effective for annual periods beginning after December 15, 2019. Early adoption is permitted.	N/A	We do not expect this standard to have a material impact on our consolidated financial statements.
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NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

ASU 2018-13 "Fair Value Measurement (Topic 820) Disclosure Framework- Changes to Disclosure Requirements for Fair Value Measurement"	August 2018	The standard modifies the disclosure requirements by adding, removing, and modifying certain required disclosures for fair value measurements for assets and liabilities disclosed within the fair value hierarchy. The standard is effective for fiscal years, and interim periods within those fiscal years, beginning after December 15, 2019 and early adoption is permitted.	N/A	We do not expect this standard to have a material impact on our consolidated financial statements as it modifies disclosure requirements only.
ASU 2018-14 "Compensation-Retirement Benefits - Defined Benefit Plans- General Topic (715-20): Disclosure Framework- Changes to the Disclosure Requirements for Defined Benefit Plans"	August 2018	The standard modifies the disclosure requirements by adding, removing, and modifying certain required disclosures for employers that sponsor defined benefit pension or other post-retirement benefit plans. The standard also clarifies disclosure requirements for defined benefit pension plans relating to the projected benefit obligation and accumulated benefit obligation. The standard is effective for fiscal years ending after December 15, 2019 and early adoption is permitted.	N/A	We do not expect this standard to have a material impact on our consolidated financial statements as it modifies disclosure requirements only.
ASU 2018-15 "Intangibles- Goodwill and Other- Internal Use Software (Subtopic 350-40): Customer's Accounting for Implementation Costs Incurred in a Cloud Computing Arrangement that is a Service Contract"	August 2018	The standard aligns the requirements for capitalizing implementation costs incurred in a hosting arrangement that is a service contract with the requirements for capitalizing implementation costs incurred to develop or obtain internal-use software. The standard is effective for fiscal years beginning after December 15, 2019 and early adoption is permitted.	N/A	We do not expect this standard to have a material impact on our consolidated financial statements.
ASU 2019-12 "Income Taxes (Topic 740)"	December 2019	The standard provides final guidance that simplifies the accounting for income taxes by eliminating certain exceptions to the guidance in ASC 740 related to the approach for intraperiod tax allocation, the methodology for calculating income taxes in an interim period and the recognition of deferred tax liabilities for outside basis differences. The guidance simplifies accounting for franchise taxes and enacted changes in tax laws or rates and clarifies the accounting for transactions that result in a step-up in the tax basis of goodwill. The standard is effective for fiscal years ending after December 15, 2020 and early adoption is permitted.	N/A	We are in the process of evaluating the impact that the standard will have on our consolidated financial statements.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

2. RESTRUCTURING

Fiscal 2019 Restructuring Plan. During the third quarter of fiscal 2019, we adopted and announced a targeted restructuring plan (the "Fiscal 2019 Restructuring Plan"), which included the closure of two manufacturing facilities, one in Brazil and one in England, as well as other actions including the rationalization of certain products. Fewer than 200 positions were eliminated. The Company has relocated the production of certain impacted products to other existing manufacturing operations during fiscal 2020. These restructuring actions were designed to enhance profitability and improve efficiency.

Since inception of the Fiscal 2019 Restructuring Plan we have incurred pre-tax expenses totaling \$43,851 related to these restructuring actions, of which \$31,660 was recorded as restructuring expenses and \$12,191 was recorded in cost of revenues, with a total of \$31,162, \$2,518, \$668, and \$7,798 related to the Healthcare Products, Healthcare Specialty Services, Life Sciences, and Applied Sterilization Technologies segments, respectively. Corporate related restructuring charges were \$1,705. Additional restructuring expenses related to this plan are not expected to be material to our results of operations.

The following table summarizes our total pre-tax restructuring expenses for fiscal 2020 and 2019:

Fiscal 2019 Restructuring Plan	Year Ended March 31, 2020	Year Ended March 31, 2019
Severance and other compensation related costs	\$ 1,554 \$	5,651
Accelerated depreciation and amortization	_	16,194
(Gain) on disposal of asset	(1,164)	
Asset impairment	_	4,312
Lease termination costs and other	283	4,830
Product rationalization (1)	2,470	9,721
Total restructuring expenses	\$ 3,143 \$	40,708

⁽¹⁾ Recorded in cost of revenues on the Consolidated Statements of Income.

Liabilities related to restructuring activities are recorded as current liabilities on the accompanying Consolidated Balance Sheets within "Accrued payroll and other related liabilities" and "Accrued expenses and other." The following tables summarize our restructuring liability balances:

Fiscal 2019 Restructuring Plan	ľ	March 31, 2019	Provisions	ayments / airments (1)	March 31, 2020
Severance and termination benefits	\$	4,102	\$ 1,554	\$ (4,659)	\$ 997
Lease termination obligations and other		2,029	283	(2,292)	20
Total	\$	6,131	\$ 1,837	\$ (6,951)	\$ 1,017

(1) Certain amounts reported include the impact of foreign currency movements relative to the U.S. dollar.

Fiscal 2019 Restructuring Plan	March 31, 2018	Provisions	Payments / pairments (1)	March 31, 2019
Severance and termination benefits	\$ 	\$ 5,651	\$ (1,549) \$	4,102
Lease termination obligations and other	_	4,830	(2,801)	2,029
Total	\$ 	\$ 10,481	\$ (4,350) \$	6,131

(1) Certain amounts reported include the impact of foreign currency movements relative to the U.S. dollar.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

3. GOODWILL AND INTANGIBLE ASSETS

Changes to the carrying amount of goodwill for the years ended March 31, 2020 and 2019 were as follows:

	Iealthcare Products Segment	Healthcare Specialty Services Segment	Life Sciences Segment		Applied Sterilization Technologies Segment	Total
Balance at March 31, 2018	404,674	388,025	148,816)	1,492,269	2,433,784
Goodwill acquired or allocated	(1,202)	(907)	_	-	5,341	3,232
Foreign currency translation adjustments	(6,188)	(12,208)	(1,02))	(94,671)	(114,088)
Balance at March 31, 2019	\$ 397,284	\$ 374,910	\$ 147,795	5 \$	5 1,402,939	\$ 2,322,928
Goodwill acquired or allocated	65,222	1,364	_	-	7,945	74,531
Divestitures	_	(199)	_	-	_	(199)
Foreign currency translation adjustments	(3,499)	(7,816)	762	2	(30,622)	(41,175)
Balance at March 31, 2020	\$ 459,007	\$ 368,259	\$ 148,557	′ §	1,380,262	\$ 2,356,085

See Note 18, titled "Business Acquisitions and Divestitures" for additional information regarding our recent business acquisitions and divestitures.

We evaluate the recoverability of recorded goodwill amounts annually during the third fiscal quarter, or when evidence of potential impairment exists. As a result of our annual impairment review of goodwill for fiscal years 2020, 2019 and 2018, no indicators of impairment were identified.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Information regarding our intangible assets is as follows:

	2020				20)19	
March 31,		Gross Carrying Amount		ccumulated mortization	Gross Carrying Amount		cumulated mortization
Customer relationships	\$	614,162	\$	227,581	\$ 623,774	\$	189,752
Non-compete agreements		4,646		4,012	4,693		3,945
Patents and technology		259,101		145,457	226,520		126,149
Trademarks and tradenames		62,543		39,942	63,570		38,850
Supplier relationships		54,800		12,787	54,800		10,047
Total	\$	995,252	\$	429,779	\$ 973,357	\$	368,743

Certain trademarks and tradenames obtained as a result of business combinations are indefinite-lived assets. The approximate carrying value of these assets at March 31, 2020 and March 31, 2019 was \$14,250 and \$13,000, respectively. We evaluate our indefinite-lived intangible assets annually during the third quarter, or when evidence of potential impairment exists. No impairment was recognized for fiscal year 2020. During the third quarter of fiscal 2019, management adopted a branding strategy that included phasing out the usage of a tradename associated with certain products in the Healthcare Products business segment. As a result, management recorded an impairment charge of \$16,249, which is included within the Selling, general, and administrative line of the Consolidated Statements of Income. The remaining fair value of the asset was calculated using an income approach (the relief from royalty method). The remaining fair value was not material and will be amortized over the asset's remaining useful life. Fair value calculated using this approach is classified within Level 3 of the fair value hierarchy and requires several assumptions.

Total amortization expense for intangible assets was \$74,528, \$98,747, and \$70,195 for the years ended March 31, 2020, 2019, and 2018, respectively. Based upon the current amount of intangible assets subject to amortization, the amortization expense for each of the five succeeding fiscal years is estimated to be as follows:

	2021	2022	2023	2024	2025
Estimated amortization expense	\$ 71 049	\$ 68 393	\$ 62.808	\$ 56 549	\$ 54 772

The estimated annual amortization expense presented in the preceding table has been calculated based upon March 31, 2020 currency exchange rates.

4. INVENTORIES, NET

Inventories, net consisted of the following:

March 31,	2020		2019
Raw materials	\$ 94,3	21 \$	83,009
Work in process	35,6	43	30,694
Finished goods	151,3	81	131,051
LIFO reserve	(16,9	37)	(16,757)
Reserve for excess and obsolete inventory	(16,1	49)	(19,754)
Inventories, net	\$ 248,2	59 \$	208,243

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(dollars in thousands, except per share amounts and as noted)

5. PROPERTY, PLANT AND EQUIPMENT

Information related to the major categories of our depreciable assets is as follows:

March 31,	2020	2019
Land and land improvements (1)	\$ 65,994	\$ 63,522
Buildings and leasehold improvements	531,267	480,359
Machinery and equipment	682,488	656,956
Information systems	181,112	169,711
Radioisotope	508,593	483,080
Construction in progress (1)	159,731	133,689
Total property, plant, and equipment	2,129,185	1,987,317
Less: accumulated depreciation and depletion	(1,017,330)	(955,735)
Property, plant, and equipment, net	\$ 1,111,855	\$ 1,031,582

⁽¹⁾ Land is not depreciated. Construction in progress is not depreciated until placed in service.

Depreciation and depletion expense were \$122,707, \$127,174 and \$108,137, for the years ended March 31, 2020, 2019, and 2018, respectively.

Asset Retirement Obligations

We have incurred asset retirement obligations (ARO) associated with the future disposal of these assets once depleted. Recognition of ARO includes: the present value of a liability and offsetting asset, the subsequent accretion of that liability and depletion of the asset, and the periodic review of the ARO liability estimates and discount rates used in the analysis.

The following table summarizes the activity in the liability for asset retirement obligations.

	Asset etirement oligations
Balance at March 31, 2018	\$ 11,639
Liabilities incurred during the period	1,033
Accretion expense and change in estimate	385
Foreign currency and other	(671)
Balance at March 31, 2019	\$ 12,386
Liabilities incurred during the period	94
Liabilities settled during the period	(168)
Accretion expense and change in estimate	453
Foreign currency and other	(251)
Balance at March 31, 2020	\$ 12,514

6. DEBT

Indebtedness as of March 31, 2020 and 2019 was as follows:

	2020	2019
Credit Agreement	\$ 275,449	\$ 301,846
Private Placement	878,409	884,967
Deferred financing fees	(3,337)	(3,619)
Other	_	33
Total long term debt	\$ 1,150,521	\$ 1,183,227

On March 23, 2018, STERIS UK and certain of its subsidiaries entered into a Credit Agreement (the "Credit Agreement") with various financial institutions as lenders, and JPMorgan Chase Bank, N.A., as Administrative Agent. STERIS Ireland

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

subsequently became a borrower and guarantor under the Credit Agreement. The Credit Agreement replaced a bank credit facility dated March 31, 2015. The Credit Agreement provides up to \$1,000,000 of credit, in the form of a revolver facility, which may be utilized for revolving credit borrowings, swing line borrowings and letters of credit, with sublimits for swing line borrowings and letters of credit. The revolver facility may be increased in specified circumstances by up to \$500,000. The Credit Agreement will mature on March 23, 2023, and all unpaid borrowings, together with accrued and unpaid interest thereon, are repayable on that date. The Credit Agreement contains leverage and interest coverage covenants. Borrowings may be taken in U.S. dollars, euros, and pounds sterling and certain other specified currencies and bear interest at our option based upon either the Base Rate or the Eurocurrency Rate, plus the Applicable Margin in effect from time to time under the Credit Agreement. The Applicable Margin is determined based on the ratio of Consolidated Total Debt to Consolidated EBITDA (as such terms are defined in the Credit Agreement). Interest on Base Rate Advances is payable quarterly in arrears and interest on Eurocurrency Rate Advances is payable at the end of the relevant interest period therefor, but in no event less frequently than every three months. Borrowings at closing were used to repay outstanding balances of debt outstanding under the former bank credit facility dated March 31, 2015 that was scheduled to mature on March 31, 2020 and for other general corporate purposes. The Credit Agreement was amended in March 2019, in connection with the Redomiciliation to permit the Redomiciliation. The amendments did not effect any material changes in the terms of the Credit Agreement regarding borrowings or the issuance of letters of credit.

As of March 31, 2020 a total of \$275,449 of Credit Agreement and Swing Line Facility borrowings were outstanding under the Credit Agreement, based on currency exchange rates as of March 31, 2020.

Our outstanding Senior Notes at March 31, 2020 and 2019 were as follows:

	Applicable Note Purchase Agreement	Maturity Date	U.S. Dollar Value at March 31, 2020	U.S. Dollar Value at March 31, 2019
\$35,000 Senior notes at 6.43%	2008 Private Placement	August 2020	35,000	35,000
\$91,000 Senior notes at 3.20%	2012 Private Placement	December 2022	91,000	91,000
\$80,000 Senior notes at 3.35%	2012 Private Placement	December 2024	80,000	80,000
\$25,000 Senior notes at 3.55%	2012 Private Placement	December 2027	25,000	25,000
\$125,000 Senior notes at 3.45%	2015 Private Placement	May 2025	125,000	125,000
\$125,000 Senior notes at 3.55%	2015 Private Placement	May 2027	125,000	125,000
\$100,000 Senior notes at 3.70%	2015 Private Placement	May 2030	100,000	100,000
\$50,000 Senior notes at 3.93%	2017 Private Placement	February 2027	50,000	50,000
€60,000 Senior notes at 1.86%	2017 Private Placement	February 2027	66,342	67,352
\$45,000 Senior notes at 4.03%	2017 Private Placement	February 2029	45,000	45,000
€20,000 Senior notes at 2.04%	2017 Private Placement	February 2029	22,114	22,450
£45,000 Senior notes at 3.04%	2017 Private Placement	February 2029	55,767	58,702
€19,000 Senior notes at 2.30%	2017 Private Placement	February 2032	21,008	21,328
£30,000 Senior notes at 3.17%	2017 Private Placement	February 2032	37,178	39,135
Total Senior Notes			\$ 878,409	\$ 884,967

On February 27, 2017, STERIS UK issued and sold an aggregate principal amount of \$95,000, €99,000, and £75,000, of senior notes in a private placement to certain institutional investors in an offering that was exempt from the registration requirements of the Securities Act of 1933. These notes have maturities of between 10 years and 15 years from the issue date. The agreement governing these notes contains leverage and interest coverage covenants.

On May 15, 2015, STERIS Corporation issued and sold \$350,000 of senior notes, in a private placement to certain institutional investors in an offering that was exempt from the registration requirements of the Securities Act of 1933. These notes have maturities of 10 years to 15 years from the issue date. The agreement governing these notes contains leverage and interest coverage covenants.

The agreements governing certain senior notes issued and sold in February 2013, December 2012, and August 2008, were amended and restated in their entirety on March 31, 2015. All of these notes were issued and sold in private placements to

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

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certain institutional investors in offerings that were exempt from the registration requirements of the Securities Act of 1933. The amended and restated agreements, which have been consolidated into a single agreement for the 2013 and 2012 notes, and a separate single agreement for the 2008 notes, contain leverage and interest coverage covenants.

All of the note agreements were amended in March 2019, in connection with the Redomiciliation. The amendments waived certain repurchase rights of the note holders and increased the size of certain baskets to more closely align with Credit Agreement baskets.

At March 31, 2020, we were in compliance with all financial covenants associated with our indebtedness.

The combined annual aggregate amount of maturities of our outstanding debt by fiscal year is as follows:

2021 (1)	\$ 35,000
2022	_
2023	366,449
2024	_
2025 and thereafter	752,409
Total	\$ 1,153,858

⁽¹⁾ This amount represents a senior note that matures in August 2020. In accordance with ASU 470-10-45, we have presented the note as a long-term liability based on our intention to refinance the note on a long-term basis under our credit facility.

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(dollars in thousands, except per share amounts and as noted)

7. ADDITIONAL CONSOLIDATED BALANCE SHEET INFORMATION

Additional information related to our Consolidated Balance Sheet is as follows:

arch 31,	2020	2019
Accrued payroll and other related liabilities:		
Compensation and related items	\$ 42,205	\$ 37,25
Accrued vacation/paid time off	9,917	10,19
Accrued bonuses	53,041	40,19
Accrued employee commissions	19,298	17,85
Other post-retirement benefits obligations-current portion	1,488	1,63
Other employee benefit plans' obligations-current portion	2,312	1,93
Total accrued payroll and other related liabilities	\$ 128,261	\$ 109,05
Accrued expenses and other:		
Deferred revenues	\$ 53,299	\$ 55,33
Service liabilities	47,505	42,10
Self-insured and related risk reserves-current portion	7,342	6,53
Accrued dealer commissions	15,827	15,28
Accrued warranty	7,381	7,19
Asset retirement obligation-current portion	2,671	2,65
Other	58,158	58,66
Total accrued expenses and other	\$ 192,183	\$ 187,70
Other liabilities:		
Self-insured risk reserves-long-term portion	\$ 17,452	\$ 14,44
Other post-retirement benefits obligations-long-term portion	9,880	10,9
Defined benefit pension plans obligations-long-term portion	10,987	16,16
Other employee benefit plans obligations-long-term portion	2,333	4,71
Accrued long-term income taxes	11,959	13,5
Asset retirement obligation-long-term portion	9,843	9,73
Contingent consideration obligations- long term portion	15,358	5,95
Other	12,534	12,37
Total other liabilities	\$ 90,346	\$ 87,81

8. INCOME TAXES

The Tax Cuts and Jobs Act (the "TCJA") was enacted on December 22, 2017. The TCJA reduced the maximum U.S. federal corporate income tax rate to 21.0%, required companies to pay a one-time transition tax on earnings of certain foreign subsidiaries that were previously tax deferred and created new taxes on certain foreign sourced earnings. The Company applied the guidance in Staff Accounting Bulletin No. 118, Income Tax Accounting Implications of the Tax Cut and Jobs Act when accounting for the enactment-date effects of the TCJA.

