

**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549**

FORM 10-Q



**Quarterly Report Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934
for the quarterly period ended July 2, 2023**

or



**Transition Report Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934
for the transition period from to**

Commission file number 1-3215

Johnson & Johnson

(Exact name of registrant as specified in its charter)

New Jersey

(State or other jurisdiction of
incorporation or organization)

22-1024240

(I.R.S. Employer
Identification No.)

One Johnson & Johnson Plaza
New Brunswick, New Jersey 08933
(Address of principal executive offices)

Registrant's telephone number, including area code (732) 524-0400

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. ☒ Yes ☐ No

Indicate by check mark whether the registrant has submitted electronically every Interactive Data File required to be submitted pursuant to Rule 405 of Regulation S-T during the preceding 12 months (or for such shorter period that the registrant was required to submit such files). ☒ Yes ☐ No

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, a smaller reporting company, or an emerging growth company. See the definitions of "large accelerated filer," "accelerated filer," "smaller reporting company," and "emerging growth company" in Rule 12b-2 of the Exchange Act.

Large accelerated filer	<input checked="" type="checkbox"/>	Accelerated filer	<input type="checkbox"/>
Non-accelerated filer	<input type="checkbox"/>	Smaller reporting company	<input type="checkbox"/>
Emerging growth company	<input type="checkbox"/>		

If an emerging growth company, indicated by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). ☐ Yes ☒ No

SECURITIES REGISTERED PURSUANT TO SECTION 12(b) OF THE ACT

Title of each class	Trading Symbol	Name of each exchange on which registered
Common Stock, Par Value \$1.00	JNJ	New York Stock Exchange
0.650% Notes Due May 2024	JNJ24C	New York Stock Exchange
5.50% Notes Due November 2024	JNJ24BP	New York Stock Exchange
1.150% Notes Due November 2028	JNJ28	New York Stock Exchange

Indicate the number of shares outstanding of each of the issuer's classes of common stock, as of the latest practicable date.

On July 24, 2023, 2,598,969,594 shares of Common Stock, \$1.00 par value, were outstanding.

JOHNSON & JOHNSON AND SUBSIDIARIES

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CAUTIONARY NOTE REGARDING FORWARD-LOOKING STATEMENTS

This Quarterly Report on Form 10-Q and Johnson & Johnson's other publicly available documents contain "forward-looking statements" within the meaning of the safe harbor provisions of the United States Private Securities Litigation Reform Act of 1995. Management and representatives of Johnson & Johnson and its subsidiaries (the Company) also may from time to time make forward-looking statements. Forward-looking statements do not relate strictly to historical or current facts and reflect management's assumptions, views, plans, objectives and projections about the future. Forward-looking statements may be identified by the use of words such as "plans," "expects," "will," "anticipates," "estimates," and other words of similar meaning in conjunction with, among other things: discussions of future operations, expected operating results, financial performance; impact of planned acquisitions and dispositions; impact and timing of restructuring initiatives including associated cost savings and other benefits; the Company's strategy for growth; product development activities; regulatory approvals; market position and expenditures.

Because forward-looking statements are based on current beliefs, expectations and assumptions regarding future events, they are subject to uncertainties, risks and changes that are difficult to predict and many of which are outside of the Company's control. Investors should realize that if underlying assumptions prove inaccurate, or known or unknown risks or uncertainties materialize, the Company's actual results and financial condition could vary materially from expectations and projections expressed or implied in its forward-looking statements. Investors are therefore cautioned not to rely on these forward-looking statements. Risks and uncertainties include, but are not limited to:

Risks Related to Product Development, Market Success and Competition

- Challenges and uncertainties inherent in innovation and development of new and improved products and technologies on which the Company's continued growth and success depend, including uncertainty of clinical outcomes, additional analysis of existing clinical data, obtaining regulatory approvals, health plan coverage and customer access, and initial and continued commercial success;
- Challenges to the Company's ability to obtain and protect adequate patent and other intellectual property rights for new and existing products and technologies in the United States and other important markets;
- The impact of patent expirations, typically followed by the introduction of competing generic, biosimilar or other products and resulting revenue and market share losses;
- Increasingly aggressive and frequent challenges to the Company's patents by competitors and others seeking to launch competing generic, biosimilar or other products and increased receptivity of courts, the United States Patent and Trademark Office and other decision makers to such challenges, potentially resulting in loss of market exclusivity and rapid decline in sales for the relevant product sooner than expected;
- Competition in research and development of new and improved products, processes and technologies, which can result in product and process obsolescence;
- Competition to reach agreement with third parties for collaboration, licensing, development and marketing agreements for products and technologies;
- Competition based on cost-effectiveness, product performance, technological advances and patents attained by competitors; and
- Allegations that the Company's products infringe the patents and other intellectual property rights of third parties, which could adversely affect the Company's ability to sell the products in question and require the payment of money damages and future royalties.