We consider the tax expense recorded for the TCJA to be complete at this time. However, it is possible that additional legislation, regulations, interpretations and/or guidance may be issued in the future that may result in additional adjustments to the tax expense recorded related to the TCJA. We have continued to monitor these as they are published. While none have resulted in material adverse impacts through fiscal 2020, there are certain items, which were not yet considered law as of March 31, 2020, that if finalized as proposed, could result in an adverse impact on our consolidated financial statements. Specifically, full retroactive application to April 1, 2019 of certain of the regulations relating to §267A, would require recognition of income tax expense up to \$15,000 related to the period April 1, 2019 through December 4, 2019 when we restructured certain of our intercompany financing arrangements. This potential impact contains significant uncertainty and

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

could be impacted by various factors, including any differences between the proposed and finalized regulations, issued in April 2020, and their retroactive application.

Income from continuing operations before income taxes was as follows:

Years Ended March 31,	2020	2019	2018
United States operations	\$ 325,522	\$ 235,405	\$ 203,872
Ireland operations	29,543	13,693	11,837
Other locations operations	143,616	120,372	139,273
	\$ 498,681	\$ 369,470	\$ 354,982

The components of the provision for income taxes related to income from continuing operations consisted of the following:

Years Ended March 31,	2020	2019	2018
Current:			
United States federal	\$ 42,032 \$	29,943 \$	47,728
United States state and local	9,971	12,484	7,727
Ireland	5,036	2,627	2,596
Other locations	24,600	26,824	26,742
	 81,639	71,878	84,793
Deferred:			
United States federal	10,073	5,775	(15,728)
United States state and local	2,363	2,836	2,656
Ireland	(899)	(546)	(280)
Other locations	(2,300)	(15,549)	(8,081)
	 9,237	(7,484)	(21,433)
Total Provision for Income Taxes	\$ 90,876 \$	64,394 \$	63,360

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The total provision for income taxes can be reconciled to the tax computed at the Ireland statutory tax rate for 2020 and 2019, and the United Kingdom statutory rate for 2018 as follows:

Years Ended March 31,	2020	2019	2018
National statutory tax rate	12.5 %	12.5 %	19.0 %
Increase (decrease) in accruals for uncertain tax positions	(0.3)%	— %	0.1 %
U.S. state and local taxes, net of federal income tax benefit	2.0 %	3.1 %	2.3 %
Increase in valuation allowances	0.5 %	0.4 %	0.1 %
U.S. research and development credit	(0.5)%	(0.6)%	(0.5)%
U.S. foreign income tax credit	(0.6)%	(0.2)%	(0.2)%
Difference in non-Ireland tax rates	6.9 %	4.5 %	%
Difference in non-United Kingdom tax rates	— %	— %	4.1 %
U.S. manufacturing deduction	— %	— %	(0.8)%
Excess tax benefit for equity compensation	(2.8)%	(2.2)%	(1.8)%
Tax rate changes on deferred tax assets and liabilities	0.1 %	(0.6)%	(10.3)%
U.S. transition tax on foreign earnings	%	(0.3)%	4.9 %
U.S. tax reform impact, GILTI and FDII	0.1 %	0.3 %	%
Acquisitions and divestitures	%	— %	0.5 %
Capitalized acquisition, redomiciliation costs	0.1 %	0.5 %	%
All other, net	0.2 %	— %	0.4 %
Total Provision for Income Taxes	18.2 %	17.4 %	17.8 %

Unrecognized Tax Benefits. We classify uncertain tax positions and related interest and penalties as long-term liabilities within "Other liabilities" in our accompanying Consolidated Balance Sheets, unless they are expected to be paid within 12 months, in which case, the uncertain tax positions would be classified as current liabilities within "Accrued income taxes." We recognize interest and penalties related to unrecognized tax benefits within "Income tax expense" in our accompanying Consolidated Statements of Income.

A reconciliation of the beginning and ending balances of the total amounts of unrecognized tax benefits is as follows:

	2020		2019
Unrecognized Tax Benefits Balance at April 1	\$ 2,3	14 \$	2,500
Increases for tax provisions of current year	1	76	178
Decreases for tax provisions of prior year	(1,5	70)	(186)
Other, including currency translation	(45)	(178)
Unrecognized Tax Benefits Balance at March 31	\$ 8	75 \$	2,314

We recognized interest and penalties related to uncertain tax positions in the provision for income taxes. As of March 31, 2020, and 2019 we had \$243 and \$360 accrued for interest and penalties, respectively. If all unrecognized tax benefits were recognized, the net impact on the provision for income tax expense would be \$1,118. The decrease in unrecognized tax benefits from prior year is due to the release of expired positions. It is reasonably possible that during the next 12 months, there will be no material reductions in unrecognized tax benefits as a result of the expiration of various statutes of limitations or other matters.

We operate in numerous taxing jurisdictions and are subject to regular examinations by various United States federal, state and local, as well as foreign jurisdictions. We are no longer subject to United States federal examinations for years before fiscal 2016 and, with limited exceptions, we are no longer subject to United States state and local, or non-United States, income tax examinations by tax authorities for years before fiscal 2015. We remain subject to tax authority audits in various jurisdictions wherever we do business.

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In May 2019, we received two notices of proposed tax adjustment from the U.S. Internal Revenue Service (the "IRS") regarding the deductibility of interest paid on certain intercompany debt. The notices relate to fiscal years 2016 and 2017. In September 2019, we received another notice of proposed adjustment for the same issue, for the 2018 fiscal year. The IRS adjustments would result in a cumulative tax liability of approximately \$40,000. Notices have not been received for subsequent periods. We are contesting the IRS's assertions, and intend to pursue available remedies such as appeals and litigation, if necessary. We have not established reserves related to these notices. An unfavorable outcome is not expected to have a material adverse impact on our consolidated financial position but could be material to our consolidated results of operations and cash flows for any one period.

We estimate that the tax benefit from our Costa Rican Tax Holiday is \$1,900 (or \$0.02 per fully diluted share), annually. The Tax Holiday runs fully exempt, from income tax, through 2025 and partially exempt through 2029.

Deferred Taxes. The significant components of the deferred tax assets and liabilities recorded in our accompanying balance sheets at March 31, 2020 and 2019 were as follows:

arch 31,	2020	2019
Deferred Tax Assets:		
Post-retirement benefit accrual	\$ 2,871	\$ 3,14
Compensation	12,560	14,27
Net operating loss carryforwards	16,149	19,19
Accrued expenses	5,490	4,85
Insurance	3,620	3,18
Deferred income	11,316	7,50
Bad debt	1,820	1,38
Pension	2,273	3,36
Operating leases (1)	28,945	-
Other	6,024	7,70
Deferred Tax Assets	 91,068	64,62
Less: Valuation allowance	13,891	13,47
Total Deferred Tax Assets	 77,177	51,14
Deferred Tax Liabilities:		
Depreciation and depletion	68,179	61,06
Operating leases (1)	29,268	-
Intangibles	129,951	128,47
Other	2,078	2,19
Total Deferred Tax Liabilities	 229,476	191,73
Net Deferred Tax Assets (Liabilities)	\$ (152,299)	\$ (140,59

⁽¹⁾ For more information regarding our operating leases, see Note 10 titled, "Commitments and Contingencies".

At March 31, 2020, we had U.S. federal operating loss carryforwards of \$10,942, which remain subject to a 20 year carryforward period. Additionally, we had non-U.S. operating loss carry forwards of \$41,450. Although the majority of the non-U.S. carryforwards have indefinite expiration periods, those carryforwards that have definite expiration periods will expire if unused between fiscal years 2021 and 2041. In addition, we have recorded pre-valuation allowance tax benefits of \$2,042 related to state operating loss carryforwards. If unused, these state operating loss carryforwards will expire between fiscal years 2021 and 2040. At March 31, 2020, we had \$2,547 of tax credit carryforwards. These credit carryforwards can be used through fiscal 2030.

We review the need for a valuation allowance against our deferred tax assets. A valuation allowance of \$13,891 has been applied to a portion of the net deferred tax assets because we do not believe it is more-likely-than-not that we will receive future benefit. The valuation allowance increased during fiscal 2020 by \$413.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Other than the tax expense recorded for the one-time transition tax on unremitted earnings of non-US subsidiaries, no additional provision has been made for income taxes on undistributed earnings of foreign subsidiaries as the Company's position is that these amounts continue to be indefinitely reinvested. The amount of undistributed earnings of subsidiaries was approximately \$1,600,000 at March 31, 2020. It is not practicable to estimate the additional income taxes and applicable withholding taxes that would be payable on the remittance of such undistributed earnings.

In October 2015, the Organization for Economic Cooperation and Development (OECD), in conjunction with the G20, finalized broad-based international tax policy guidelines that involve transfer pricing and other international tax subjects. While some member jurisdictions automatically adopt the new OECD guidelines, most member countries can adopt the guidelines only by new law or regulations. We are currently adopting processes to comply with the reporting requirements specified by the guidelines and are evaluating the other parts of the guidelines.

9. BENEFIT PLANS

In the United States, we sponsor an unfunded post-retirement welfare benefits plan for two groups of United States retirees. Benefits under this plan include retiree life insurance and retiree medical insurance, including prescription drug coverage.

During the second quarter of fiscal 2009, we amended our United States post-retirement welfare benefits plan, reducing the benefits to be provided to retirees under the plan and increasing their share of the costs. The amendments resulted in a decrease of \$46,001 in the accumulated post-retirement benefit obligation. The impact of this change was recognized in our Consolidated Balance Sheets in fiscal 2009 and is being amortized as a component of the annual net periodic benefit cost over a period of approximately thirteen years.

We sponsor several defined benefit pension schemes outside the United States: two in the UK, one in the Netherlands, two in Germany, and one in Switzerland. The Synergy Health plc Retirement Benefit Scheme is a defined benefit (final salary) funded pension scheme. In previous years, Synergy sponsored a funded defined benefit arrangement in the Netherlands. This was a separate fund holding the pension scheme assets to meet long-term pension liabilities for past and present employees. Accrual of benefits ceased under the scheme effective January 1, 2013. The Synergy Radeberg and Synergy Allershausen Schemes are unfunded defined pension schemes and are closed to new entrants. The Synergy Daniken Scheme is a defined benefit funded pension scheme. As a result of our fiscal 2018 acquisition of Harwell Dosimeters Ltd, we also sponsor in the Harwell Dosimeters Ltd Retirement Benefits Scheme which is a defined benefit funded pension scheme.

We recognize the funded status of our defined benefit pension and post-retirement benefit plans in our Consolidated Balance Sheets, with a corresponding adjustment to accumulated other comprehensive income, net of tax. The funded status is measured as of March 31 each year and is calculated as the difference between the fair value of plan assets and the benefit obligation (which is the projected benefit obligation for pension plans and the accumulated post-retirement benefit obligation for post-retirement benefit plans). Accumulated comprehensive income (loss) represents the net unrecognized actuarial losses and unrecognized prior service cost. These amounts will be recognized in net periodic benefit cost as they are amortized. We will recognize future changes to the funded status of these plans in the year the change occurs, through other comprehensive income.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Obligations and Funded Status. The following table reconciles the funded status of the defined benefit pension plans and the other post-retirement benefits plan to the amounts recorded on our Consolidated Balance Sheets at March 31, 2020 and 2019, respectively. Benefit obligation balances presented in the following table reflect the projected benefit obligations for our defined benefit pension plans and the accumulated other post-retirement benefit obligation for our post-retirement benefits plan. The measurement date of our defined benefit pension plans and other post-retirement benefits plan is March 31, for both periods presented.

	Other Defin Pension		Otl Post-Re Benefi	tirement		
	2020	2019	2020	2019		
Change in Benefit Obligations:						
Benefit Obligations at Beginning of Year	\$ 133,672	\$ 148,848	\$ 12,551	\$ 14,100		
Service cost	1,380	2,394	_	_		
Prior service cost	_	831	_	_		
Interest cost	2,955	3,255	408	457		
Actuarial loss (gain)	(3,736)	(4,402)	181	(106)		
Benefits and expenses	(6,466)	(6,150)	(1,772)	(1,900)		
Employee contributions	1,046	743	_	_		
Impact of foreign currency exchange rate changes	(5,661)	(11,847)	_	_		
Benefit Obligations at End of Year	123,190	133,672	11,368	12,551		
Change in Plan Assets:						
Fair Value of Plan Assets at Beginning of Year	117,504	119,441	_	_		
Actual return on plan assets	228	6,543	_	_		
Employer contributions	5,071	5,005	1,772	1,900		
Employee contributions	1,045	742	_	_		
Benefits and expenses paid	(6,466)	(6,150)	(1,772)	(1,900)		
Impact of foreign currency exchange rate changes	(5,179)	(8,077)	_	_		
Fair Value of Plan Assets at End of Year	112,203	117,504				
Funded Status of the Plans	\$ (10,987)	\$ (16,168)	\$ (11,368)	\$ (12,551)		

Amounts recognized in the consolidated balance sheets consist of the following:

	Other Defined Benefit Pension Plans 2020 2019			C	Other Post- Benefit			
				2019	2020			2019
Current liabilities	\$	_	\$	_	\$	(1,488)	\$	(1,633)
Noncurrent liabilities		(10,987)		(16,168)		(9,880)		(10,918)
	\$	(10,987)	\$	(16,168)	\$	(11,368)	\$	(12,551)

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

The pre-tax amount of unrecognized actuarial net loss and unamortized prior service cost included in accumulated other comprehensive (loss) income at March 31, 2020, was approximately \$14,405 and \$7,463, respectively. During fiscal 2021, we will amortize the following pre-tax amounts from accumulated other comprehensive income:

	Defined Benefit Pension Plans	Other Post- Retirement Benefits Plan		
Actuarial loss	\$ 20	\$ 482		
Prior Service Cost	69	(3,263)		

Defined benefit plans with an accumulated benefit obligation and projected benefit obligation exceeding the fair value of plan assets had the following plan assets and obligations at March 31, 2020 and 2019:

	Pension Plans
	2020 2019
Aggregate fair value of plan assets	\$ 112,203 \$ 117,504
Aggregate accumulated benefit obligations	120,084 130,669
Aggregate projected benefit obligations	123,190 132,672

Components of Net Periodic Benefit Cost and Other Amounts Recognized in Other Comprehensive

Income. Components of the annual net periodic benefit cost of our defined benefit pension plans and our other post-retirement benefits plan were as follows:

	Other Defined Benefit Pension Plans					Other Pos	t-Re	etirement Ber	nefit	s Plan	
		2020		2019		2018	2020		2019		2018
Service cost	\$	1,380	\$	2,394	\$	2,402	\$ 	\$	_	\$	
Interest cost		2,876		3,139		3,262	409		457		519
Expected return on plan assets		(4,735)		(4,930)		(4,835)	_		_		_
Prior service cost recognition		69		51		_	(3,263)		(3,263)		(3,263)
Net amortization and deferral		9		474		126	482		552		648
Net periodic benefit (credit) cost	\$	(401)	\$	1,128	\$	955	\$ (2,372)	\$	(2,254)	\$	(2,096)
Recognized in other comprehensive loss (income) before tax:											
Net loss (gain) occurring during year	\$	890	\$	(6,545)	\$	(697)	\$ (181)	\$	106	\$	501
Amortization of prior service credit		(78)		781		_	3,263		3,263		3,263
Amortization of net loss		_		(468)		(126)	(482)		(552)		(648)
Total recognized in other comprehensive loss (income)		812		(6,232)		(823)	2,600		2,817		3,116
Total recognized in total benefits cost and other comprehensive loss (income)	\$	411	\$	(5,104)	\$	132	\$ 228	\$	563	\$	1,020

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Assumptions Used in Calculating Benefit Obligations and Net Periodic Benefit Cost. The following table presents significant assumptions used to determine the projected benefit obligations at March 31:

	2020	2019
Discount Rate:		
Synergy Health plc Retirement Benefits Scheme	2.40%	2.50%
Isotron BV Pension Plan	1.60%	1.20%
Synergy Health Daniken AG	0.20%	0.85%
Synergy Health Radeberg	1.60%	1.60%
Synergy Health Allershausen	0.50%	1.60%
Harwell Dosimeters Ltd Retirement Benefits Scheme	2.45%	2.35%
Other post-retirement plan	3.00%	3.50%

The following table presents significant assumptions used to determine the net periodic benefit costs for the years ended March 31:

	2020	2019	2018
Discount Rate:			
Synergy Health plc Retirement Benefits Scheme	2.50%	2.50%	2.60%
Isotron BV Pension Plan	1.20%	1.60%	1.60%
Synergy Health Daniken AG	0.20%	0.95%	0.65%
Synergy Health Radeberg	1.60%	1.60%	1.50%
Synergy Health Allershausen	1.75%	1.60%	1.50%
Harwell Dosimeters Ltd Retirement Benefits Scheme	2.45%	2.55%	2.55%
Other post-retirement plan	3.50%	3.50%	3.50%
Expected Return on Plan Assets:			
Synergy Health plc Retirement Benefits Scheme	4.80%	5.02%	4.97%
Isotron BV Pension Plan	1.20%	1.60%	1.60%
Synergy Health Daniken AG	0.65%	1.20%	1.40%

The net periodic benefit cost and the actuarial present value of projected benefit obligations are based upon assumptions that we review on an annual basis. These assumptions may be revised annually based upon an evaluation of long-term trends, as well as market conditions that may have an impact on the cost of providing benefits.

We develop our expected long-term rate of return on plan assets assumptions by evaluating input from third-party professional advisers, taking into consideration the asset allocation of the portfolios and the long-term asset class return expectations.

We develop our discount rate assumptions by evaluating input from third-party professional advisers, taking into consideration the current yield on country specific investment grade long-term bonds which provide for similar cash flow streams as our projected obligations.

We have made assumptions regarding healthcare costs in computing our other post-retirement benefit obligation. The assumed rates of increase generally decline ratably over a five-year period from the assumed current year healthcare cost trend rate to the assumed long-term healthcare cost trend rate noted below.

	2020	2019	2018
Healthcare cost trend rate – medical	6.75%	6.75%	7.00%
Healthcare cost trend rate – prescription drug	6.75%	6.75%	7.00%
Long-term healthcare cost trend rate	4.50%	4.50%	4.50%

To determine the healthcare cost trend rates, we evaluate a combination of information, including ongoing claims cost monitoring, annual statistical analyses of claims data, reconciliation of forecasted claims against actual claims, review of trend

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

assumptions of other plan sponsors and national health trends, and adjustments for plan design changes, workforce changes, and changes in plan participant behavior.

A one-percentage-point change in assumed healthcare cost trend rates (including medical, prescription drug, and long-term rates) would have had the following effect on our other post-retirement benefit obligation at March 31, 2020:

		Increase		entage Point		
	-			Deci	rease	
Effect on total service and interest cost components		\$		\$		
Effect on other post-retirement benefit obligation			7		(6)	

Plan Assets. The investment policies for our plans are generally established by the local pension plan trustees and seek to maintain the plans' ability to meet liabilities and to comply with local minimum funding requirements. Plan assets are invested in diversified portfolios that provide adequate levels of return at an acceptable level of risk. The investment policies are reviewed at least annually and revised, as deemed appropriate to ensure that the objectives are being met. At March 31, 2020, the targeted allocation for the plans were approximately 75% equity investments and 25% fixed income investments.

Financial instruments included in pension plan assets are categorized into three tiers. These tiers include a fair value hierarchy of three levels, based on the degree of subjectivity inherent in the valuation methodology as follows:

- Level 1 Quoted prices for identical assets in active markets.
- Level 2 Quoted prices for similar assets in active markets with inputs that are observable, either directly or indirectly.

Fair Value Measurements at March 31, 2020

Level 3 - Unobservable prices or inputs in which little or no market data exists.

The fair value of our pension benefits plan assets at March 31, 2020 and 2019 by asset category is as follows:

(In thousands)		Total	fo	Quoted Prices in ctive Markets for Identical Assets (Level 1)		Significant Other Observable Inputs (Level 2)		Significant Other Unobservable Inputs (Level 3)
Cash	\$	302	\$	302	\$	_	\$	_
Insured annuities		14,522		_		14,522		_
Insurance contracts		4,345		_		_		4,345
Common and collective trusts valued at net asset value:								
Equity security trusts		47,187		_		_		_
Debt security trusts		45,847		_		_		_
Total Plan Assets	\$	112,203	\$	302	\$	14,522	\$	4,345
	Fair Value Measurements at March 31, 2019							
(In thousands)		Total	A	Quoted Prices in Active Markets for Identical Assets		Significant Other Observable Inputs (Level 2)		Significant Other Unobservable Inputs
Cash	<u> </u>	450) \$	(Level 1)	<u> </u>			(Level 3)
Insured annuities	Ą	14,720		430	, ф -	14,720		.
Insurance contracts		5,089				14,720	_	5,089
Common and collective trusts valued at net asset value:		3,00	,					3,007
Equity security trusts		73,532	,			_	_	_
Debt security trusts		23,713				_	_	
Total Plan Assets	\$	117,504		450	\$	14,720)	\$ 5,089
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NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Collective investment trusts are measured at fair value using the net asset value per share practical expedient. These trusts have not been categorized in the fair value hierarchy and are being presented in the tables above to permit a reconciliation of the fair value hierarchy to the total plan assets.

The fair value measurement of plan assets using significant unobservable inputs (Level 3) changed during fiscal year 2020 due to the following:

	Insura contr		
Balance at March 31, 2018	\$	5,484	
Gains (losses) related to assets still held at year-end		29	
Transfers out of Level 3		(132)	
Foreign currency		(292)	
Balance at March 31, 2019	\$	5,089	
Gains (losses) related to assets still held at year-end		62	
Transfers out of Level 3		(664)	
Foreign currency		(142)	
Balance at March 31, 2020	\$	4,345	

Cash Flows. We contribute amounts to our defined benefit pension plans at least equal to the minimum amounts required by applicable employee benefit laws and local tax laws. We expect to make contributions of approximately \$3,839 during fiscal 2021.

Based upon the actuarial assumptions utilized to develop our benefit obligations at March 31, 2020, the following benefit payments are expected to be made to plan participants:

	Other Defined Benefit Pension Plans	Other Post- Retirement Benefits Plan		
2021	\$ 5,872	\$ 1,510		
2022	6,025	1,392		
2023	6,600	1,252		
2024	6,336	1,115		
2025	6,518	1,007		
2026-2031	35,292	3,726		

The Medicare Prescription Drug, Improvement and Modernization Act of 2003 (the "Act") provides a prescription drug benefit for Medicare beneficiaries, a benefit we provide to Medicare eligible retirees covered by our post-retirement benefits plan. We have concluded that the prescription drug benefit provided in our post-retirement benefit plan is considered to be actuarially equivalent to the benefit provided under the Act and thus qualifies for the subsidy under the Act. Benefits are subject to a per capita per month cost cap and any costs above the cap become the responsibility of the retiree. Under the plan, the subsidy is applied to reduce the retiree responsibility. As a result, the expected future subsidy no longer reduces our accumulated post-retirement benefit obligation and net periodic benefit cost. We collected subsidies totaling approximately \$708 and \$706, during fiscal 2020 and fiscal 2019, respectively, which reduced the retiree responsibility for costs in excess of the caps established in the post-retirement benefit plan.

Defined Contribution Plans. We maintain a 401(k) defined contribution plan for eligible U.S. employees, a 401(k) defined contribution plan for eligible Puerto Rico employees and similar savings plans for certain employees in Canada, United Kingdom, Ireland, and Finland. We provide a match on a specified portion of an employee's contribution. The U.S. plan assets are held in trust and invested as directed by the plan participants. The Canadian plan assets are held by insurance companies. The aggregate fair value of the U.S. plan assets was \$668,960 at March 31, 2020. At March 31, 2020, the U.S. plan held 555,080 STERIS ordinary shares with a fair value of \$77,695. We paid dividends of \$855, \$826, and \$781 to the plan and participants on STERIS shares held by the plan for the years ended March 31, 2020, 2019, and 2018, respectively. We contributed approximately \$27,818, \$25,935, and \$24,037, to the defined contribution plans for the years ended March 31, 2020, 2019, and 2018, respectively.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

We also maintain a domestic non-qualified deferred compensation plan covering certain employees, which formerly allowed for the deferral of compensation for an employee-specified term or until retirement or termination. There have been no employee contributions made to this plan since fiscal 2012. The Plan was amended in fiscal 2012 to disallow deferrals of salary payable in 2012 and subsequent calendar years and of commissions and other incentive compensation payable in respect of the 2013 and subsequent fiscal years. We hold investments in mutual funds to satisfy future obligations of the plan. We account for these assets as available-for-sale securities and they are included in "Other assets" on our accompanying Consolidated Balance Sheets, with a corresponding liability for the plan's obligation recorded in "Accrued expenses and other." The aggregate value of the assets was \$1,273 and \$1,400 at March 31, 2020 and March 31, 2019, respectively. Realized gains and losses on these investments are recorded in "Interest and miscellaneous income" within "Non-operating expenses" on our accompanying Consolidated Statements of Income. Changes in the fair value of the assets are recorded in other comprehensive income on our accompanying balance sheets.

10. COMMITMENTS AND CONTINGENCIES

We are, and will likely continue to be, involved in a number of legal proceedings, government investigations, and claims, which we believe generally arise in the course of our business, given our size, history, complexity, and the nature of our business, products, Customers, regulatory environment, and industries in which we participate. These legal proceedings, investigations and claims generally involve a variety of legal theories and allegations, including, without limitation, personal injury (e.g., slip and falls, burns, vehicle accidents), product liability or regulation (e.g., based on product operation or claimed malfunction, failure to warn, failure to meet specification, or failure to comply with regulatory requirements), product exposure (e.g., claimed exposure to chemicals, asbestos, contaminants, radiation), property damage (e.g., claimed damage due to leaking equipment, fire, vehicles, chemicals), commercial claims (e.g., breach of contract, economic loss, warranty, misrepresentation), financial (e.g., taxes, reporting), employment (e.g., wrongful termination, discrimination, benefits matters), and other claims for damage and relief.

We believe we have adequately reserved for our current litigation and claims that are probable and estimable, and further believe that the ultimate outcome of these pending lawsuits and claims will not have a material adverse effect on our consolidated financial position or results of operations taken as a whole. Due to their inherent uncertainty, however, there can be no assurance of the ultimate outcome or effect of current or future litigation, investigations, claims or other proceedings (including without limitation the matters discussed below). For certain types of claims, we presently maintain insurance coverage for personal injury and property damage and other liability coverages in amounts and with deductibles that we believe are prudent, but there can be no assurance that these coverages will be applicable or adequate to cover adverse outcomes of claims or legal proceedings against us.

On May 31, 2012, our Albert Browne Limited subsidiary received a warning letter from the FDA regarding chemical indicators manufactured in the United Kingdom. These devices are intended for the monitoring of certain sterilization and other processes. The FDA warning letter stated that the agency had concerns regarding operational business processes. In the second half of calendar 2019, the FDA conducted a comprehensive inspection of the Albert Browne facility in question. In a May 12, 2020 email, the FDA provided the Company with a copy of the Inspection Report. In that same email the FDA advised the Company that the email would serve as a "No Action Indicated" notice and that it was finalizing a Warning Letter Closeout to be provided to the Company. These actions bring this matter to a favorable conclusion for the Company.

Civil, criminal, regulatory or other proceedings involving our products or services could possibly result in judgments, settlements or administrative or judicial decrees requiring us, among other actions, to pay damages or fines or effect recalls, or be subject to other governmental, Customer or other third party claims or remedies, which could materially effect our business, performance, prospects, value, financial condition, and results of operations.

For additional information regarding these matters, see the risks and uncertainties described under the title "product related regulations and claims" in Item 1A. of this Annual Report on Form 10-K.

From time to time, STERIS is also involved in legal proceedings as a plaintiff involving contract, patent protection, and other claims asserted by us. Gains, if any, from these proceedings are recognized when they are realized.

We are subject to taxation from United States federal, state and local, and foreign jurisdictions. Tax positions are settled primarily through the completion of audits within each individual jurisdiction or the closing of statutes of limitation. Changes in applicable tax law or other events may also require us to revise past estimates. We describe income taxes further in Note 8 to our consolidated financial statements titled, "Income Taxes" in this Annual Report on Form 10-K.

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Additional information regarding our contingencies is included in Item 7 of Part II titled, "Management's Discussion and Analysis of Financial Conditions and Results of Operations under "Contingencies".

As of March 31, 2020 and 2019, our commercial commitments totaled \$80,230 and \$73,765, respectively. Commercial commitments include standby letters of credit, letters of credit required as security under our self-insured risk retention policies, and other potential cash outflows resulting from an event that requires payment by us. Approximately \$12,474 and \$7,794 of the March 31, 2020 and 2019 totals, respectively, relate to letters of credit required as security under our self-insured risk retention policies.

As of March 31, 2020, we had minimum purchase commitments with suppliers for raw material purchases totaling \$63,054. As of March 31, 2020, we also had commitments of \$91,077 for long term construction contracts.

Leases

We lease manufacturing, warehouse and office space, service facilities, vehicles, equipment and communication systems. Certain leases contain options that provide us with the ability to extend the lease term. Such options are included in the lease term when it is reasonably certain that the option will be exercised. We made an accounting policy election to not recognize lease assets or lease liabilities for leases with a lease term of twelve months or less.

We determine if an agreement contains a lease and classify our leases as operating or finance at the lease commencement date. Finance leases are generally those leases for which we will pay substantially all the underlying asset's fair value or will use the asset for all or a major part of its economic life, including circumstances in which we will ultimately own the asset. Lease assets arising from finance leases are included in property, plant and equipment, net and the liabilities are included in other liabilities. For finance leases, we recognize interest expense using the effective interest method and we recognize amortization expense on the lease asset over the shorter of the lease term or the useful life of the asset. Our finance leases are not material as of March 31, 2020 and for the twelve month period then ended.

Operating lease assets and liabilities are recognized at the commencement date of the lease based on the present value of lease payments over the lease term. Lease assets represent the right to use an underlying asset for the lease term and lease liabilities represent the obligation to make lease payments arising from the lease. As most leases do not provide an implicit interest rate, we estimate an incremental borrowing rate to determine the present value of lease payments. Our estimated incremental borrowing rate reflects a secured rate based on recent debt issuances, our estimated credit rating, lease term, as well as publicly available data for instruments with similar characteristics. For operating leases, we recognize lease cost on a straight-line basis over the term of the lease. When accounting for leases, we combine payments for leased assets, related services and other components of a lease.

The components of operating lease expense are as follows:

	M	Iarch 31, 2020
Fixed operating lease expense	\$	28,252
Variable operating lease expense		5,449
Total operating lease expense	\$	33,701

Year Ended

Supplemental cash flow information related to operating leases is as follows:

	Ma	Ended rch 31,
Cash paid for amounts included in the measurement of operating lease liabilities	\$	27,613
Right-of-use assets obtained in exchange for operating lease obligations, net	\$	44,636

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

Maturities of lease liabilities at March 31, 2020 are as follows:

	M	
2021	\$	25,302
2022		21,064
2023		17,271
2024		14,045
2025 and thereafter		96,249
Total operating lease payments		173,931
Less imputed interest		40,008
Total operating lease liabilities	\$	133,923

In the preceding table, the future minimum annual rentals payable under noncancelable leases denominated in foreign currencies have been calculated using March 31, 2020 foreign currency exchange rates.

Supplemental information related to operating leases is as follows:

	March 31,
	2020
Weighted-average remaining lease term of operating leases	11.5 years
Weighted-average discount rate of operating leases	4.4%

Prior to the adoption of ASU 2016-02, "Leases" (Topic 842) future minimum annual rentals payable under noncancelable operating lease agreements in excess of one year as of March 31, 2019 were as follows:

	March 31, 2019	
2020	\$ 24,00	8
2021	18,56	7
2022	13,91	7
2023	11,92	9
2024 and thereafter	93,93	9
Total minimum lease payments	\$ 162,36	0

In the preceding table, the future minimum annual rentals payable under noncancelable leases denominated in foreign currencies have been calculated using March 31, 2019 foreign currency exchange rates.

11. BUSINESS SEGMENT INFORMATION

We operate and report in four reportable business segments: Healthcare Products, Healthcare Specialty Services, Life Sciences, and Applied Sterilization Technologies. Corporate is presented separately and contains the costs that are associated with being a publicly traded company and certain other corporate costs.

Our Healthcare Products segment offers infection prevention and procedural solutions for healthcare providers worldwide, including consumable products, equipment maintenance and installation services, and capital equipment.

Our Healthcare Specialty Services segment provides a range of specialty services for healthcare providers including hospital sterilization services and instrument and scope repairs.

Our Life Sciences segment offers consumable products, equipment maintenance and specialty services for pharmaceutical manufacturers and research facilities, and capital equipment.

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(dollars in thousands, except per share amounts and as noted)

Our Applied Sterilization Technologies ("AST") segment provides contract sterilization and testing services for medical device and pharmaceutical manufacturers.

We disclose a measure of segment income that is consistent with the way management operates and views the business. The accounting policies for reportable segments are the same as those for the consolidated Company. In fiscal 2019, we ceased the allocation of certain corporate costs to our segments to align with internal management measures. The prior period operating income measures have been recast for comparability.

For the year ended March 31, 2020, revenues from a single Customer did not represent ten percent or more of any reportable segment's revenues.

Years Ended March 31,	2020	2019	2018
Revenues:			
Healthcare Products	\$ 1,423,198	\$ 1,338,428	\$ 1,276,054
Healthcare Specialty Services	563,611	510,057	469,065
Life Sciences	416,939	378,558	361,590
Applied Sterilization Technologies	627,147	555,127	513,287
Total revenues	\$ 3,030,895	\$ 2,782,170	\$ 2,619,996
Operating income (loss):			
Healthcare Products	356,419	323,684	294,162
Healthcare Specialty Services	64,217	64,222	58,458
Life Sciences	144,088	132,129	123,889
Applied Sterilization Technologies	 270,917	221,828	196,297
Total reportable segments	835,641	741,863	672,806
Corporate	(207,015)	(184,900)	(162,999)
Total operating income before adjustments	\$ 628,626	\$ 556,963	\$ 509,807
Less: Adjustments			
Amortization of acquired intangible assets (1)	71,675	86,878	67,793
Acquisition and integration related charges (2)	8,225	8,901	16,211
Redomiciliation and tax restructuring costs (3)	3,699	8,783	_
(Gain) on fair value adjustment of acquisition related contingent consideration (1)	_	(842)	(593)
Net loss (gain) on divestiture of businesses (1)	1,770	(1,370)	14,547
Amortization of property "step up" to fair value (1)	2,392	2,440	1,599
Restructuring charges (4)	3,143	40,708	103
Impact of the U.S. Tax Cuts and Jobs Act (5)		_	10,264
COVID-19 incremental costs (6)	749		_
Total operating income	\$ 536,973	\$ 411,465	\$ 399,883

⁽¹⁾ For more information regarding our recent acquisitions and divestitures see Note 18 titled, "Business Acquisitions and Divestitures". Amortization of purchased intangible assets fiscal 2019 total includes an impairment charge of \$16,249, see Note 3 titled, "Goodwill and Intangible Assets", for more information.

⁽²⁾ Acquisition and integration related charges include transaction costs and integration expenses associated with acquisitions.

⁽³⁾ Costs incurred in connection with the Redomiciliation and subsequent tax restructuring.

⁽⁴⁾ For more information regarding our restructuring activities see Note 2 titled, "Restructuring".

⁽⁵⁾ Represents a one-time special employee bonus paid to most U.S. employees and associated professional fees.

⁽⁶⁾ COVID-19 incremental costs includes the additional costs attributable to COVID-19 such as enhanced cleaning protocols, personal protective equipment for our employees, event cancellation fees, and payroll costs associated with our response to COVID-19, net of any government subsidies available.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

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Assets include the current and long-lived assets directly attributable to the segment based on the management of the location or on utilization. Certain corporate assets were allocated to the reportable segments based on revenues. Assets attributed to sales and distribution locations are only allocated to the Healthcare Products and Life Sciences segments.

Individual facilities, equipment, and intellectual properties are utilized for production by both the Healthcare Products and Life Sciences segments at varying levels over time. As a result, an allocation of total assets, capital expenditures, and depreciation and amortization is not meaningful to the individual performance of the Healthcare Products and Life Sciences segments. Therefore, their respective amounts are reported together.

March 31,		2020	2019
Assets:			
Healthcare Products and Life Sciences		\$ 1,809,636	\$ 1,611,852
Healthcare Specialty Services		895,741	805,349
Applied Sterilization Technologies		2,720,205	2,655,870
Total assets		\$ 5,425,582	\$ 5,073,071
Years Ended March 31,	2020	2019	2018
Capital Expenditures			
Healthcare Products and Life Sciences	\$ 44,029	\$ 49,688	\$ 52,767
Healthcare Specialty Services	40,619	39,950	16,497
Applied Sterilization Technologies	129,868	100,077	96,193
Total Capital Expenditures	\$ 214,516	\$ 189,715	\$ 165,457
Depreciation, Depletion, and Amortization			
Healthcare Products and Life Sciences (1) (2)	\$ 59,150	\$ 81,264	\$ 52,025
Healthcare Specialty Services	33,043	33,392	29,269
Applied Sterilization Technologies (1)	105,042	111,265	97,038
Total Depreciation, Depletion, and Amortization	\$ 197,235	\$ 225,921	\$ 178,332

⁽¹⁾ The fiscal 2020 and 2019 totals include the impact of Restructuring see Note 2 titled, "Restructuring" for additional information.