Risks Related to Product Liability, Litigation and Regulatory Activity

- Product efficacy or safety concerns, whether or not based on scientific evidence, potentially resulting in product withdrawals, recalls, regulatory action on the part of the United States Food and Drug Administration (or international counterparts), declining sales, reputational damage, increased litigation expense and share price impact;
 - The impact, including declining sales and reputational damage, of significant litigation or government action adverse to the Company, including product liability claims and allegations related to pharmaceutical marketing practices and contracting strategies;
 - The impact of an adverse judgment or settlement and the adequacy of reserves related to legal proceedings, including patent litigation, product liability, personal injury claims, securities class actions, government investigations, employment and other legal proceedings;
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- Increased scrutiny of the healthcare industry by government agencies and state attorneys general resulting in investigations and prosecutions, which carry the risk of significant civil and criminal penalties, including, but not limited to, debarment from government business;
- Failure to meet compliance obligations in compliance agreements with governments or government agencies, which could result in significant sanctions;
- Potential changes to applicable laws and regulations affecting United States and international operations, including relating to: approval of new products; licensing and patent rights; sales and promotion of healthcare products; access to, and reimbursement and pricing for, healthcare products and services; environmental protection; and sourcing of raw materials;
- Compliance with local regulations and laws that may restrict the Company's ability to manufacture or sell its products in relevant markets, including requirements to comply with medical device reporting regulations and other requirements such as the European Union's Medical Devices Regulation;
- Changes in domestic and international tax laws and regulations, increasing audit scrutiny by tax authorities around the world and exposures to additional tax liabilities potentially in excess of existing reserves; and
- The issuance of new or revised accounting standards by the Financial Accounting Standards Board and regulations by the Securities and Exchange Commission.

Risks Related to the Company's Strategic Initiatives, Healthcare Market Trends and the Planned Separation of the Company's Consumer Health Business

- Pricing pressures resulting from trends toward healthcare cost containment, including the continued consolidation among healthcare providers and other market participants, trends toward managed care, the shift toward governments increasingly becoming the primary payers of healthcare expenses, significant new entrants to the healthcare markets seeking to reduce costs and government pressure on companies to voluntarily reduce costs and price increases;
- Restricted spending patterns of individual, institutional and governmental purchasers of healthcare products and services due to economic hardship and budgetary constraints;
- Challenges to the Company's ability to realize its strategy for growth including through externally sourced innovations, such as development collaborations, strategic acquisitions, licensing and marketing agreements, and the potential heightened costs of any such external arrangements due to competitive pressures;
- The potential that the expected strategic benefits and opportunities from any planned or completed acquisition or divestiture by the Company may not be realized or may take longer to realize than expected;
- The potential that the expected benefits and opportunities related to past and ongoing restructuring actions may not be realized or may take longer to realize than expected;
- The Company's ability to consummate the planned full separation of the Company's Consumer Health business on a timely basis or at all;
- The Company's ability to successfully separate the Company's Consumer Health business and realize the anticipated benefits from the planned full separation; and
- Kenvue, Inc.'s (Kenvue) ability to succeed as a standalone publicly traded company.

Risks Related to Economic Conditions, Financial Markets and Operating Internationally

- The risks associated with global operations on the Company and its customers and suppliers, including foreign governments in countries in which the Company operates;
 - The impact of inflation and fluctuations in interest rates and currency exchange rates and the potential effect of such fluctuations on revenues, expenses and resulting margins;
 - Potential changes in export/import and trade laws, regulations and policies of the United States and other countries, including any increased trade restrictions or tariffs and potential drug reimportation legislation;
 - The impact on international operations from financial instability in international economies, sovereign risk, possible imposition of governmental controls and restrictive economic policies, and unstable international governments and legal systems;
 - The impact of global public health crises and pandemics;
 - Changes to global climate, extreme weather and natural disasters that could affect demand for the Company's products and services, cause disruptions in manufacturing and distribution networks, alter the availability of goods and services within the supply chain, and affect the overall design and integrity of the Company's products and operations; and
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- The impact of armed conflicts and terrorist attacks in the United States and other parts of the world, including social and economic disruptions and instability of financial and other markets.