Financial information for each of our United States and international geographic areas is presented in the following table. Revenues are based on the location of these operations and their Customers. Property, plant and equipment, net are those assets that are identified within the operations in each geographic area.

March 31,		2020	2019
Property, Plant, and Equipment, Net			
Ireland		\$ 47,459	\$ 41,137
United States		632,333	577,113
Other locations		432,063	413,332
Property, Plant, and Equipment, Net		\$ 1,111,855	\$ 1,031,582
Years Ended March 31,	2020	2019	2018
Revenues:			
Ireland	\$ 63,821	\$ 56,784	\$ 48,246
United States	2,211,722	1,976,814	1,836,414
Other locations	755,352	748,572	735,336
Total Revenues	\$ 3,030,895	\$ 2,782,170	\$ 2,619,996

⁽²⁾ The fiscal 2019 total includes an impairment charge see Note 3 titled, "Goodwill and Intangible Assets", for additional information.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

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Years Ended March 31,	2020	2019	2018
Healthcare Products:			
Capital equipment	\$ 592,436	568,811	527,402
Consumables	454,518	414,969	412,495
Service	376,244	354,648	336,157
Total Healthcare Products Revenues	\$ 1,423,198	\$ 1,338,428	\$ 1,276,054
Total Healthcare Specialty Services Revenues	\$ 563,611	\$ 510,057	\$ 469,065
Life Sciences:			
Capital equipment	\$ 112,747	102,714	100,555
Consumables	185,904	161,780	150,656
Service	118,288	114,064	110,379
Total Life Sciences Revenues	\$ 416,939	\$ 378,558	\$ 361,590
Applied Sterilization Technologies Service Revenues	\$ 627,147	\$ 555,127	\$ 513,287
Total Revenues	\$ 3,030,895	\$ 2,782,170	\$ 2,619,996

Effective April 1, 2020, and consistent with the way management will operate and view the business, the current Healthcare Products and Healthcare Specialty Services segments will be combined and reported as one segment, simply called Healthcare. Going forward we will operate and report in three business segments: Healthcare, Life Sciences and Applied Sterilization Technologies. Corporate will continue to be presented separately and contain the costs that are associated with being a publicly traded company and certain other corporate costs.

12. SHARES AND PREFERRED SHARES

Ordinary Shares

In connection with the Redomiciliation, STERIS UK shareholders received STERIS plc shares pursuant to a scheme of arrangement under UK law. Each STERIS UK ordinary shareholder received one ordinary share, par value \$75.00, of STERIS plc for each STERIS UK ordinary share held, which STERIS UK shares were canceled. On May 3, 2019, the par value of STERIS plc shares issued pursuit to the scheme of arrangement was reduced to \$0.001 per share.

We calculate basic earnings per share based upon the weighted average number of shares outstanding. We calculate diluted earnings per share based upon the weighted average number of shares outstanding plus the dilutive effect of share equivalents calculated using the treasury stock method. The following is a summary of shares and share equivalents outstanding used in the calculations of basic and diluted earnings per share:

Years ended March 31,	2020	2019	2018
Denominator (shares in thousands):		,	
Weighted average shares outstanding—basic	84,778	84,577	85,028
Dilutive effect of share equivalents	863	891	685
Weighted average shares outstanding and share equivalents—diluted	85,641	85,468	85,713

Options to purchase the following number of shares were outstanding but excluded from the computation of diluted earnings per share because the combined exercise prices, unamortized fair values, and assumed tax benefits upon exercise were greater than the average market price for the shares during the periods, so including these options would be anti-dilutive:

Years ended March 31,	2020	2019	2018
Number of ordinary share options (shares in thousands)	285	352	393

Additional Authorized Shares

The Company has an additional authorized share capital of 50,000,000 preferred shares of \$0.001 par value each, plus 25,000 deferred ordinary shares of €1.00 par value each, in order to satisfy minimum statutory capital requirements for all Irish public limited companies.

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13. REPURCHASE OF ORDINARY SHARES

On August 9, 2016, STERIS UK announced that its Board of Directors had authorized the purchase of up to \$300,000 (net of taxes, fees and commissions) of our ordinary shares. As a result of the Redomiciliation, that share repurchase authorization terminated.

On May 7, 2019, our Board of Directors authorized the continuation of the share repurchase program resulting in a share repurchase authorization of \$78,979 (net of taxes, fees and commissions).

On July 30, 2019, our Board of Directors approved an increase to the May 7, 2019 authorization of an additional amount of \$300,000 (net of taxes, fees and commissions). As of March 31, 2020, there was approximately \$338,979 (net of taxes, fees and commissions) of remaining availability under the authorization.

Under the authorizations, the Company may repurchase its shares from time to time through open market purchases, including 10b5-1 plans. Any repurchase program may be activated, suspended or discontinued at any time.

During fiscal 2020, we repurchased 273,259 of our ordinary shares for the aggregate amount of \$40,000 (net of fees and commissions) pursuant to the 2019 authorizations. During fiscal 2019, we repurchased 651,093 of our ordinary shares for the aggregate amount of \$72,082 (net of fees and commissions) pursuant to the 2016 authorization. During fiscal 2018, we repurchased 664,963 of our ordinary shares for the aggregate amount of \$58,939 (net of fees and commissions) pursuant to the 2016 authorization.

During fiscal 2020, we obtained 122,884 of our ordinary shares in the aggregate amount of \$11,235 in connection with share based compensation award programs. During fiscal 2019, we obtained 112,356 of our ordinary shares in the aggregate amount of \$8,262 in connection with share based compensation award programs. During fiscal 2018, we obtained 127,903 of our ordinary shares in the aggregate amount of \$7,014 in connection with share based compensation award programs.

14. SHARE-BASED COMPENSATION

We maintain a long-term incentive plan that makes available shares for grants, at the discretion of the Board of Directors or Compensation Committee of the Board of Directors, to officers, directors, and key employees in the form of stock options, restricted shares, restricted share units, stock appreciation rights and share grants. We satisfy share award incentives through the issuance of new ordinary shares.

Stock options provide the right to purchase our shares at the market price on the date of grant, or for options granted to employees in fiscal 2019 and thereafter, 110% of the market price on the date of grant, subject to the terms of the plan and agreements. Generally, one-fourth of the stock options granted to employees become exercisable for each full year of employment following the grant date. Stock options granted generally expire 10 years after the grant date, or in some cases earlier if the option holder is no longer employed by us. Restricted shares and restricted share units generally cliff vest after a four year period or vest in tranches of one-fourth of the number granted for each year of employment after the grant date. As of March 31, 2020, 3,961,998 shares remained available for grant under the long-term incentive plan.

The fair value of share-based stock option compensation awards was estimated at their grant date using the Black-Scholes-Merton option pricing model. This model was developed for use in estimating the fair value of traded options that have no vesting restrictions and are fully transferable, characteristics that are not present in our option grants. If the model permitted consideration of the unique characteristics of employee stock options, the resulting estimate of the fair value of the stock options could be different. The value of the portion of the award that is ultimately expected to vest is recognized as expense over the requisite service periods in our Consolidated Statements of Income. The expense is classified as cost of goods sold or selling, general and administrative expenses in a manner consistent with the employee's compensation and benefits.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

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The following weighted-average assumptions were used for options granted during fiscal 2020, fiscal 2019 and fiscal 2018:

	Fiscal 2020	Fiscal 2019	Fiscal 2018
Risk-free interest rate	2.26%	2.64%	2.01%
Expected life of options	6.2 years	6.2 years	5.7 years
Expected dividend yield of stock	1.22%	1.47%	1.58%
Expected volatility of stock	20.27%	19.91%	22.08%

The risk-free interest rate is based upon the U.S. Treasury yield curve. The expected life of options is reflective of historical experience, vesting schedules and contractual terms. The expected dividend yield of stock represents our best estimate of the expected future dividend yield. The expected volatility of stock is derived by referring to our historical stock prices over a time frame similar to that of the expected life of the grant. An estimated forfeiture rate of 2.77%, 2.37% and 2.25% was applied in fiscal 2020, 2019 and 2018 respectively. This rate is calculated based upon historical activity and represents an estimate of the granted options not expected to vest. If actual forfeitures differ from this calculated rate, we may be required to make additional adjustments to compensation expense in future periods. The assumptions used above are reviewed at the time of each significant option grant, or at least annually.

A summary of share option activity is as follows:

	Number of Options	Weighted Average Exercise Price	Average Remaining Contractual Term	ggregate ntrinsic Value
Outstanding at March 31, 2019	2,104,685	72.82		
Granted	345,138	147.22		
Exercised	(613,086)	57.29		
Forfeited	(40,611)	122.61		
Outstanding at March 31, 2020	1,796,126	91.29	6.8 years	\$ 89,800
Exercisable at March 31, 2020	922,708	69.52	5.6 years	\$ 65,136

We estimate that 857,860 of the non-vested stock options outstanding at March 31, 2020 will ultimately vest.

The aggregate intrinsic value in the table above represents the total pre-tax difference between the \$139.97 closing price of our ordinary shares on March 31, 2020 over the exercise prices of the stock options, multiplied by the number of options outstanding or outstanding and exercisable, as applicable. The aggregate intrinsic value is not recorded for financial accounting purposes and the value changes daily based on the daily changes in the fair market value of our ordinary shares.

The total intrinsic value of stock options exercised during the years ended March 31, 2020, 2019 and 2018 was \$57,683, \$25,371 and \$16,096, respectively. Net cash proceeds from the exercise of stock options were \$34,731, \$13,308 and \$11,093 for the years ended March 31, 2020, 2019 and 2018, respectively. The tax benefit from stock option exercises was \$16,440, \$8,306 and \$6,581 for the years ended March 31, 2020, 2019 and 2018, respectively.

The weighted average grant date fair value of stock option grants was \$23.52, \$18.12 and \$15.51 for the years ended March 31, 2020, 2019 and 2018, respectively.

Stock appreciation rights ("SARS") carry generally the same terms and vesting requirements as stock options except that they are settled in cash upon exercise and therefore, are classified as liabilities. The fair value of the outstanding SARS as of March 31, 2020, 2019 and 2018 was \$544, \$889, and \$1,437, respectively. The fair value of outstanding SARS is revalued at each reporting date and the related liability and expense are adjusted appropriately.

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A summary of the non-vested restricted share activity is presented below:

	Number of Restricted Shares	Number of Restricted Share Units	We	ighted-Average Grant Date Fair Value
Non-vested at March 31, 2019	676,373	33,219	\$	80.86
Granted	156,901	14,553		135.86
Vested	(221,606)	(14,999)		74.63
Forfeited	(35,838)	(1,879)		93.56
Non-vested at March 31, 2020	575,830	30,894	\$	98.07

Restricted shares granted are valued based on the closing stock price at the grant date. The value of restricted shares and units that vested during fiscal 2020 was \$17,657.

As of March 31, 2020, there was a total of \$42,056 in unrecognized compensation cost related to non-vested share-based compensation granted under our share-based compensation plans. We expect to recognize the cost over a weighted average period of 2.1 years.

15. FINANCIAL AND OTHER GUARANTEES

We generally offer a limited parts and labor warranty on capital equipment. The specific terms and conditions of those warranties vary depending on the product sold and the countries where we conduct business. We record a liability for the estimated cost of product warranties at the time product revenues are recognized. The amounts we expect to incur on behalf of our Customers for the future estimated cost of these warranties are recorded as a current liability on the accompanying Consolidated Balance Sheets. Factors that affect the amount of our warranty liability include the number and type of installed units, historical and anticipated rates of product failures, and material and service costs per claim. We periodically assess the adequacy of our recorded warranty liabilities and adjust the amounts as necessary.

Changes in our warranty liability during the periods presented are as follows:

Years Ended March 31,	2020	2019	2018
Balance, Beginning of Year	\$ 7,194 \$	6,872 \$	6,861
Warranties issued during the period	12,311	11,177	12,305
Settlements made during the period	(12,124)	(10,855)	(12,294)
Balance, End of Year	\$ 7,381 \$	7,194 \$	6,872

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16. DERIVATIVES AND HEDGING

From time to time, we enter into forward contracts to hedge potential foreign currency gains and losses that arise from transactions denominated in foreign currencies, including inter-company transactions. We may also enter into commodity swap contracts to hedge price changes in nickel that impact raw materials included in our cost of revenues. We do not use derivative financial instruments for speculative purposes. These contracts are not designated as hedging instruments and do not receive hedge accounting treatment; therefore, changes in their fair value are not deferred but are recognized immediately in the Consolidated Statements of Income. At March 31, 2020, we held a foreign currency forward contract to buy 6.0 million Canadian dollars. At March 31, 2020, we held commodity swap contracts to buy 715.2 thousand pounds of nickel.

		Asset Der	ivatives		Liability D	eriv	atives
	Fa	ir Value at	Fa	ir Value at	Fair Value at		Fair Value at
Balance Sheet Location	Mar	ch 31, 2020	Mai	rch 31, 2019	March 31, 2020		March 31, 2019
Prepaid & Other	\$	124	\$	552	\$ _	\$	_
Accrued expenses and other	\$	_	\$		\$ 912	\$	278

The following table presents the impact of derivative instruments and their location within the Consolidated Statements of Income:

		Amount of (l	oss) g	gain recogniz	zed in	income
	Location of (loss) gain recognized in	Ye	ars Eı	nded March	31,	
	income	2020		2019		2018
Foreign currency forward contracts	Selling, general and administrative	\$ 798	\$	235	\$	(1,357)
Commodity swap contracts	Cost of revenues	\$ (660)	\$	434	\$	373

Additionally, we hold our debt in multiple currencies to fund our operations and investments in certain subsidiaries. We designate portions of non-functional currency denominated intercompany loans as hedges of portions of net investments in foreign operations. Net debt designated as non-derivative net investment hedging instruments totaled \$45,765 at March 31, 2020. These hedges are designed to be fully effective and any associated gain or loss is recognized in Accumulated Other Comprehensive Income and will be reclassified to income in the same period when a gain or loss related to the net investment in the foreign operation is included in income.

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17. FAIR VALUE MEASUREMENTS

Fair value is defined as the price that would be received to sell an asset or that would be paid to transfer a liability in an orderly transaction between market participants at the measurement date. We estimate the fair value of financial assets and liabilities using available market information and generally accepted valuation methodologies. The inputs used to measure fair value are classified into three tiers. These tiers include Level 1, defined as observable inputs such as quoted prices in active markets; Level 2, defined as inputs other than quoted prices in active markets that are either directly or indirectly observable; and Level 3, defined as unobservable inputs in which little or no market data exists, therefore requiring the entity to develop its own assumptions. The following table shows the fair value of our financial assets and liabilities at March 31, 2020 and March 31, 2019:

							Fai	ir Value M	easur	ements	3			
	Carryin	ıg V	Value	Quoted in Active for Identic	N	Iarkets		Significa Observab				Signifi Unobser Inpu	vable	
				Lev	el	1		Leve	2 1			Level	3	
At March 31,	 2020		2019	2020		2019		2020	20	19		2020	20	19
Assets:												'		
Cash and cash equivalents	\$ 319,581	\$	220,633	\$ 319,581	\$	220,633	\$	_	\$	_	\$	_ \$,	_
Forward and swap contracts (1)	124		552	_		_		124		552		_		_
Equity investments (2)	9,624		13,873	9,624		13,873		_		_		_		_
Other investments	2,507		2,545	2,507		2,545		_		_		_		_
Liabilities:														
Forward and swap contracts (1)	\$ 912	\$	278	\$ _	\$	_	\$	912	\$	278	\$	— \$,	_
Deferred compensation plans (2)	1,475		1,564	1,475		1,564		_		_		_		_
Long term debt (3)	1,150,521		1,183,227	_		_		1,143,978	1,20	0,558		_		_
Contingent consideration obligations (4)	15,988		5,950	_		_		_		_		15,988		5,950

⁽¹⁾ The fair values of forward and swap contracts are based on period-end forward rates and reflect the value of the amount that we would pay or receive for the contracts involving the same notional amounts and maturity dates.

⁽²⁾ We maintain a frozen domestic non-qualified deferred compensation plan covering certain employees, which allowed for the deferral of payment of previously earned compensation for an employee-specified term or until retirement or termination. Amounts deferred can be allocated to various hypothetical investment options (compensation deferrals have been frozen under the plan). We hold investments to satisfy the future obligations of the plan. Employees who made deferrals are entitled to receive distributions of their hypothetical account balances (amounts deferred, together with earnings (losses)). We also hold an investment in the common stock of Servizi Italia, S.p.A, a leading provider of integrated linen washing and outsourced sterile processing services to hospital Customers. Beginning in fiscal 2019, changes in the fair value of these investments are recorded in the "Interest income and miscellaneous expense line" of the Consolidated Statement of Income. During fiscal 2020 and fiscal 2019 we recorded losses of \$3,579 and \$2,731, respectively, related to these investments.

⁽³⁾ We estimate the fair value of our long-term debt using discounted cash flow analyses, based on our current incremental borrowing rates for similar types of borrowing arrangements.

⁽⁴⁾ Contingent consideration obligations arise from prior business acquisitions. The fair values are based on discounted cash flow analyses reflecting the possible achievement of specified performance measures or events and captures the contractual nature of the contingencies, commercial risk, and the time value of money. Contingent consideration obligations are classified in the consolidated balance sheets as accrued expense (short-term) and other liabilities (long-term), as appropriate based on the contractual payment dates.

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The changes in Level 3 assets and liabilities measured at fair value on a recurring basis are summarized as follows:

	ontingent isideration
Balance at March 31, 2018	\$ 8,068
Payments	(691)
Reductions and adjustments	(1,466)
Foreign currency translation adjustments	 39
Balance at March 31, 2019	\$ 5,950
Additions	9,907
Foreign currency translation adjustments	131
Balance at March 31, 2020	\$ 15,988

Additions and payments of contingent consideration obligations during fiscal year 2020 and 2019 were primarily related to our fiscal year 2020 and 2019 acquisitions. Refer to Note 18, "Business Acquisitions and Divestitures" for more information.

18. BUSINESS ACQUISITIONS AND DIVESTITURES

Fiscal 2020 Acquisitions

During fiscal 2020, we completed several tuck-in acquisitions which continued to expand our product and service offerings in the Healthcare Products, Healthcare Specialty Services and Applied Sterilization Technologies segments. The aggregate purchase price associated with these transactions was approximately \$120,537, net of cash acquired and including potential contingent consideration of \$9,830 and deferred consideration of \$893.

Fiscal 2019 Acquisitions

During fiscal 2019, we completed a minor purchase to expand our service offerings in the Applied Sterilization Technologies segment. The total purchase price was \$13,313, and was financed with both cash on hand and with credit facility borrowings. Purchase price allocations will be finalized within a measurement period not to exceed one year from closing.

Fiscal 2018 Acquisitions

We completed several minor purchases that continued to expand our product and service offerings in the Healthcare Products, Healthcare Specialty Services and Applied Sterilization Technologies segments. The aggregate purchase price associated with these transactions was approximately \$52,292, net of cash acquired and including contingent consideration of \$5,018. The purchase price for the acquisitions was financed with both cash on hand and with credit facility borrowings.

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Fair Value of Assets Acquired and Liabilities Assumed

The table below summarizes the allocation of the purchase price to the net assets acquired based on fair values at the acquisition dates for our fiscal 2020, 2019 and 2018 acquisitions.

	Fiscal Year 2020	Fiscal Year 2019	Fiscal Year 2018
(dollars in thousands)	All Acquisitions (1)	All Acquisitions	All Acquisitions
Cash	\$ 8,811	\$ —	\$ 235
Accounts receivable	10,331	750	1,464
Inventory	8,999	51	2,289
Property, plant and equipment	9,241	2,004	3,381
Lease right-of-use assets, net	4,462	-	-
Other assets	1,133	479	126
Intangible assets	36,500	4,070	17,404
Goodwill	74,531	6,614	32,384
Total Assets	154,008	13,968	57,283
Current liabilities	(20,659)	(146)	(2,077)
Non-current liabilities	(4,000)	(509)	(2,679)
Total Liabilities	(24,659)	(655)	(4,756)
Net Assets	\$ 129,349	\$ 13,313	\$ 52,527

⁽¹⁾ Purchase price allocation is still preliminary as of March 31, 2020, as valuations have not been finalized.

Acquisition related transaction and integration costs totaled \$8,225, \$8,901, and \$16,211 for the fiscal years ended March 31, 2020, 2019, and 2018, respectively. These costs are included in Selling, general, and administrative expenses in the Consolidated Statements of Income.

Divestitures

Fiscal 2020

During fiscal 2020, we sold the operations of our Healthcare Specialty Services business that were located in China. We recorded proceeds of \$439, net of cash divested, and recognized a pre-tax loss on the sale of \$2,365 in the selling, general and administrative expense line of the Consolidated Statements of Income. The business generated annual revenues of approximately \$5,000.

Fiscal 2018

Synergy Health Healthcare Consumable Solutions

On November 20, 2017, we sold our Synergy Health Healthcare Consumable Solutions ("HCS") business to Vernacare. Annual revenues for the HCS business were approximately \$40,000 and were included in the Healthcare Products segment. We recorded proceeds of \$8,891, net of cash divested, including a working capital adjustment. We also recognized a pre-tax loss on the sale, subject to final working capital adjustments, of \$12,972 in Selling, general, and administrative expense in the Consolidated Statement of Income.

Loans Receivable

In connection with an equity investment of \$4,955, we agreed to provide a credit facility of up to approximately \$10,000 for a term of up to seven years ending in 2025. The loan carries an interest rate of 4% compounded daily and interest is payable annually. Outstanding borrowings under the agreement totaled \$7,084 at March 31, 2020 and \$7,465 at March 31, 2019.

In connection with the fiscal 2017 divestiture of Synergy Health Netherlands Linen Management Services, we entered into a loan agreement to provide financing of up to epsilon15,000 for a term of up to 15 years. The loan carried an interest rate of 4% for the first four years and 12% thereafter. The loan was renegotiated during the third quarter of fiscal 2020. According to the new terms of the loan agreement, the outstanding balance at October 31, 2019, of epsilon7,300, will be repaid in six equal annual

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installments beginning on October 18, 2022. The loan carries an interest rate of 4% for the first four years and 8% thereafter. Outstanding borrowings under the agreement totaled \$8,072 (or €7,300) at March 31, 2020 and \$8,494 (or €7,550) at March 31, 2019.

Amounts for loan receivables as noted above are recorded in the "Other assets" line of our Consolidated balance sheets. Interest income is not material.

19. RECLASSIFICATIONS OUT OF ACCUMULATED OTHER COMPREHENSIVE INCOME (LOSS)

 $\alpha \cdot \alpha$

Amounts in Accumulated Other Comprehensive Income (Loss) are presented net of the related tax. Foreign Currency Translation is not adjusted for income taxes. Accumulated other comprehensive income (loss) shown in our Consolidated Statements of Shareholders' Equity and changes in our balances, net of tax, for the years ended March 31, 2020, 2019 and 2018 were as follows:

		Avai	in (Loss) lable for curities ⁽	Sale		ined Benef Plans ⁽²⁾	iit	Foreign Currency Translation ⁽³⁾			Total Acc Compre		
	20	020	2019	2018	2020	2019	2018	2020	2019	2018	2020	2019	2018
Beginning Balance	\$	_	\$ 1,970	\$ 178	\$ (4,204)	\$ (6,742) \$	(2,355)	\$ (155,574)	\$ 16,457	\$ (238,525)	\$(159,778) \$	11,685	\$(240,702)
Other Comprehensive Income (Loss) before reclassifications		_	_	1,703	1,505	3,920	(2,291)	(73,076)	(172,031)	254,982	(71,571)	(168,111)	254,394
Reclassified from Accumulated Other Comprehensive Income (Loss)		_	_	89	(4,114)	(1,382)	(2,096)	_	_	_	(4,114)	(1,382)	(2,007)
Net current- period Other Comprehensive Income (Loss)		_	_	1,792	(2,609)	2,538	(4,387)	(73,076)	(172,031)	254,982	(75,685)	(169,493)	252,387
Cumulative adjustment to Retained Earnings (4)	\$	_	\$(1,970))\$ —	s –	s — s	· —	s —	\$ —	s —	s — s	5 (1,970)	s —
Ending Balance	\$	_	\$ —	\$ 1,970	\$ (6,813)	\$ (4,204) \$	(6,742)	\$ (228,650)	\$(155,574)	\$ 16,457	\$(235,463) \$	(159,778)	\$ 11,685

⁽¹⁾ Realized gain (loss) on available for sale securities is reported in the Interest income and miscellaneous expense line of the Consolidated Statements of Income for fiscal 2018.

⁽²⁾ Amortization (gain) of defined benefit plan items are reported in the Interest income and miscellaneous expense line of our Consolidated Statements of Income.

⁽³⁾ The effective portion of gain or loss on net debt designated as non-derivative net investment hedging instruments is recognized in Accumulated Other Comprehensive Income and is reclassified to income in the same period when a gain or loss related to the net investment is included in income.

⁽⁴⁾ As a result of the adoption of ASC 2016-01 we recorded a cumulative effect adjustment to our opening fiscal 2019 retained earnings balance that increased retained earnings and decreased accumulated other comprehensive income. See Note 1 titled, "Nature of Operations and Summary of Significant Accounting Policies" for further details.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

20. QUARTERLY RESULTS (UNAUDITED)

quarters Ended		March 31,	D	ecember 31,	September 30,			June 30,
Fiscal 2020								
Revenues:								
Product	\$	393,592	\$	363,795	\$	337,666	\$	307,735
Service		429,399		410,466		399,174		389,068
Total Revenues		822,991		774,261		736,840		696,803
Cost of Revenues:								
Product		210,538		195,105		183,600		160,959
Service		248,393		247,803		234,573		230,001
Total Cost of Revenues		458,931		442,908		418,173		390,960
Gross Profit		364,060		331,353		318,667		305,843
Percentage of Revenues		44.2%		42.8%		43.2%		43.9%
Restructuring Expenses		6		(448)		(274)		1,389
Net Income Attributable to Shareholders	\$	123,316	\$	104,930	\$	94,769	\$	84,590
Basic Income Per Ordinary Share Attributable to Shareholders:								
Net income	\$	1.45	\$	1.24	\$	1.12	\$	1.00
Diluted Income Per Ordinary Share Attributable to Shareholders:								
Net income	\$	1.44	\$	1.23	\$	1.11	\$	0.99
77. 18040								
Fiscal 2019								
Revenues:	_		_		_		_	
Product	\$	374,937	\$	327,639	\$	314,659	\$	278,790
Service	_	393,276		368,599		364,302		359,968
Total Revenues		768,213		696,238		678,961		638,758
Cost of Revenues:								
Product		201,357		182,229		172,107		146,602
Service	_	232,140		227,012		222,190		223,106
Total Cost of Revenues		433,497		409,241		394,297		369,708
Gross Profit		334,716		286,997		284,664		269,050
Percentage of Revenues		43.6%		41.2 %		41.9 %		42.1 %
Restructuring Expenses		4,840		26,147				
Net Income Attributable to Shareholders	\$	108,745	\$	47,858	\$	77,457	\$	69,991
Basic Income Per Ordinary Share Attributable to Shareholders:								
Net income	\$	1.29	\$	0.57	\$	0.92	\$	0.83
Diluted Income Per Ordinary Share Attributable to Shareholders:								
Net income	\$	1.27	\$	0.56	\$	0.91	\$	0.82

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

21. SUBSEQUENT EVENTS

The COVID-19 pandemic began to impact our business late in fiscal 2020. The coronavirus pandemic and related public health recommendations and mandated precautions to mitigate the spread of COVID-19, including deferral of medical procedures and treatments and shelter-in-place orders or similar measures, is negatively affecting, and is expected to continue to affect some of our operations which would impact our financial position and cash flows in fiscal 2021. We have experienced and expect to continue to experience unpredictable fluctuations in demand for certain of our products and services, including some products and services that are experiencing increased demand.

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

(dollars in thousands, except per share amounts and as noted)

SCHEDULE II – VALUATION AND QUALIFYING ACCOUNTS

Description	В	salance at seginning of Period	1	Charges to Costs and Expenses	1	Charges to Other Accounts	De	eductions	В	alance at End of Period
(in thousands)				_						
Year ended March 31, 2020										
Deducted from asset accounts:										
Allowance for trade accounts receivable (1)	\$	9,645	\$	6,760	\$	(247) ⁽³⁾	\$	(4,107) ⁽⁴⁾	\$	12,051
Inventory valuation reserve		19,754		(4,105) (2)		500 (3)		_		16,149
Deferred tax asset valuation allowance		13,478		3,327		(1,927) ⁽³⁾		(987)		13,891
Recorded within liabilities:										
Casualty loss reserves	\$	19,742	\$	6,000	\$	3,007	\$	(5,521)	\$	23,228
Year ended March 31, 2019										
Deducted from asset accounts:										
Allowance for trade accounts receivable (1)	\$	12,472	\$	356	\$	(327) ⁽³⁾	\$	(2,856) (4)	\$	9,645
Inventory valuation reserve		19,639		$(673)^{(2)}$		788 ⁽³⁾				19,754
Deferred tax asset valuation allowance		13,596		4,055		(1,653) ⁽³⁾		(2,520)		13,478
Recorded within liabilities:										
Casualty loss reserves	\$	20,949	\$	4,456	\$	(1,158)	\$	(4,505)	\$	19,742
Year ended March 31, 2018										
Deducted from asset accounts:										
Allowance for trade accounts receivable (1)	\$	10,357	\$	2,183	\$	1,925 ⁽³⁾	\$	(1,993) (4)	\$	12,472
Inventory valuation reserve		17,854		2,446 (2)		$(661)^{(3)}$		_		19,639
Deferred tax asset valuation allowance		16,366		3,535		209 (3)		(6,514)		13,596
Recorded within liabilities:										
Casualty loss reserves	\$	22,718	\$	5,713	\$	(2,563)	\$	(4,919)	\$	20,949

⁽¹⁾ Net allowance for doubtful accounts and allowance for sales and returns.
(2) Provision for excess and obsolete inventory, net of inventory written off.
(3) Change in foreign currency exchange rates and acquired reserves.
(4) Uncollectible accounts written off, net of recoveries.

ITEM 9. CHANGES IN AND DISAGREEMENTS WITH ACCOUNTANTS ON ACCOUNTING AND FINANCIAL DISCLOSURE

None.

ITEM 9A. CONTROLS AND PROCEDURES

EVALUATION OF DISCLOSURE CONTROLS AND PROCEDURES

Our management, including the Principal Executive Officer ("PEO") and Principal Financial Officer ("PFO"), has evaluated the effectiveness of our disclosure controls and procedures, as defined in Exchange Act Rules 13a-15(e) and 15d-15(e), as of the end of the period covered by this Annual Report on Form 10-K. Based on this evaluation, the PEO and PFO have determined that, as of the end of the period covered by this Annual Report on Form 10-K, our disclosure controls and procedures were effective.

CHANGES IN INTERNAL CONTROLS

During the quarter ended March 31, 2020, there were no changes in our internal control over financial reporting that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

MANAGEMENT'S REPORT ON INTERNAL CONTROL OVER FINANCIAL REPORTING

Management is responsible for establishing and maintaining adequate internal control over financial reporting, as such term is defined in the Exchange Act Rules 13a-15(f). Under the supervision and with the participation of management, including the PEO and PFO, we conducted an evaluation of the effectiveness of internal control over financial reporting as of March 31, 2020 based on the framework in 2013 Internal Control-Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission. Based on this evaluation under this framework, management concluded that the internal control over financial reporting was effective as of March 31, 2020. Our evaluation of internal control over financial reporting did not include the internal controls of the entities that were acquired during fiscal 2020. Total assets of the acquired businesses (inclusive of acquired intangible assets and goodwill) represented approximately 4% of our total assets as of March 31, 2020 and approximately 1% of our total revenues for the year ended March 31, 2020. Based on this evaluation under this framework, management concluded that the internal control over financial reporting was effective as of March 31, 2020.

The independent registered public accounting firm that audited the financial statements has issued an attestation report on internal control over financial reporting.

REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

To the Shareholders and the Board of Directors of STERIS plc

Opinion on Internal Control Over Financial Reporting

We have audited STERIS plc and subsidiaries' internal control over financial reporting as of March 31, 2020, based on criteria established in Internal Control-Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission (2013 framework) (the COSO criteria). In our opinion, STERIS plc and subsidiaries (the Company) maintained, in all material respects, effective internal control over financial reporting as of March 31, 2020, based on the COSO criteria.

As indicated in the accompanying Management's Report on Internal Control Over Financial Reporting, management's assessment of and conclusion on the effectiveness of internal control over financial reporting did not include the internal controls of the entities that were acquired during the year ended March 31, 2020, which are included in the fiscal 2020 consolidated financial statements of the Company and constituted approximately 4% of total assets as of March 31, 2020 and approximately 1% of total revenues for the year then ended. Our audit of internal control over financial reporting of the Company also did not include an evaluation of the internal control over financial reporting of the entities that were acquired during the year ended March 31, 2020.

We also have audited, in accordance with the standards of the Public Company Accounting Oversight Board (United States) (PCAOB), the consolidated balance sheets of the Company as of March 31, 2020 and 2019, the related consolidated statements of income, comprehensive income, shareholders' equity and cash flows for each of the three years in the period ended March 31, 2020, and the related notes and the financial statement schedule listed in the Index at Item 15(a) and our report dated May 29, 2020 expressed an unqualified opinion thereon.

Basis for Opinion

The Company's management is responsible for maintaining effective internal control over financial reporting and for its assessment of the effectiveness of internal control over financial reporting included in the accompanying Management's Report on Internal Control Over Financial Reporting. Our responsibility is to express an opinion on the Company's internal control over financial reporting based on our audit. We are a public accounting firm registered with the PCAOB and are required to be independent with respect to the Company in accordance with the U.S. federal securities laws and the applicable rules and regulations of the Securities and Exchange Commission and the PCAOB.

We conducted our audit in accordance with the standards of the PCAOB. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether effective internal control over financial reporting was maintained in all material respects.

Our audit included obtaining an understanding of internal control over financial reporting, assessing the risk that a material weakness exists, testing and evaluating the design and operating effectiveness of internal control based on the assessed risk, and performing such other procedures as we considered necessary in the circumstances. We believe that our audit provides a reasonable basis for our opinion.

Definition and Limitations of Internal Control Over Financial Reporting

A company's internal control over financial reporting is a process designed to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles. A company's internal control over financial reporting includes those policies and procedures that (1) pertain to the maintenance of records that, in reasonable detail, accurately and fairly reflect the transactions and dispositions of the assets of the company; (2) provide reasonable assurance that transactions are recorded as necessary to permit preparation of financial statements in accordance with generally accepted accounting principles, and that receipts and expenditures of the company are being made only in accordance with authorizations of management and directors of the company; and (3) provide reasonable assurance regarding prevention or timely detection of unauthorized acquisition, use, or disposition of the company's assets that could have a material effect on the financial statements.

Because of its inherent limitations, internal control over financial reporting may not prevent or detect misstatements. Also, projections of any evaluation of effectiveness to future periods are subject to the risk that controls may become inadequate because of changes in conditions, or that the degree of compliance with the policies or procedures may deteriorate.

/s/ Ernst & Young LLP Cleveland, Ohio May 29, 2020

ITEM 9B. OTHER INFORMATION

None.

ITEM 10. DIRECTORS, EXECUTIVE OFFICERS AND CORPORATE GOVERNANCE

This Annual Report on Form 10-K incorporates by reference the information appearing under the caption "Nominees for Election as Directors," "Delinquent Section 16(a) Reports," "Board Meetings and Committees," "Shareholder Nominations of Directors and Nominee Criteria" and "Shareholder Proposals" of our definitive proxy statement to be filed with the SEC in connection with our 2020 Annual Meeting of Shareholders (the "Proxy Statement").

Our executive officers serve for a term of one year from the date of election to the next organizational meeting of the Board of Directors and until their respective successors are elected and qualified, except in the case of death, resignation, or removal. Information concerning our executive officers is contained in Item 1 of Part 1 of this Annual Report under the heading "Information about our Executive Officers", and is incorporated herein by reference. We have adopted a code of ethics, our Code of Business Conduct for Employees, that applies to our CEO and CFO and Principal Accounting Officer as well as all of our other employees. We have also adopted a code of ethics, our Director Code of Ethics, which applies to the members of the Company's Board of Directors, including our CEO. Our Code of Business Conduct for Employees and the Director Code of Ethics can be found on our Investor Relations website at www.steris-ir.com. Any amendments or waivers of either of these codes will be made available on this website.

ITEM 11. EXECUTIVE COMPENSATION

This Annual Report on Form 10-K incorporates by reference the information appearing beginning under the captions "Executive Compensation," "Non-Employee Director Compensation" and "Miscellaneous Matters" of the Proxy Statement.

ITEM 12. SECURITY OWNERSHIP OF CERTAIN BENEFICIAL OWNERS AND MANAGEMENT AND RELATED STOCKHOLDER MATTERS

This Annual Report on Form 10-K incorporates by reference the information appearing under the captions "Ownership of Voting Securities" of the Proxy Statement.

The table below presents information concerning all equity compensation plans and individual equity compensation arrangements in effect as of our fiscal year ended March 31, 2020.