Risks Related to Supply Chain and Operations

- Difficulties and delays in manufacturing, internally, through third-party providers or otherwise within the supply chain, that may lead to voluntary or involuntary business interruptions or shutdowns, product shortages, withdrawals or suspensions of products from the market, and potential regulatory action;
- Interruptions and breaches of the Company's information technology systems or those of the Company's vendors, which could result in reputational, competitive, operational or other business harm as well as financial costs and regulatory action;
- Reliance on global supply chains and production and distribution processes that are complex and subject to increasing regulatory requirements that may adversely affect supply, sourcing and pricing of materials used in the Company's products; and
- The potential that the expected benefits and opportunities related to restructuring actions may not be realized or may take longer to realize than expected, including due to any required approvals from applicable regulatory authorities.

Investors also should carefully read the Risk Factors described in Item 1A of the Company's Annual Report on Form 10-K for the fiscal year ended January 1, 2023, for a description of certain risks that could, among other things, cause the Company's actual results to differ materially from those expressed in its forward-looking statements. Investors should understand that it is not possible to predict or identify all such factors and should not consider the risks described above to be a complete statement of all potential risks and uncertainties. The Company does not undertake to publicly update any forward-looking statement that may be made from time to time, whether as a result of new information or future events or developments.

Part I — FINANCIAL INFORMATION

Item 1 — FINANCIAL STATEMENTS

JOHNSON & JOHNSON AND SUBSIDIARIES
CONSOLIDATED BALANCE SHEETS
(Unaudited; Dollars in Millions Except Share and Per Share Data)

	July 2, 2023	January 1, 2023
ASSETS		
Current assets:		
Cash and cash equivalents (Note 4)	\$ 21,183	14,127
Marketable securities	7,322	9,392
Accounts receivable, trade, less allowances \$205 (2022, \$203)	16,777	16,160
Inventories (Note 2)	12,888	12,483
Prepaid expenses and other	2,397	3,132
Total current assets	60,567	55,294
Property, plant and equipment at cost	51,218	49,253
Less: accumulated depreciation	(30,642)	(29,450)
Property, plant and equipment, net	20,576	19,803
Intangible assets, net (Note 3)	46,246	48,325
Goodwill (Note 3)	45,440	45,231
Deferred taxes on income (Note 5)	8,779	9,123
Other assets	10,078	9,602
Total assets	\$ 191,686	187,378
LIABILITIES AND SHAREHOLDERS' EQUITY		
Current liabilities:		
Loans and notes payable	\$ 11,701	12,771
Accounts payable	10,443	11,703
Accrued liabilities	10,605	11,456
Accrued rebates, returns and promotions	15,672	14,417
Accrued compensation and employee related obligations	3,062	3,328
Accrued taxes on income (Note 5)	2,687	2,127
Total current liabilities	54,170	55,802
Long-term debt (Note 4)	33,901	26,888
Deferred taxes on income (Note 5)	3,627	6,374
Employee related obligations (Note 6)	6,461	6,767
Long-term taxes payable (Note 5)	2,536	4,306
Other liabilities	14,582	10,437
Total liabilities	\$ 115,277	110,574
Commitments and Contingencies (Note 11)		
Shareholders' equity:		
Common stock — par value \$1.00 per share (authorized 4,320,000,000 shares; issued 3,119,843,000 shares)	\$ 3,120	3,120
Accumulated other comprehensive income (loss) (Note 7)	(13,135)	(12,967)
Retained earnings and Additional paid-in capital	129,381	128,345
Less: common stock held in treasury, at cost (521,700,000 and 506,246,000 shares)	44,217	41,694
Total Johnson & Johnson shareholders' equity	75,149	76,804
Equity attributable to non-controlling interest (Note 12)	1,260	—
Total equity	\$ 76,409	76,804
Total liabilities and shareholders' equity	\$ 191,686	187,378

See Notes to Consolidated Financial Statements

JOHNSON & JOHNSON AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF EARNINGS
(Unaudited; Dollars & Shares in Millions Except Per Share Amounts)

	July 2, 2023	Fiscal Second Quarter Ended Percent to Sales	July 3, 2022	Percent to Sales
Sales to customers (Note 9)	\$ 25,530	100.0 %	\$ 24,020	100.0 %
Cost of products sold	8,212	32.2	7,919	33.0
Gross profit	17,318	67.8	16,101	67.0
Selling, marketing and administrative expenses	6,665	26.1	6,226	25.9
Research and development expense	3,829	15.0	3,703	15.4
Interest income	(369)	(1.5)	(64)	(0.3)
Interest expense, net of portion capitalized	346	1.4	38	0.2
Other (income) expense, net*	(60)	(0.2)	273	1.1
Restructuring	145	0.5	85	0.4
Earnings before provision for taxes on income	6,762	26.5	5,840	24.3
Provision for taxes on income (Note 5)	1,618	6.4	1,026	4.3
NET EARNINGS	\$ 5,144	20.1 %	\$ 4,814	20.0 %
NET EARNINGS PER SHARE (Note 8)				
Basic	\$ 1.98		\$ 1.83	
Diluted	\$ 1.96		\$ 1.80	
AVG. SHARES OUTSTANDING				
Basic	2,598.4		2,629.6	
Diluted	2,625.7		2,667.9	

*Fiscal second quarter of 2023 Other (income) expense, net includes \$37 million related to the 10.4% non-controlling interest in Kenvue from the time of the initial public offering on May 8, 2023 through the end of the fiscal second quarter.

See Notes to Consolidated Financial Statements

JOHNSON & JOHNSON AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF EARNINGS
(Unaudited; Dollars & Shares in Millions Except Per Share Amounts)

	July 2, 2023	Fiscal Six Months Ended Percent to Sales	July 3, 2022	Percent to Sales
Sales to customers (Note 9)	\$ 50,276	100.0 %	\$ 47,446	100.0 %
Cost of products sold	16,607	33.0	15,517	32.7
Gross profit	33,669	67.0	31,929	67.3
Selling, marketing and administrative expenses	12,803	25.5	12,164	25.6
Research and development expense	7,392	14.7	7,165	15.1
In-process research and development impairments	49	0.1	610	1.3
Interest income	(604)	(1.2)	(86)	(0.2)
Interest expense, net of portion capitalized	561	1.1	48	0.1
Other (income) expense, net*	7,168	14.3	171	0.4
Restructuring	275	0.5	155	0.3
Earnings before provision for taxes on income	6,025	12.0	11,702	24.7
Provision for taxes on income (Note 5)	949	1.9	1,739	3.7
NET EARNINGS	\$ 5,076	10.1 %	\$ 9,963	21.0 %
NET EARNINGS PER SHARE (Note 8)				
Basic	\$ 1.95		\$ 3.79	
Diluted	\$ 1.93		\$ 3.73	
AVG. SHARES OUTSTANDING				
Basic	2,601.9		2,629.4	
Diluted	2,630.7		2,669.2	

* Fiscal six months of 2023 Other (income) expense, net includes \$37 million related to the 10.4% non-controlling interest in Kenvue from the time of the initial public offering on May 8, 2023 through the end of the fiscal second quarter.

See Notes to Consolidated Financial Statements

JOHNSON & JOHNSON AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF COMPREHENSIVE INCOME
(Unaudited; Dollars in Millions)

	Fiscal Second Quarter Ended		Fiscal Six Months Ended	
	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022
Net earnings	\$ 5,144	4,814	\$ 5,076	9,963
Other comprehensive income (loss), net of tax				
Foreign currency translation	(715)	(151)	(896)	(705)
Securities:				
Unrealized holding gain (loss) arising during period	4	(20)	21	(33)
Reclassifications to earnings	—	—	—	—
Net change	4	(20)	21	(33)
Employee benefit plans:				
Prior service cost amortization during period	(36)	(20)	(71)	(73)
Gain (loss) amortization during period	(34)	86	(67)	303
Net change	(70)	66	(138)	230
Derivatives & hedges:				
Unrealized gain (loss) arising during period	(137)	145	433	(50)
Reclassifications to earnings	(139)	(126)	(136)	(227)
Net change	(276)	19	297	(277)
Other comprehensive income (loss)	(1,057)	(86)	(716)	(785)
Comprehensive income	\$ 4,087	4,728	\$ 4,360	9,178

See Notes to Consolidated Financial Statements

The tax effects in other comprehensive income/(loss) for the fiscal second quarter were as follows for 2023 and 2022, respectively: Foreign Currency Translation: \$32 million and \$533 million; Securities: \$1 million and \$6 million; Employee Benefit Plans: \$21 million and \$84 million; Derivatives & Hedges: \$74 million and \$5 million.

The tax effects in other comprehensive income/(loss) for the fiscal six months were as follows for 2023 and 2022, respectively: Foreign Currency Translation: \$266 million and \$678 million; Securities: \$6 million and \$9 million; Employee Benefit Plans: \$43 million and \$65 million; Derivatives & Hedges: \$80 million and \$73 million.