Plan Category	Number of securities to be issued upon exercise of outstanding options, warrants and rights	Weighted-average exercise price of outstanding options, warrants and rights (\$)	Number of securities remaining available for future issuance under equity compensation plans (excluding securities reflected in column (a))
	(a)	(b)	(c)
Equity compensation plans approved by security holders	1,796,126	\$91.29	3,961,998
Equity compensation plans not approved by security holders	_	_	_
Total	1,796,126	\$91.29	3,961,998

ITEM 13. CERTAIN RELATIONSHIPS AND RELATED TRANSACTIONS, AND DIRECTOR INDEPENDENCE

RELATED PERSON TRANSACTIONS

This Annual Report on Form 10-K incorporates by reference the information beginning under the captions "Governance Generally", "Board Meetings and Committees" and "Miscellaneous Matters" of the Proxy Statement.

ITEM 14. PRINCIPAL ACCOUNTANT FEES AND SERVICES

This Annual Report on Form 10-K incorporates by reference the information relating to principal accountant fees and services appearing under the caption "Independent Registered Public Accounting Firm" of the Proxy Statement.

ITEM 15. EXHIBITS AND FINANCIAL STATEMENT SCHEDULE

LIST OF CONSOLIDATED FINANCIAL STATEMENTS AND FINANCIAL STATEMENT SCHEDULE

(a) (1) The following consolidated financial statements of STERIS plc and subsidiaries are included in Item 8:

Consolidated Balance Sheets – March 31, 2020 and 2019.

Consolidated Statements of Income – Years ended March 31, 2020, 2019, and 2018.

Consolidated Statements of Comprehensive Income – Years ended March 31, 2020, 2019, and 2018.

Consolidated Statements of Cash Flows – Years ended March 31, 2020, 2019, and 2018.

Consolidated Statements of Shareholders' Equity - Years ended March 31, 2020, 2019, and 2018.

Notes to Consolidated Financial Statements.

(a) (2) The following consolidated financial statement schedule of STERIS plc and subsidiaries is included in Item 8:

Schedule II - Valuation and Qualifying Accounts

All other schedules for which provision is made in the applicable accounting regulation of the SEC are not required under the related instructions or are inapplicable and, therefore, have been omitted.

(a) (3) Exhibits

Exhibit Number	Exhibit Description
3.1	STERIS plc Amended Memorandum and Articles of Association (filed as Exhibit 3.1 to STERIS plc Form 10-K for the fiscal year ended March 31, 2019 (Commission File No. 001-38848) and incorporated herein by reference).*
4.1	Description of Securities Registered Under Section 12 of the Securities Exchange Act of 1934.
10.1	STERIS plc 2006 Long-Term Equity Incentive Plan, as Assumed, Amended and Restated Effective March 28, 2019 (filed as Exhibit 10.1 to STERIS plc Form 8-K filed March 28, 2019 (Commission File No. 001-38848) and incorporated herein by reference).*
10.2	STERIS Corporation Form of Nonqualified Stock Option Agreement for Nonemployee Directors (filed as Exhibit 10.4 to Form 10-Q for the fiscal quarter ended June 30, 2008 (Commission File No. 1-14643), and incorporated herein by reference).*
10.3	STERIS Corporation Form of Nonqualified Stock Option Agreement for Employees (filed as Exhibit 10.2 to Form 10-Q for the fiscal quarter ended June 30, 2011 (Commission File No. 1-14643), and incorporated herein by reference).*
10.4	Amendment to STERIS Corporation Nonqualified Stock Option Agreement (filed as Exhibit 10.11 to Form 10-Q for the fiscal quarter ended December 31, 2012 (Commission File No. 1-14643), and incorporated herein by reference).*
10.5	STERIS Corporation Form of Nonqualified Stock Option Agreement for Nonemployee Directors (filed as Exhibit 10.12 to Form 10-Q for the fiscal quarter ended December 31, 2012 (Commission File No. 1-14643), and incorporated herein by reference).*
10.6	STERIS Corporation Form of Nonqualified Stock Option Agreement for Employees (filed as Exhibit 10.13 to Form 10-Q for the fiscal quarter ended December 31, 2012 (Commission File No. 1-14643), and incorporated herein by reference).*
10.7	STERIS Corporation Form of Nonqualified Stock Option Agreement for Employees (filed as Exhibit 10.14 to Form 10-Q for the fiscal quarter ended December 31, 2012 (Commission File No. 1-14643), and incorporated herein by reference).*
10.8	STERIS Corporation Form of Career Restricted Stock Unit Agreement for Nonemployee Directors (filed as Exhibit 10.33 to Form 10-K for the fiscal year ended March 31, 2013 (Commission File No. 1-14643), and incorporated by reference).*

- 10.9 STERIS Corporation Form of Nonqualified Stock Option Agreement for Nonemployee Directors (filed as Exhibit 10.34 to Form 10-K for the fiscal year ended March 31, 2013 (Commission File No. 1-14643), and incorporated by reference).*
- 10.10 STERIS plc Form of Nonqualified Stock Option Agreement for Employees (filed as Exhibit 10.2 to STERIS plc Form 10-Q for the fiscal quarter ended December 31, 2015 (Commission File No. 1-37614) and incorporated herein by reference).*
- 10.11 STERIS plc Form of Nonqualified Stock Option Agreement for Nonemployee Directors (filed as Exhibit 10.20 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614) and incorporated herein by reference).*
- 10.12 STERIS plc Form of Nonqualified Stock Agreement for Employees (filed as Exhibit 10.16 to STERIS plc Form 10-K for the fiscal year ended March 31, 2018 (Commission File No. 1-37614) and incorporated herein by reference).*
- Amendment to STERIS plc Nonqualified Stock Option Agreement (filed as Exhibit 10.4 to STERIS plc Form 10-Q for the fiscal quarter ended September 30, 2018 (Commission File No. 1-37614) and incorporated herein by reference).*
- 10.14 Form of STERIS plc Nonqualified Stock Option Agreement for Employees (filed as Exhibit 10.2 to STERIS plc Form 10-Q for the fiscal quarter ended September 30, 2018 (Commission File No. 1-37614) and incorporated herein by reference).*
- 10.15 Form of STERIS plc Nonqualified Stock Option Agreement for Employees (filed as Exhibit 10.3 to STERIS plc Form 10-Q for the fiscal quarter ended September 30, 2019 (Commission File No. 001-38848) and incorporated herein by reference).*
- 10.16 STERIS plc Form of Restricted Stock Agreement for Employees (filed as Exhibit 10.3 to STERIS plc Form 10-Q for the fiscal quarter ended December 31, 2015 (Commission File No. 1-37614) and incorporated herein by reference).*
- 10.17 STERIS plc Form of Career Restricted Stock Agreement for Nonemployee Directors (filed as Exhibit 10.21 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614) and incorporated herein by reference).*
- 10.18 STERIS plc Form of Performance Restricted Stock Agreement for Employees (filed as Exhibit 10.1 to STERIS plc Form 8-K filed June 1, 2017 (Commission File No. 1-37614), and incorporated herein by reference).*
- 10.19 STERIS plc Form of Restricted Stock Agreement for Employees (filed as Exhibit 10.3 to STERIS plc Form 10-Q for the fiscal quarter ended September 30, 2018 (Commission File No. 1-37614), and incorporated herein by reference).*
- 10.20 Form of STERIS plc Restricted Stock Agreement for Employees (filed as Exhibit 10.2 to STERIS plc Form 10-Q for the fiscal quarter ended September 30, 2019 (Commission File No. 001-38848) and incorporated herein by reference).*
- Description of STERIS plc Non-Employee Director Compensation Program (filed as Exhibit 10.1 to STERIS plc Form 10-Q for the fiscal quarter ended September 30, 2019 (Commission File No. 001-38848) and incorporated herein by reference).*
- 10.22 STERIS Corporation Deferred Compensation Plan Document (filed as Exhibit 10.1 to Form 8-K filed September 1, 2006 (Commission File No. 1-14643), and incorporated herein by reference).*
- STERIS Corporation Deferred Compensation Plan Document (as Amended and Restated Effective January 1, 2009) (filed as Exhibit 10.1 to Form 10-Q for the fiscal quarter ended December 31, 2008 (Commission File No. 1-14643), and incorporated herein by reference).*
- Amended and Restated Adoption Agreement related to STERIS Corporation Deferred Compensation Plan (filed as Exhibit 10.2 to Form 10-Q filed for the fiscal quarter ended December 31, 2008 (Commission File No. 1-14643), and incorporated herein by reference).*
- Amendment No. 1 to STERIS Corporation Deferred Compensation Plan Document (as Amended and Restated Effective January 1, 2009) dated November 4, 2011 (filed as Exhibit 10.1 to Form 10-Q for the fiscal quarter ended December 31, 2011 (Commission File No. 1-14643), and incorporated herein by reference).*

- STERIS plc Management Incentive Compensation Plan (As Amended and Restated Effective March 28, 2019) (filed as Exhibit 10.2 to STERIS plc Form 8-K filed March 28, 2019 (Commission File No. 001-38848), and incorporated herein by reference).*
- Amendment No. 1 to STERIS plc Management Incentive Compensation Plan (As Assumed, Amended and Restated Effective March 28, 2019).*
- Form of Make-Whole Payment and Repayment Conditions Agreement Between Former STERIS Corporation Non-Employee Directors and STERIS Corporation (filed as Exhibit 10.32 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614) and incorporated herein by reference).*
- Form of Make-Whole Payment and Repayment Conditions Agreement Between STERIS Corporation Executive Officers and STERIS Corporation (filed as Exhibit 10.33 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614) and incorporated herein by reference).*
- 10.30 STERIS plc Senior Executive Severance Plan, As Adopted effective March 28, 2019 (filed as Exhibit 10.3 to STERIS plc 8-K filed March 28, 2019 (Commission File No. 001-38848), and incorporated herein by reference).*
- Form of Indemnification Agreement between STERIS Corporation and each of its directors and certain executive officers (filed as Exhibit 10.31 to Form 10-K for the fiscal year ended March 31, 2010 (Commission File No. 1-14643), and incorporated herein by reference).
- Form of Deed of Indemnity for STERIS plc Directors and executive officers (filed as Exhibit 10.5 to STERIS plc Form 10-Q for the fiscal quarter ended December 31, 2015 (Commission File No. 1-37614), and incorporated herein by reference).
- Form of Deed of Indemnity for STERIS plc directors and executive officers (filed as Exhibit 10.4 to STERIS plc Form 8-K filed March 28, 2019 (Commission File No. 001-38848), and incorporated herein by reference).
- Agreement dated as of April 23, 2008 by and among STERIS Corporation, Richard C. Breeden, Robert H. Fields, and the Breeden Investors identified therein (filed as Exhibit 10.1 to Form 8-K filed April 24, 2008 (Commission File No. 1-14643), and incorporated herein by reference).
- 10.35 Agreement dated November 4, 2011 between STERIS Corporation and Bank of America, N.A. providing Transfer and Advised Line for Letters of Credit (filed as Exhibit 10.2 to Form 10-Q for the fiscal quarter ended December 31, 2011 (Commission File No. 1-14643), and incorporated herein by reference).
- 10.36 Credit Agreement, dated as of March 23, 2018, by and among STERIS Corporation and STERIS plc, as borrowers, various U.S. and U.K. subsidiaries of STERIS plc, as guarantors, various financial institutions, as lenders and JPMorgan Chase Bank, N.A., as Administrative Agent (filed as Exhibit 10.1 to STERIS plc Form 8-K filed March 26, 2018 (Commission File No. 1-37614), and incorporated herein by reference).
- First Amendment dated March 5, 2019 to the Credit Agreement, dated as of March 23, 2018, by and among STERIS Corporation and STERIS plc, as borrowers and guarantors, various U.S. and U.K. Subsidiaries of STERIS plc, as guarantors, various financial institutions, as lenders, and JPMorgan Chase Bank, N.A., as Administrative Agent (filed as Exhibit 10.1 to form 8-K filed March 5, 2019 (Commission File No. 001-14643), and incorporated herein by reference).
- Borrower Joinder Agreement dated March 28, 2019 among STERIS plc and Synergy Health Limited and JPMorgan Chase Bank, N.A., as Administrative Agent (filed as Exhibit 10.45 to Form 10-K filed May 30, 2019 (Commission File No. 001-38848), and incorporated herein by reference).
- Guarantor Joinder Agreement dated March 28, 2019 by STERIS plc and STERIS Emerald IE Limited in favor of JPMorgan Chase Bank, N.A., as Administrative Agent (filed as Exhibit 10.46 to Form 10-K filed May 30, 2019 (Commission File No. 001-38848),and incorporated herein by reference).
- First Amendment, dated as of March 31, 2015, to Note Purchase Agreement dated as of August 15, 2008, among STERIS Corporation and each of the institutions party thereto (filed as Exhibit 10.5 to Form 8-K filed April 2, 2015 (Commission File No. 1-14643), and incorporated herein by reference).

- Second Amendment dated as of March 5, 2019 to the Amended and Restated Note Purchase Agreement dated as of March 31, 2015, as amended by that certain First Amendment dated as of January 23, 2017, by and among STERIS Corporation and each of the purchasers listed in Schedule A thereto, (filed as Exhibit 10.2 to Form 8-K filed March 5, 2019 (Commission File No. 001-37614), and incorporated herein by reference).
- Affiliate Guaranty, dated as of March 31, 2015, by STERIS Corporation and each of American Sterilizer Company, Integrated Medical Systems International, Inc., STERIS Europe, Inc., STERIS Inc., United States Endoscopy Group, Inc., Isomedix Inc. and Isomedix Operations Inc., of the August 15, 2008 Note Purchase Agreements, as amended and restated, and Notes issued pursuant thereto (filed as Exhibit 10.6 to Form 8-K filed April 2, 2015 (Commission File No. 1-14643), and incorporated herein by reference).
- Guaranty Supplement dated September 9, 2015 by General Econopak, Inc. and STERIS Corporation of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation August 15, 2008 Note Purchase Agreements as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.10 to STERIS plc Form 10-Q for the fiscal quarter ending December 31, 2015 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated November 2, 2015 by Solar New US Holding Co, LLC, Solar New US Parent Co, LLC and Solar New US Acquisition Co, LLC and STERIS Corporation of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation August 15, 2008 Note Purchase Agreements, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.52 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated January 12, 2016 by Synergy Health Holdings Limited, Synergy Health Sterilisation UK Limited, Synergy Health (UK) Limited, Synergy Health Investments Limited and Synergy Health US Holdings Limited of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation August 15, 2008 Note Purchase Agreements, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.53 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated August 8, 2017 by Synergy Health AST, LLC, Synergy Health US Holdings, Inc., and Synergy Health North America, Inc. of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation August 15, 2008 Note Purchase Agreements, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.2 to STERIS plc Form 10-Q for the fiscal quarter ending September 30, 2017 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated March 28, 2019 by STERIS plc and STERIS Emerald IE Limited and STERIS Corporation of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation August 15, 2008 Note Purchase Agreements, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.54 to Form 10-K filed May 30, 2019 (Commission File No. 001-38848), and incorporated herein by reference).
- First Amendment, dated as of March 31, 2015, to Note Purchase Agreements dated as of December 4, 2012, among STERIS Corporation and each of the institutions party thereto (filed as Exhibit 10.7 to Form 8-K filed April 2, 2015 (Commission File No. 1-14643), and incorporated herein by reference).
- Second Amendment dated as of March 5, 2019 to the Amended and Restated Note Purchase Agreement dated as of March 31, 2015, as amended by that certain First Amendment dated as of January 23, 2017, by and among STERIS Corporation and each of the purchasers listed in Schedule A thereto, (filed as Exhibit 10.3 to Form 8-K filed March 5, 2019 (Commission File No. 1-37614), and incorporated herein by reference).
- Affiliate Guaranty, dated as of March 31, 2015, by STERIS Corporation and each of American Sterilizer Company, Integrated Medical Systems International, Inc., STERIS Europe, Inc., STERIS Inc., United States Endoscopy Group, Inc., Isomedix Inc. and Isomedix Operations Inc., of the December 4, 2012 Note Purchase Agreements, as amended and restated, and Notes issued pursuant thereto (filed as Exhibit 10.8 to Form 8-K filed April 2, 2015 (Commission File No. 1-14643), and incorporated herein by reference).

- Guaranty Supplement dated September 9, 2015 by General Econopak, Inc. and STERIS Corporation of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation December 4, 2012 Note Purchase Agreements, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.11 to STERIS plc Form 10-Q for the fiscal quarter ended December 31, 2015 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated November 2, 2015 by Solar New US Holding Co, LLC, Solar New US Parent Co, LLC and Solar New US Acquisition Co, LLC and STERIS Corporation of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation December 4, 2012 Note Purchase Agreements, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.57 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated January 12, 2016 by Synergy Health Holdings Limited, Synergy Health Sterilisation UK Limited, Synergy Health (UK) Limited, Synergy Health Investments Limited and Synergy Health US Holdings Limited of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation December 4, 2012 Note Purchase Agreements, as amended and restated and of the Notes issued pursuant thereto (filed as Exhibit 10.58 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated August 8, 2017 by Synergy Health AST, LLC, Synergy Health US Holdings, Inc., and Synergy Health North America, Inc. of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation December 4, 2012 Note Purchase Agreements, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.3 to STERIS plc Form 10-Q for the fiscal quarter ending September 30, 2017 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated March 28, 2019 by STERIS plc and STERIS Emerald IE Limited and STERIS Corporation of Affiliate Guaranty dated as of March 31, 2015 of STERIS Corporation December 4, 2012 Note Purchase Agreements, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.62 to Form 10-K filed May 30, 2019 (Commission File No. 001-38848), and incorporated herein by reference).
- Note Purchase Agreement dated as of May 15, 2015, among STERIS Corporation and each of the institutions party thereto (filed as Exhibit 10.1 to Form 8-K of STERIS Corporation filed May 18, 2015 (Commission File No. 1-14643), and incorporated herein by reference).
- Second Amendment dated as of March 5, 2019 to the Note Purchase Agreement dated as of May 15, 2015, as amended by that certain First Amendment dated as of January 23, 2017, by and among STERIS Corporation and each of the purchasers listed in Schedule A thereto, (filed as Exhibit 10.4 to Form 8-K filed March 5, 2019 (Commission File No. 1-37614), and incorporated herein by reference).
- Affiliate Guaranty, dated as of May 15, 2015, by STERIS Corporation and each of American Sterilizer Company, Integrated Medical Systems International, Inc., STERIS Europe, Inc., STERIS Inc., United States Endoscopy Group, Inc., Isomedix Inc. and Isomedix Operations Inc., of STERIS Corporation May 15, 2015 Note Purchase Agreement and Notes issued pursuant thereto (filed as Exhibit 10.2 to Form 8-K of STERIS Corporation filed May 18, 2015 (Commission File No. 1-14643), and incorporated herein by reference).
- Guaranty Supplement dated September 9, 2015 by General Econopak, Inc. and STERIS Corporation of Affiliate Guaranty dated as of May 15, 2015 of STERIS Corporation May 15, 2015 Note Purchase Agreement and of the Notes issued pursuant thereto (filed as Exhibit 10.12 to STERIS plc Form 10-Q for the fiscal quarter ended December 31, 2015 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated November 2, 2015 by Solar New US Holding Co, LLC, Solar New US Parent Co, LLC and Solar New US Acquisition Co, LLC and STERIS Corporation of Affiliate Guaranty dated as of May 15, 2015 of STERIS Corporation May 15, 2015 Note Purchase Agreement and of the Notes issued pursuant thereto (filed as Exhibit 10.62 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614), and incorporated herein by reference).