JOHNSON & JOHNSON AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF EQUITY
(Unaudited; Dollars in Millions)

Fiscal Second Quarter Ended July 2, 2023

	Total	Retained Earnings and Additional paid-in capital	Accumulated Other Comprehensive Income	Common Stock Issued Amount	Treasury Stock Amount	Non-controlling interest (NCI)
Balance, April 2, 2023	\$ 70,869	124,558	(12,626)	3,120	(44,183)	—
Net earnings	5,144	5,144	—	—	—	—
Cash dividends paid (\$1.19 per share)	(3,092)	(3,092)	—	—	—	—
Employee compensation and stock option plans	649	301	—	—	348	—
Repurchase of common stock	(381)	—	—	—	(381)	—
Other	(1)	—	—	—	(1)	—
Kenvue IPO	4,278	2,470	548	—	—	1,260 *
Other comprehensive income (loss), net of tax	(1,057)	—	(1,057)	—	—	—
Balance, July 2, 2023	\$ 76,409	129,381	(13,135)	3,120	(44,217)	1,260

Fiscal Six Months Ended July 2, 2023

	Total	Retained Earnings and Additional paid-in capital	Accumulated Other Comprehensive Income	Common Stock Issued Amount	Treasury Stock Amount	Non-controlling interest (NCI)
Balance, January 1, 2023	\$ 76,804	128,345	(12,967)	3,120	(41,694)	—
Net earnings	5,076	5,076	—	—	—	—
Cash dividends paid (\$2.32 per share)	(6,034)	(6,034)	—	—	—	—
Employee compensation and stock option plans	944	(476)	—	—	1,420	—
Repurchase of common stock	(3,918)	—	—	—	(3,918)	—
Other	(25)	—	—	—	(25)	—
Kenvue IPO	4,278	2,470	548	—	—	1,260 *
Other comprehensive income (loss), net of tax	(716)	—	(716)	—	—	—
Balance, July 2, 2023	\$ 76,409	129,381	(13,135)	3,120	(44,217)	1,260

*Includes \$37 million recorded in net earnings related to the 10.4% non-controlling interest of Kenvue.

Fiscal Second Quarter Ended July 3, 2022

	Total	Retained Earnings and Additional paid-in capital	Accumulated Other Comprehensive Income	Common Stock Issued Amount	Treasury Stock Amount
Balance, April 3, 2022	\$ 74,709	124,380	(13,757)	3,120	(39,034)
Net earnings	4,814	4,814	—	—	—
Cash dividends paid (\$1.13 per share)	(2,971)	(2,971)	—	—	—
Employee compensation and stock option plans	864	(7)	—	—	871
Repurchase of common stock	(973)	—	—	—	(973)
Other comprehensive income (loss), net of tax	(86)	—	(86)	—	—
Balance, July 3, 2022	\$ 76,357	126,216	(13,843)	3,120	(39,136)

Fiscal Six Months Ended July 3, 2022

	Total	Retained Earnings	Accumulated Other Comprehensive Income	Common Stock Issued Amount	Treasury Stock Amount
Balance, January 2, 2022	\$ 74,023	123,060	(13,058)	3,120	(39,099)
Net earnings	9,963	9,963	—	—	—
Cash dividends paid (\$2.19 per share)	(5,758)	(5,758)	—	—	—
Employee compensation and stock option plans	1,464	(1,049)	—	—	2,513
Repurchase of common stock	(2,550)	—	—	—	(2,550)
Other comprehensive income (loss), net of tax	(785)	—	(785)	—	—
Balance, July 3, 2022	\$ 76,357	126,216	(13,843)	3,120	(39,136)

See Notes to Consolidated Financial Statements

JOHNSON & JOHNSON AND SUBSIDIARIES
CONSOLIDATED STATEMENTS OF CASH FLOWS
(Unaudited; Dollars in Millions)

	Fiscal Six Months Ended	
	July 2, 2023	July 3, 2022
CASH FLOWS FROM OPERATING ACTIVITIES		
Net earnings	\$ 5,076	9,963
Adjustments to reconcile net earnings to cash flows from operating activities:		
Depreciation and amortization of property and intangibles	3,814	3,513
Stock based compensation	688	644
Asset write-downs	388	747
Net gain on sale of assets/businesses	(47)	(213)
Deferred tax provision	(2,342)	(2,349)
Credit losses and accounts receivable allowances	—	(3)
Changes in assets and liabilities, net of effects from acquisitions and divestitures:		
Increase in accounts receivable	(599)	(1,386)
Increase in inventories	(741)	(1,257)
Decrease in accounts payable and accrued liabilities	(1,061)	(1,170)
(Increase)/Decrease in other current and non-current assets	(1,144)	3,527
Increase/(Decrease) in other current and non-current liabilities	3,407	(2,456)
NET CASH FLOWS FROM OPERATING ACTIVITIES	7,439	9,560
CASH FLOWS FROM INVESTING ACTIVITIES		
Additions to property, plant and equipment	(1,987)	(1,470)
Proceeds from the disposal of assets/businesses, net (Note 10)	116	314
Acquisitions, net of cash acquired (Note 10)	—	(523)
Purchases of investments	(9,688)	(22,048)
Sales of investments	11,877	17,634
Credit support agreements activity, net	(798)	(10)
Other (primarily licenses and milestones)	19	(170)
NET CASH USED BY INVESTING ACTIVITIES	(461)	(6,273)
CASH FLOWS FROM FINANCING ACTIVITIES		
Dividends to shareholders	(6,034)	(5,758)
Repurchase of common stock	(3,918)	(2,550)
Proceeds from short-term debt (Note 4)	12,221	4,371
Repayment of short-term debt	(13,611)	(2,201)
Proceeds from long-term debt, net of issuance costs (Note 4)	7,674	2
Repayment of long-term debt	(501)	(2,132)
Proceeds from the exercise of stock options/employee withholding tax on stock awards, net	254	820
Credit support agreements activity, net	(126)	813
Proceeds from Kenvue initial public offering (Note 12)	4,241	—
Other	(53)	(11)
NET CASH FROM/(USED BY) FINANCING ACTIVITIES	147	(6,646)
Effect of exchange rate changes on cash and cash equivalents	(69)	(145)
Increase/(Decrease) in cash and cash equivalents	7,056	(3,504)
Cash and Cash equivalents beginning of period	14,127	14,487
CASH AND CASH EQUIVALENTS, END OF PERIOD	\$ 21,183	10,983
Acquisitions		
Fair value of assets acquired	\$ —	621
Fair value of liabilities assumed	—	(98)
Net cash paid for acquisitions	\$ —	523

See Notes to Consolidated Financial Statements

NOTES TO CONSOLIDATED FINANCIAL STATEMENTS

NOTE 1 — The accompanying unaudited interim consolidated financial statements and related notes should be read in conjunction with the audited Consolidated Financial Statements of Johnson & Johnson and its subsidiaries (the Company) and related notes as contained in the Company's Annual Report on Form 10-K for the fiscal year ended January 1, 2023. The unaudited interim financial statements include all adjustments (consisting only of normal recurring adjustments) and accruals necessary in the judgment of management for a fair statement of the results for the periods presented.

For details regarding the Kenvue separation see Note 12 to the Consolidated Financial Statements.

Columns and rows within tables may not add due to rounding. Percentages have been calculated using actual, non-rounded figures.

New Accounting Standards

The Company assesses the adoption impacts of recently issued accounting standards by the Financial Accounting Standards Board on the Company's financial statements as well as material updates to previous assessments, if any, from the Company's Annual Report on Form 10-K for the fiscal year ended January 1, 2023.

Recently Adopted Accounting Standards

ASU 2022-04: Liabilities-Supplier Finance Programs (Topic 405-50) – Disclosure of Supplier Finance Program Obligations

The Company adopted the standard as of the beginning of fiscal year 2023, which requires that a buyer in a supplier finance program disclose additional information about the program for financial statement users.

The Company has agreements for supplier finance programs with third-party financial institutions. These programs provide participating suppliers the ability to finance payment obligations from the Company with the third-party financial institutions. The Company is not a party to the arrangements between the suppliers and the third-party financial institutions. The Company's obligations to its suppliers, including amounts due, and scheduled payment dates (which have general payment terms of 90 days), are not affected by a participating supplier's decision to participate in the program.

As of July 2, 2023, and January 1, 2023, \$0.9 billion and \$1.0 billion, respectively, were valid obligations under the program. The obligations are presented as Accounts payable on the Consolidated Balance Sheets.

Recently Issued Accounting Standards**Not Adopted as of July 2, 2023**

There were no new material accounting standards issued in the fiscal six months of 2023.

Reclassification

Certain prior period amounts have been reclassified to conform to current year presentation.

NOTE 2 — INVENTORIES

(Dollars in Millions)

	July 2, 2023	January 1, 2023
Raw materials and supplies	\$ 2,419	2,070
Goods in process	1,944	1,700
Finished goods	8,525	8,713
Total inventories	\$ 12,888	12,483

NOTE 3 — INTANGIBLE ASSETS AND GOODWILL

Intangible assets that have finite useful lives are amortized over their estimated useful lives. The latest annual impairment assessment of goodwill and indefinite lived intangible assets was completed in the fiscal fourth quarter of 2022. Future impairment tests for goodwill and indefinite lived intangible assets will be performed annually in the fiscal fourth quarter, or sooner, if warranted.