- Guaranty Supplement dated January 12, 2016 by Synergy Health Holdings Limited, Synergy Health Sterilisation UK Limited, Synergy Health (UK) Limited, Synergy Health Investments Limited and Synergy Health US Holdings Limited of STERIS Corporation May 15, 2015 Note Purchase Agreement and of the Notes issued pursuant thereto (filed as Exhibit 10.63 to STERIS plc Form 10-K for the year ended March 31, 2016 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated August 8, 2017 by Synergy Health AST, LLC, Synergy Health US Holdings, Inc., and Synergy Health North America, Inc. of STERIS Corporation May 15, 2015 Note Purchase Agreement and of the Notes issued pursuant thereto (filed as Exhibit 10.4 to STERIS plc Form 10-Q for the fiscal quarter ending September 30, 2017 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated March 28, 2019 by STERIS plc and STERIS Emerald IE Limited and STERIS Corporation of Affiliate Guaranty dated as of May 15, 2015 of STERIS Corporation May 15, 2015 Note Purchase Agreement, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.70 to Form 10-K filed May 30, 2019 (Commission File No. 001-38848), and incorporated herein by reference).
- Note Purchase Agreement dated as of January 23, 2017, among STERIS plc and each of the institutions party thereto (filed as Exhibit 10.1 to Form 8-K filed January 26, 2017 (Commission File No. 1-37614), and incorporated herein by reference).
- First Amendment dated as of March 5, 2019 to the Note Purchase Agreement dated as of January 23, 2017, by and among STERIS plc and each of the purchasers listed in Schedule A thereto, (filed as Exhibit 10.5 to Form 8-K filed March 5, 2019 (Commission File No. 1-37614), and incorporated herein by reference).
- Affiliated Guaranty, dated as of January 23, 2017, by STERIS plc and each of the American Sterilizer Company, Integrated Medical Systems International, Inc., Isomedix Inc., Isomedix Operations Inc., Solar New US Holding Co, LLC, Solar New US Parent Co, LLC, Solar US Acquisition Co, LLC, STERIS Barrier Products Solutions, Inc., STERIS Corporation, STERIS Europe, Inc., STERIS Inc., Synergy Health Holdings Limited, Synergy Health Limited, Synergy Health Sterilisation UK Limited, Synergy Health (UK) Limited, Synergy Health Investments Limited, Synergy Health US Holdings Limited, and United States Endoscopy Group, Inc., of STERIS plc January 23, 2017 Note Purchase Agreement and Notes issued pursuant thereto (filed as Exhibit 10.2 to Form 8-K filed January 26, 2017 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated August 8, 2017 by Synergy Health AST, LLC, Synergy Health US Holdings, Inc. and Synergy Health North America, Inc., of Affiliate Guaranty dated as January 23, 2017 of STERIS plc January 23, 2017 Note Purchase Agreement, and of the Notes issued pursuant thereto (filed as Exhibit 10.5 to STERIS plc Form 10-Q for the fiscal quarter ending September 30, 2017 (Commission File No. 1-37614), and incorporated herein by reference).
- Guaranty Supplement dated March 28, 2019 by STERIS plc and STERIS Emerald IE Limited and STERIS Limited of Affiliate Guaranty dated as of January 23, 2017 of STERIS plc January 23, 2017 Note Purchase Agreement, as amended and restated, and of the Notes issued pursuant thereto (filed as Exhibit 10.75 to Form 10-K filed May 30, 2019 (Commission File No. 001-38848), and incorporated herein by reference).
- 10.69 Stock Purchase Agreement dated July 16, 2012 by and among STERIS Corporation, United States Endoscopy Group, Inc. and the shareholders party thereto (filed as Exhibit 2.1 to Form 8-K filed August 15, 2012 (Commission File No. 1-14643), and incorporated herein by reference).
- 10.70 Stock Purchase Agreement dated March 31, 2014 by and among STERIS Corporation, Integrated Medical Systems International, Inc. and the shareholders party thereto (filed as Exhibit 2.1 to Form 8-K filed May 9, 2014 (Commission File No. 1-14643), and incorporated herein by reference).
- 10.71 Stock Purchase Agreement dated June 23, 2015 by and among STERIS Corporation, General Econopak, Inc. and each of the Stockholders of General Econopak, Inc. (filed as Exhibit 10.1 to STERIS Corporation Form 10-Q for the fiscal quarter ended June 30, 2015 (Commission File No. 1-14643), and incorporated herein by reference).
- 21.1 Subsidiaries of STERIS plc.
- 23.1 Consent of Independent Registered Public Accounting Firm.

24.1	Power of Attorney		
31.1	Certification of the Principal Executive Officer Pursuant to Exchange Act Rule 13a-14(a)/15d-14(a).		
31.2	Certification of the Principal Financial Officer Pursuant to Exchange Act Rule 13a-14(a)/15d-14(a).		
32.1	Certification of the Principal Executive Officer and the Principal Financial Officer Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.		
101.SCH	Inline Schema Document.		
101.CAL	Inline Calculation Linkbase Document.		
101.DEF	Inline Definition Linkbase Document.		
101.LAB	Inline Labels Linkbase Document.		
101.PRE	Inline Presentation Linkbase Document.		
104	Cover Page Interactive Data File (formatted as Inline XBRL and contained in Exhibits 101).		
*	A management contract or compensatory plan or arrangement required to be filed as an exhibit hereto.		

SIGNATURES

Pursuant to the requirements of Sections 13 or 15(d) of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized, on the date indicated.

STERIS plc (Registrant)

Date: May 29, 2020 By: /s

By: /S/ KAREN L. BURTON

Karen L. Burton

Vice President, Controller, and Chief Accounting Officer

Pursuant to the requirements of the Securities Exchange Act of 1934, this report has been signed below by the following persons on behalf of the Registrant and in the capacities and on the date indicated.

<u>SIGNATURE</u>	TITLE	<u>DATE</u>
/S/ WALTER M ROSEBROUGH, JR. Walter M Rosebrough, Jr.	President, Chief Executive Officer and Director	May 29, 2020
/S/ MICHAEL J. TOKICH Michael J. Tokich	Senior Vice President and Chief Financial Officer (Principal Financial Officer)	May 29, 2020
/S/ KAREN L. BURTON	Vice President, Controller and Chief Accounting Officer	May 29, 2020
Karen L. Burton *	Chairman and Director	May 29, 2020
Mohsen M. Sohi *	Director	May 29, 2020
Richard C. Breeden *	Director	May 29, 2020
Cynthia L. Feldmann *	Director	May 29, 2020
David B. Lewis	Director	May 29, 2020
Jacqueline B. Kosecoff *		•
Nirav R. Shah	Director	May 29, 2020
* Richard M. Steeves	Director	May 29, 2020

^{*} The undersigned, by signing his name hereto, does sign and execute this Annual Report on Form 10-K pursuant to the Powers of Attorney executed by the above-named directors of the Registrant and filed with the Securities and Exchange Commission on behalf of such directors.

Date: May 29, 2020 By: /S/ J. ADAM ZANGERLE

J. Adam Zangerle,

Attorney-in-Fact for Directors

SUBSIDIARIES OF STERIS PLC

STERIS plc has no parent company. As of March 31, 2020, its direct and indirect subsidiaries⁽¹⁾ were as follows:

Albert Browne Limited England & Wales
American Sterilizer Company Pennsylvania

Bioster Mottahedoon Egypt SAE

Birkova Products

Bizworth Gammarad Sdn Bhd

Black Diamond Video, Inc.

Bryton Corporation

Egypt

Indiana

California

Indiana

CLBV Limited England & Wales

Controlled Environment Certification Services, Inc.

Ohio

Diagmed Healthcare Limited England & Wales
Dover UK I Limited England & Wales
Dover UK II Limited England & Wales
Dover UK III Limited England & Wales

Electron Beam Sdn Bhd Malaysia

Eschmann Holdings Limited England & Wales

Genii, Inc.

Minnesota

Harwell Dosimeters Limited England & Wales

Herotron E-Beam Service GmbH

Hungaroptics kft

Isomedix Inc.

Belaware

Isomedix Operations Inc.

Delaware

Isotron Limited England & Wales

Medisafe America, L.L.C. Florida

Medisafe Holdings LimitedEngland & WalesMedisafe UK LimitedEngland & Wales

PeriOptimum, Inc.

SATYAtek S.A.

Switzerland

Sercon Indústria E Comércio De Aparelhos Médicos E Hospitalares Ltda.

Brazil

Shiloh Limited England & Wales Shiloh Properties Limited England & Wales

Solar New US Holding Co, LLC

Solar New US Parent Co, LLC

Solar US Acquisition Co, LLC

Delaware

STE Hong Kong Limited

Hong Kong

STE UK HoldCo Limited

England & Wales

STE UK Sub HoldCo Limited

England & Wales

STE No. Two Corporation Delaware

Sterile Supplies Limited England & Wales

STERIS AB Sweden

Canada STERIS Applied Sterilization Technologies ULC STERIS Asia Pacific, Inc. Delaware STERIS AST CZ s.r.o. Czech Republic STERIS AST d.o.o. Slovenia STERIS AST SK s.r.o. Slovakia STERIS Barrier Products Solutions, Inc. Pennsylvania STERIS Brazil Holdings, LLC Delaware STERIS (BVI) I Limited British Virgin Islands STERIS Canada Sales ULC Canada STERIS Canada ULC Canada STERIS CH Limited England & Wales STERIS China Holdings Limited Hong Kong **STERIS** Corporation Ohio Costa Rica STERIS Corporation de Costa Rica, S.A. STERIS Deutschland GmbH Germany STERIS Dover AST Holdings Limited England & Wales STERIS Dover Canada Holdings Limited England & Wales STERIS Dover Limited England & Wales Ireland STERIS Emerald IE Limited STERIS Enterprises LLC Russia STERIS Europe, Inc. Delaware STERIS FinCo S.à r.l. Luxembourg STERIS FinCo II S.à r.l. Luxembourg STERIS GmbH Switzerland STERIS Holdings B.V. Netherlands STERIS Iberia, S.A. Spain STERIS IMS Canada Inc. Canada STERIS IMS Limited England & Wales STERIS Inc. Delaware India STERIS (India) Private Limited STERIS Instrument Management Services, Inc. Delaware STERIS Ireland Limited Ireland Ireland STERIS Irish FinCo Unlimited Company Ireland STERIS Irish FinCo II Unlimited Company STERIS Isomedix Puerto Rico, LLC Puerto Rico STERIS Japan Inc. Japan STERIS LLC Delaware STERIS Laboratories, Inc. Minnesota STERIS Latin America, Inc. Delaware STERIS Luxembourg Finance S.à r.l. Luxembourg STERIS Luxembourg Holding S.à r.l. Luxembourg

Republic of Mauritius

Mexico

STERIS Mauritius Limited

STERIS Mexico, S. de R.L. de C.V.

STERIS NV Belgium STERIS Personnel Services, Inc. Delaware STERIS Personnel Services Mexico, S. de R.L. de C.V. Mexico STERIS S.r.l. Italy STERIS SAS France STERIS SEA Sdn. Bhd. Malaysia Korea STERIS Solutions Korea Limited STERIS Solutions S. de R.L. de C.V. Mexico STERIS (Shanghai) Trading Co., Ltd. China STERIS Singapore Pte Ltd Singapore STERIS Solutions Limited England & Wales STERIS Solutions Pte. Limited Singapore STERIS S.p.A. Italy STERIS TOMOE (Thailand) Ltd. Thailand STERIS UK Holding Limited England & Wales STERIS-Austar Pharmaceutical Systems Hong Kong Limited Hong Kong STERIS-Austar Pharmaceutical Systems (Shanghai) Limited China Strategic Technology Enterprises, Inc. Delaware Synergy Health Allershausen GmbH Germany Synergy Health Amsterdam B.V. The Netherlands Synergy Health AST, LLC Delaware Synergy Health AST S.r.l. Costa Rica Synergy Health Däniken AG Switzerland Synergy Health Ede B.V. The Netherlands Synergy Health France SAS France Synergy Health Holding B.V. The Netherlands Synergy Health Holdings Limited England & Wales Synergy Health Investments Limited England & Wales Synergy Health Ireland Limited Ireland Synergy Health Limited England & Wales The Netherlands Synergy Health Logistics B.V. Synergy Health Marseille SAS France Synergy Health Nederland B.V. The Netherlands Synergy Health Radeberg GmbH Germany Synergy Health Sterilisation UK Limited England & Wales

China Synergy Health (Suzhou) Limited

China STERIS Sterilization Technologies (Suzhou) Ltd.

Synergy Health Systems Limited England & Wales

Thailand Synergy Health (Thailand) Limited Synergy Health True North, LLC New York

Synergy Health (UK) Limited England & Wales

Synergy Health US Holdings, Inc. Delaware

Synergy Health Utrecht B.V. The Netherlands Synergy Health Westport Limited

Synergy Sterilisation KL (M) Sdn Bhd

Synergy Sterilisation Kulim (M) Sdn Bhd

Synergy Sterilisation (M) Sdn Bhd

Synergy Sterilisation (M) Sdn Bhd

Synergy Sterilisation Rawang (M) Sdn Bhd

Synergy Sterilisation South Africa (Proprietary) Limited

Ireland

Malaysia

Malaysia

Synergy Sterilisation South Africa (Proprietary) Limited

South Africa

United States Endoscopy Group, Inc.

Vernon and Co. LimitedEngland & WalesVernon-Carus LimitedEngland & Wales

Ohio

The names of one or more subsidiaries which, considered in the aggregate as a single subsidiary, would not constitute at the end of fiscal 2020 a "significant subsidiary" within the meaning of Rule 1-02(w) of Regulation S-X have been excluded.

Consent of Independent Registered Public Accounting Firm

We consent to the incorporation by reference in the following Registration Statements:

- (1) Registration Statement (Form S-8, No. 333-230557) of STERIS plc pertaining to the STERIS Corporation 401(k) Plan, and
- (2) Registration Statement (Form S-8, No. 333-230558) of STERIS plc pertaining to the STERIS plc 2006 Long-Term Equity Incentive Plan (As Assumed, Amended and Restated Effective March 28, 2019);

of our reports dated May 29, 2020, with respect to the consolidated financial statements and schedule of STERIS plc and subsidiaries (STERIS) and the effectiveness of internal control over financial reporting of STERIS included in this Annual Report (Form 10-K) of STERIS for the year ended March 31, 2020.

/s/ Ernst & Young LLP

Cleveland, Ohio May 29, 2020

STERIS PLC POWER OF ATTORNEY FORM 10-K

Each of the undersigned hereby makes, constitutes, and appoints Walter M Rosebrough, Jr., Michael J. Tokich, Karen L. Burton, J. Adam Zangerle, Ronald E. Snyder, Julia Kipnis, and each of them, his or her true and lawful attorney, with full power of substitution, for and in his or her name, place, and stead, to affix, as attorney-in-fact, his or her signature in any and all capacities, to the Annual Report on Form 10-K of STERIS plc for its fiscal year ended March 31, 2020, and any and all amendments thereto to be filed with the Securities and Exchange Commission, Washington, D.C., under the provisions of the Securities Exchange Act of 1934, as amended, with power to file said Form 10-K and such amendments, and any and all other documents that may be required in connection therewith, with the Securities and Exchange Commission, hereby granting unto said attorneys-in-fact, and each of them, full power and authority to do and perform any and all acts and things requisite or appropriate in connection therewith, as fully to all intents and purposes as he or she might or could do in person, hereby ratifying and confirming all that said attorneys-in-fact or any of them may lawfully do or cause to be done by virtue hereof.

IN WITNESS WHEREOF, the undersigned have executed this Power of Attorney as of the 30th day of April, 2020.

/s/ RICHARD C. BREEDEN	/s/ CYNTHIA L. FELDMANN				
Richard C. Breeden, Director	Cynthia L. Feldmann, Director				
/s/ JACQUELINE B. KOSECOFF	/s/ DAVID B. LEWIS				
Jacqueline B. Kosecoff, Director	David B. Lewis, Director				
/s/ MOHSEN M. SOHI	/s/ RICHARD M. STEEVES				
Mohsen M. Sohi, Director	Richard M. Steeves, Director				
/s/ NIRAV R. SHAH	/s/ WALTER M ROSEBROUGH, JR				
Nirav R. Shah, Director	Walter M Rosebrough, Jr.				
	President and Chief Executive Officer				
	(Principal Executive Officer), Director				
	/s/ MICHAEL J. TOKICH				
	Michael J. Tokich				
	Senior Vice President, and Chief Financial Officer				
	(Principal Financial Officer)				
	/s/ KAREN L. BURTON				
	Karen L. Burton				
	Vice President and Controller				

(Controller and Principal Accounting Officer)

CERTIFICATION OF THE PRINCIPAL EXECUTIVE OFFICER

- I, Walter M Rosebrough, Jr., certify that:
 - 1. I have reviewed this annual report on Form 10-K of STERIS plc;
 - 2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
 - 3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
 - 4. The registrant's other certifying officer and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be
 designed under our supervision, to ensure that material information relating to the registrant, including its
 consolidated subsidiaries, is made known to us by others within those entities, particularly during the period
 in which this report is being prepared;
 - b. Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
 - c. Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
 - d. Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
 - 5. The registrant's other certifying officer and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - a. All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b. Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: May 29, 2020

/s/ Walter M Rosebrough, Jr.