(Dollars in Millions)	July 2, 2023	January 1, 2023
Intangible assets with definite lives:		
Patents and trademarks — gross	\$ 44,637	44,012
Less accumulated amortization	(24,460)	(22,266)
Patents and trademarks — net	20,177	21,746
Customer relationships and other intangibles — gross	23,055	22,987
Less accumulated amortization	(13,516)	(12,901)
Customer relationships and other intangibles — net ⁽¹⁾	9,539	10,086
Intangible assets with indefinite lives:		
Trademarks	6,834	6,807
Purchased in-process research and development	9,696	9,686
Total intangible assets with indefinite lives	16,530	16,493
Total intangible assets — net	\$ 46,246	48,325

⁽¹⁾The majority is comprised of customer relationships

Goodwill as of July 2, 2023 was allocated by segment of business as follows:

(Dollars in Millions)	Consumer Health	Pharmaceutical	MedTech	Total
Goodwill at January 1, 2023	\$ 9,184	10,184	25,863	45,231
Goodwill, related to acquisitions	—	—	—	—
Goodwill, related to divestitures	—	—	—	—
Currency translation/Other	(103)	137	175 *	209
Goodwill at July 2, 2023	\$ 9,081	10,321	26,038	45,440

*Includes purchase price allocation adjustment for Abiomed

The weighted average amortization period for patents and trademarks is 12 years. The weighted average amortization period for customer relationships and other intangible assets is 21 years. The amortization expense of amortizable intangible assets included in cost of products sold was \$1.2 billion and \$1.1 billion for the fiscal second quarters ended July 2, 2023 and July 3, 2022, respectively. The amortization expense of amortizable intangible assets included in cost of products sold was \$2.4 billion and \$2.2 billion for the fiscal six months ended July 2, 2023 and July 3, 2022, respectively. Intangible asset write-downs are included in Other (income) expense, net, with the exception of In-Process research and development which are included in the In-Process research and development impairments line.

The estimated amortization expense for approved products, before tax, for the five succeeding years is approximately:

(Dollars in Millions)	2023	2024	2025	2026	2027
	\$4,800	4,600	3,800	3,200	2,600

See Note 10 to the Consolidated Financial Statements for additional details related to acquisitions and divestitures.

NOTE 4 — FAIR VALUE MEASUREMENTS

The Company uses forward foreign exchange contracts to manage its exposure to the variability of cash flows, primarily related to the foreign exchange rate changes of future intercompany product and third-party purchases of materials denominated in a foreign currency. The Company uses cross currency interest rate swaps to manage currency risk primarily related to borrowings. Both types of derivatives are designated as cash flow hedges.

Additionally, the Company uses interest rate swaps as an instrument to manage interest rate risk related to fixed rate borrowings. These derivatives are designated as fair value hedges. The Company uses cross currency interest rate swaps and forward foreign exchange contracts designated as net investment hedges. Additionally, the Company uses forward foreign exchange contracts to offset its exposure to certain foreign currency assets and liabilities. These forward foreign exchange contracts are not designated as hedges, and therefore, changes in the fair values of these derivatives are recognized in earnings, thereby offsetting the current earnings effect of the related foreign currency assets and liabilities.

The Company does not enter into derivative financial instruments for trading or speculative purposes, or that contain credit risk related contingent features. The Company maintains credit support agreements (CSA) with certain derivative counterparties establishing collateral thresholds based on respective credit ratings and netting agreements. As of July 2, 2023, the cumulative amount of cash collateral paid by the Company under the CSA amounted to \$1.8 billion net, related to net investment and cash flow hedges. On an ongoing basis, the Company monitors counter-party credit ratings. The Company considers credit non-performance risk to be low, because the Company primarily enters into agreements with commercial institutions that have at least an investment grade credit rating. Refer to the table on significant financial assets and liabilities measured at fair value contained in this footnote for receivables and payables with these commercial institutions. As of July 2, 2023, the Company had notional amounts outstanding for forward foreign exchange contracts, cross currency interest rate swaps and interest rate swaps of \$46.9 billion, \$39.5 billion and \$10.0 billion, respectively. As of January 1, 2023, the Company had notional amounts outstanding for forward foreign exchange contracts, cross currency interest rate swaps and interest rate swaps of \$43.3 billion, \$36.2 billion and \$12.4 billion, respectively.

All derivative instruments are recorded on the balance sheet at fair value. Changes in the fair value of derivatives are recorded each period in current earnings or other comprehensive income, depending on whether the derivative is designated as part of a hedge transaction, and if so, the type of hedge transaction.

The designation as a cash flow hedge is made at the entrance date of the derivative contract. At inception, all derivatives are expected to be highly effective. Foreign exchange contracts designated as cash flow hedges are accounted for under the forward method and all gains/losses associated with these contracts will be recognized in the income statement when the hedged item impacts earnings. Changes in the fair value of these derivatives are recorded in accumulated other comprehensive income until the underlying transaction affects earnings and are then reclassified to earnings in the same account as the hedged transaction.