CERTIFICATION OF THE PRINCIPAL FINANCIAL OFFICER

I, Michael J. Tokich, certify that:

- 1. I have reviewed this annual report on Form 10-K of STERIS plc;
- 2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
- 3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
- 4. The registrant's other certifying officer and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be
 designed under our supervision, to ensure that material information relating to the registrant, including its
 consolidated subsidiaries, is made known to us by others within those entities, particularly during the period
 in which this report is being prepared;
 - Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
 - c. Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
 - d. Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
- 5. The registrant's other certifying officer and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b. Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: May 29, 2020

/s/ Michael J. Tokich

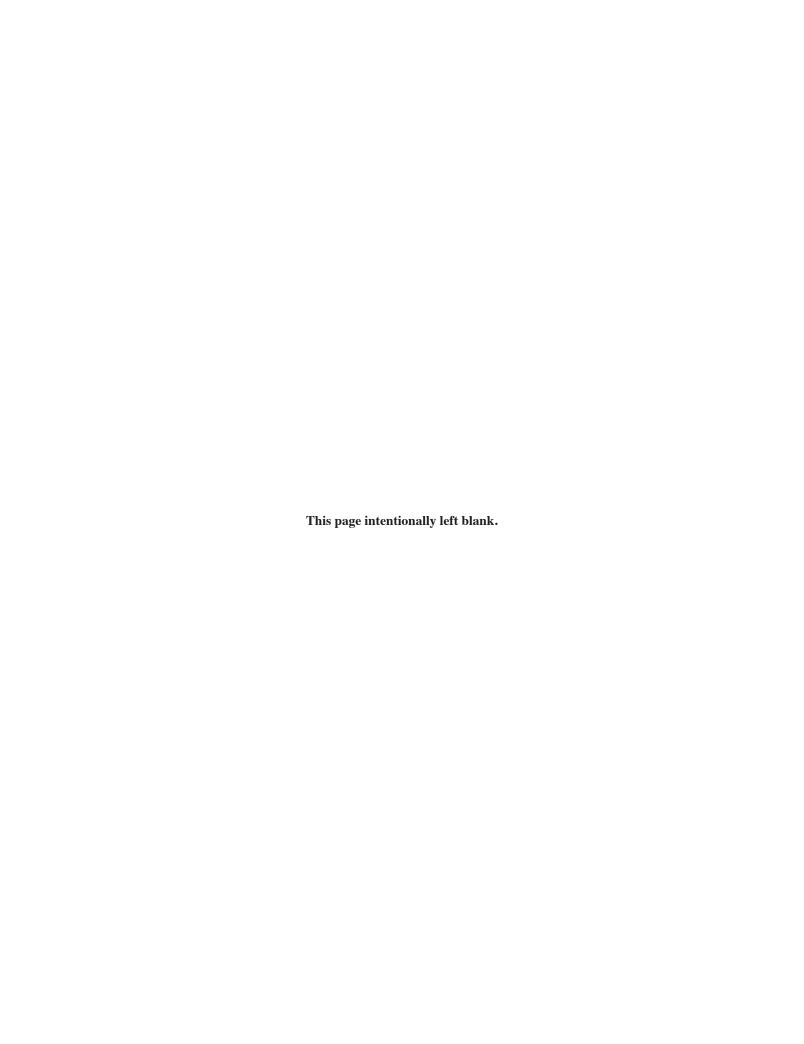
Certification Pursuant to § 906 of the Sarbanes-Oxley Act of 2002

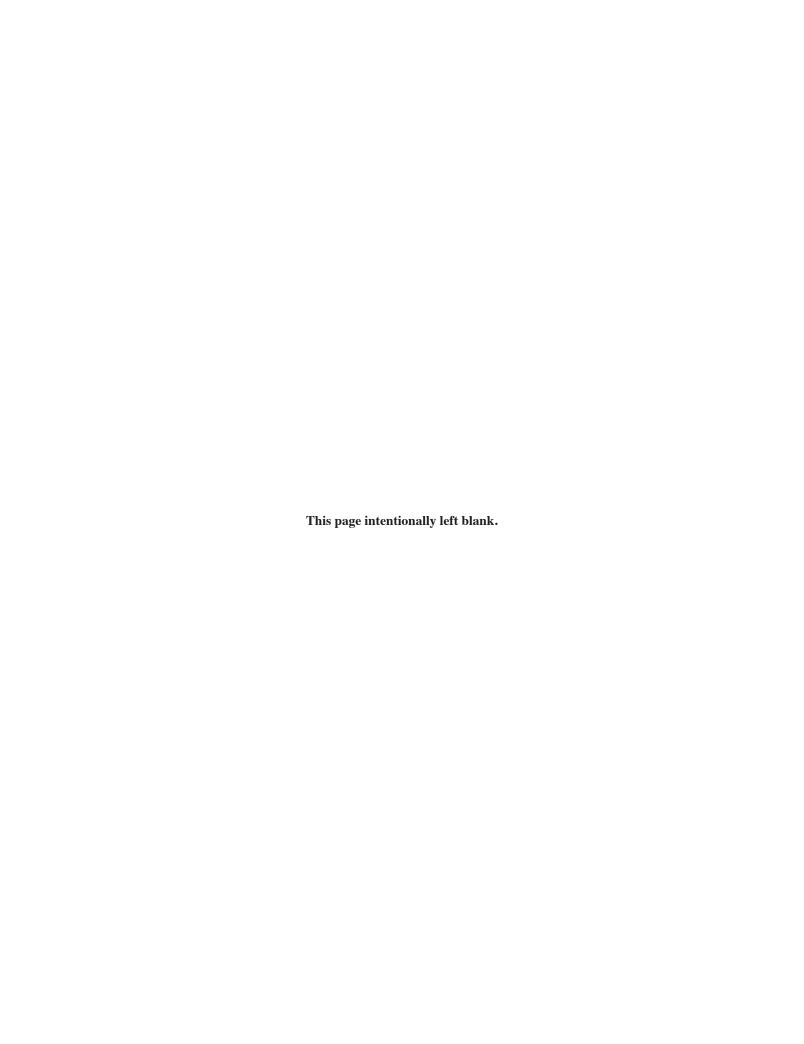
Pursuant to 18 U.S.C. § 1350, as adopted pursuant to § 906 of the Sarbanes-Oxley Act of 2002, in connection with the filing of the Form 10-K of STERIS plc (the "Company") for the fiscal year ended March 31, 2020, as filed with the Securities and Exchange Commission on the date hereof (the "Report"), each of the undersigned officers of the Company certifies, that, to such officer's knowledge:

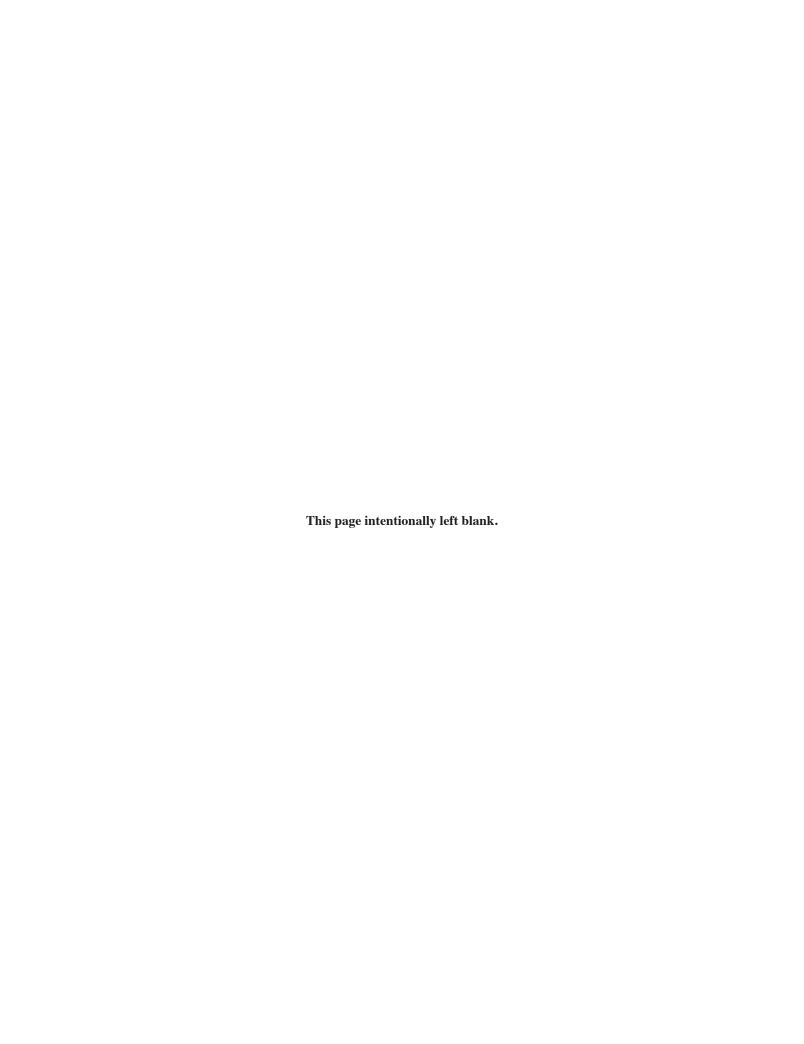
- (1) The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and
- (2) The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company as of the dates and for the periods expressed in the Report.

	/s/ Walter M Rosebrough, Jr.					
Name: Title:	Walter M Rosebrough, Jr. President and Chief Executive Officer					
	/s/ Michael J. Tokich					
Name: Title:	Michael J. Tokich Senior Vice President and Chief Financial Officer					

Dated: May 29, 2020







This Page is Not Part of STERIS plc's Form 10-K Filing

(In thousands, except per share data)

Non-GAAP Financial Measures. Non-GAAP financial measures are presented with the intent of providing greater transparency to supplemental financial information used by management and the Board of Directors in their financial analysis and operational decision making. These amounts are disclosed so that the reader has the same financial data that management uses with the belief that it will assist investors and other readers in making comparisons to our historical operating results and analyzing the underlying performance of our operations for the periods presented.

Management and the Board of Directors believe that the presentation of these non-GAAP financial measures, when considered along with our GAAP financial measures and the reconciliation to the corresponding GAAP financial measures, provide the reader with a more complete understanding of the factors and trends affecting our business than could be obtained absent this disclosure. It is important for the reader to note that the non-GAAP financial measure used may be calculated differently from, and therefore may not be comparable to, a similarly titled measure used by other companies.

Twelve months ended March 31, (unaudited)

	As reported, GAAP		Impact of Acquisitions		Impact of Divestitures		Impact of Foreign Currency Movements		GAAP Growth	Organic Growth	Constant Currency Organic Growth
	2020	2019		2020		2019		2020	2020	2020	2020
Segment revenues:											
Healthcare Products	\$ 1,423,198	\$ 1,338,428	\$	_	\$	_	\$	(4,794)	6.3%	6.3%	6.7%
Healthcare Specialty Services	563,611	510,057		_		(4,485)		(3,874)	10.5%	11.5%	12.2%
Life Sciences	416,939	378,558		_		_		(3,152)	10.1%	10.1%	11.0%
Applied Sterilization Technologies	627,147	555,127		_		_		(8,542)	13.0%	13.0%	14.5%
Total	\$ 3,030,895	\$ 2,782,170	\$		\$	(4,485)	\$	(20,362)	8.9%	9.1%	9.8%

To measure the percentage organic revenue growth, the Company removes the impact of acquisitions and divestitures that affect the comparability and trends in revenue. To measure the percentage constant currency organic revenue growth, the impact of changes in currency exchange rates and acquisitions and divestitures that affect the comparability and trends in revenue are removed. The impact of changes in currency exchange rates is calculated by translating current year results at prior year average currency exchange rates.

Twelve months ended March 31, (unaudited) Net Income Income from attributable to **Gross Profit Operations** Diluted EPS shareholders 2020 2019 2020 2019 2020 2019 2020 2019 GAAP \$ 1,319,923 \$1,175,427 \$536,973 \$ 411,465 \$407,605 \$ 304,051 4.76 3.56 Adjustments: Amortization of acquired intangible assets 1,566 721 71,675 86,878 8,225 8.901 Acquisition and integration related charges 1,583 2,145 Redomiciliation and tax restructuring costs 3,699 8,783 (Gain) on fair value adjustment of acquisition (842)related contingent consideration 1,770 (1,370)Net loss (gain) on divestiture of businesses Amortization of property "step up" to fair value 2,499 2,604 2,392 2,440 COVID-19 incremental costs 475 749 Restructuring charges 2,470 9,721 3,143 40,708 Net impact of adjustments after tax* 75,019 113,497 Net EPS impact 0.88 1.33 \$ 1,328,516 \$1,190,618 \$628,626 \$ 556,963 \$482,624 \$ 417,548 5.64 4.89 Adjusted \$

^{*} The tax expense includes both the current and deferred income tax impact of the adjustments.

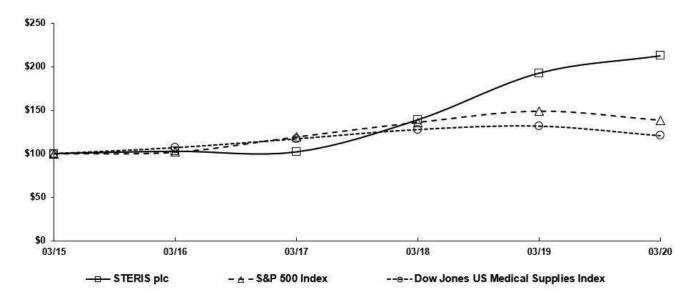
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The following table presents a financial measure which is considered to be "non-GAAP financial measures" under Securities Exchange Commission rules. Free cash flow is defined by the Company as cash flows from operating activities less purchases of property, plant, equipment and intangibles (capital expenditures) plus proceeds from the sale of property, plant, equipment and intangibles. The Company uses free cash flow as a measure to gauge its ability to pay cash dividends, fund growth outside of core operations, fund future debt principal repayments, and repurchase shares.

	Tv	Twelve Months Ended March 31,				
		2020	2019 Unaudited)			
	J)	Unaudited)				
Calculation of Free Cash Flow:						
Cash flows from operating activities	\$	590,559 \$	539,505			
Purchases of property, plant, equipment, and intangibles, net		(214,516)	(189,715)			
Proceeds from the sale of property, plant, equipment, and intangibles		4,156	5,567			
Free Cash Flow	\$	380,199 \$	355,357			

Performance Graph. The following graph shows the cumulative performance for our ordinary shares over the last five years as of March 31 of each year compared with the performance of the Standard & Poor's 500 Index and the Dow Jones U.S. Medical Supplies Index as of the same date. The graph assumes \$100 invested as of March 31, 2015 in our ordinary shares and in each of the named indices. The past performance shown in this graph does not necessarily guarantee future performance.

COMPARISON OF 5 YEAR CUMULATIVE TOTAL RETURN*



^{*\$100} invested on 3/31/15 in stock or index, including reinvestment of dividends. Fiscal year ending March 31.

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	3/15	3/16	3/17	3/18	3/19	3/20
STERIS plc	100.00	102.60	101.89	138.89	192.71	212.71
S&P 500 Index	100.00	101.78	119.26	135.95	148.86	138.47
Dow Jones US Medical Supplies Index	100.00	106.94	117.19	128.01	132.00	120.74

Corporate Information

BOARD OF DIRECTORS

Dr. Mohsen M. Sohi

Chairman of the Board, STERIS plc Chief Executive Officer, Freudenberg and Co.

Richard C. Breeden 1,2,4

Chairman and Chief Executive Officer, Breeden Capital Management LLC; Chairman, Richard C. Breeden & Co., LLC

Cynthia L. Feldmann 2,3

Former President and Founder, Jetty Lane Associates

Dr. Jacqueline B. Kosecoff 1,4

Managing Partner, Moriah Partners, LLC

David B. Lewis 2,4

Of Counsel and Former Chairman, Lewis & Munday

Walter M Rosebrough, Jr. 3

President and Chief Executive Officer, STERIS plc

Dr. Nirav R. Shah 1,3

Senior Scholar, Clinical Excellence Research Center, Stanford University

Dr. Richard Steeves 3

Former Chief Executive Officer and Director of Synergy Health plc

- ² Audit Committee Member
- ³ Compliance Committee Member
- ⁴ Nominating and Governance Committee Member

EXECUTIVE OFFICERS

Karen L. Burton

Vice President, Controller and Chief Accounting Officer

Daniel A. Carestio

Senior Vice President and Chief Operating Officer

Cary L. Majors

Senior Vice President, North America Commercial Operations

Walter M Rosebrough, Jr.

President and Chief Executive Officer

Renato G. Tamaro

Vice President and Corporate Treasurer

Michael J. Tokich

Senior Vice President and Chief Financial Officer

J. Adam Zangerle

Senior Vice President, General Counsel and Company Secretary

REGISTERED OFFICE

STERIS plc 70 Sir John Rogerson's Quay Dublin 2 Ireland

ANNUAL REPORT

Included in this Annual Report is a copy of STERIS's Form 10-K filed with the Securities and Exchange Commission for the year ended March 31, 2020. Additional copies of the Company's Form 10-K and other information are available at www.steris-ir.com or upon written request to:

Julie Winter

Senior Director, Investor Relations STERIS 5960 Heisley Road Mentor, OH 44060-1834 USA

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INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

Ernst & Young LLP

Suite 1800 950 Main Avenue Cleveland, OH 44113-7214

STOCK EXCHANGE LISTING

STERIS is listed on the New York Stock Exchange under the symbol STE.

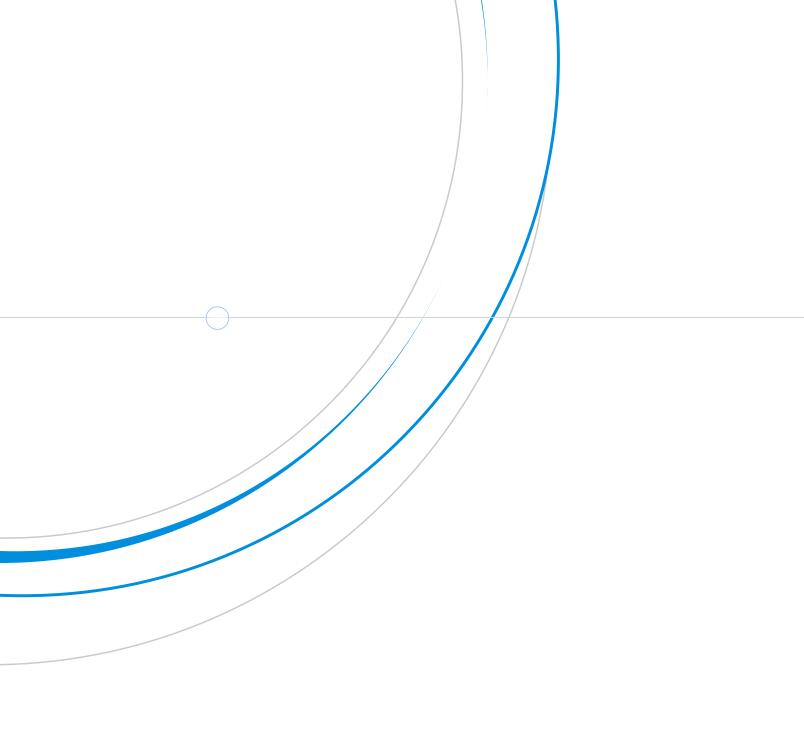
ANNUAL MEETING OF SHAREHOLDERS

The Company's 2020 annual meeting will be held Tuesday, July 28, 2020.

Portions of this Annual Report, other than the Form 10-K, have not been filed with the SEC.

Product and service descriptions and financial information herein are for illustration purposes only and do not modify or alter product warranties, labeling, instructions, or other technical literature, or the financial information contained in the Form 10-K.

¹ Compensation and Organization Development Committee Member





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