Gains and losses associated with interest rate swaps and changes in fair value of hedged debt attributable to changes in interest rates are recorded to interest expense in the period in which they occur. Gains and losses on net investment hedges are accounted for through the currency translation account within accumulated other comprehensive income. The portion excluded from effectiveness testing is recorded through interest (income) expense using the spot method. On an ongoing basis, the Company assesses whether each derivative continues to be highly effective in offsetting changes of hedged items. If and when a derivative is no longer expected to be highly effective, hedge accounting is discontinued.

The Company designated its Euro denominated notes issued in May 2016 with due dates ranging from 2022 to 2035 as a net investment hedge of the Company's investments in certain of its international subsidiaries that use the Euro as their functional currency in order to reduce the volatility caused by changes in exchange rates.

As of July 2, 2023, the balance of deferred net gain on derivatives included in accumulated other comprehensive income was \$67 million after-tax. For additional information, see the Consolidated Statements of Comprehensive Income and Note 7. The Company expects that substantially all of the amounts related to forward foreign exchange contracts will be reclassified into earnings over the next 12 months as a result of transactions that are expected to occur over that period. The maximum length of time over which the Company is hedging transaction exposure is 18 months, excluding interest rate contracts and net investment hedge contracts. The amount ultimately realized in earnings may differ as foreign exchange rates change. Realized gains and losses are ultimately determined by actual exchange rates at maturity of the derivative.

The following table is a summary of the activity related to derivatives and hedges for the fiscal second quarters ended July 2, 2023 and July 3, 2022, net of tax:

(Dollars in Millions)	July 2, 2023					July 3, 2022				
	Sales	Cost of Products Sold	R&D Expense	Interest (Income) Expense	Other (Income) Expense	Sales	Cost of Products Sold	R&D Expense	Interest (Income) Expense	Other (Income) Expense
The effects of fair value, net investment and cash flow hedging:										
Gain (Loss) on fair value hedging relationship:										
Interest rate swaps contracts:										
Hedged items	\$	—	—	—	(175)	—	—	—	—	(241)
Derivatives designated as hedging instruments		—	—	—	175	—	—	—	—	241
Gain (Loss) on net investment hedging relationship:										
Cross currency interest rate swaps contracts:										
Amount of gain or (loss) recognized in income on derivative amount excluded from effectiveness testing		—	—	—	33	—	—	—	—	44
Amount of gain or (loss) recognized in AOCI		—	—	—	33	—	—	—	—	44
Gain (Loss) on cash flow hedging relationship:										
Forward foreign exchange contracts:										
Amount of gain or (loss) reclassified from AOCI into income	(15)	56	(12)	—	3	(17)	(6)	42	—	(39)
Amount of gain or (loss) recognized in AOCI	(14)	251	7	—	18	(25)	35	69	—	(38)
Cross currency interest rate swaps contracts:										
Amount of gain or (loss) reclassified from AOCI into income		—	—	—	74	—	—	—	—	102
Amount of gain or (loss) recognized in AOCI	\$	—	—	—	(432)	—	—	—	—	60

The following table is a summary of the activity related to derivatives and hedges for the fiscal six months ended July 2, 2023 and July 3, 2022, net of tax:

(Dollars in Millions)	July 2, 2023					July 3, 2022				
	Sales	Cost of Products Sold	R&D Expense	Interest (Income) Expense	Other (Income) Expense	Sales	Cost of Products Sold	R&D Expense	Interest (Income) Expense	Other (Income) Expense
The effects of fair value, net investment and cash flow hedging:										
Gain (Loss) on fair value hedging relationship:										
Interest rate swaps contracts:										
Hedged items	\$	—	—	—	(1,104)	—	—	—	—	(772)
Derivatives designated as hedging instruments		—	—	—	1,104	—	—	—	—	772
Gain (Loss) on net investment hedging relationship:										
Cross currency interest rate swaps contracts:										
Amount of gain or (loss) recognized in income on derivative amount excluded from effectiveness testing		—	—	—	67	—	—	—	—	89
Amount of gain or (loss) recognized in AOCI		—	—	—	67	—	—	—	—	89
Gain (Loss) on cash flow hedging relationship:										
Forward foreign exchange contracts:										
Amount of gain or (loss) reclassified from AOCI into income	(3)	(90)	(25)	—	5	(34)	(58)	65	—	(57)
Amount of gain or (loss) recognized in AOCI	10	396	(29)	—	4	(3)	(59)	102	—	(111)
Cross currency interest rate swaps contracts:										
Amount of gain or (loss) reclassified from AOCI into income		—	—	—	182	—	—	—	—	222
Amount of gain or (loss) recognized in AOCI	\$	—	—	—	(15)	—	—	—	—	(68)

As of July 2, 2023, and January 1, 2023, the following amounts were recorded on the Consolidated Balance Sheet related to cumulative basis adjustment for fair value hedges

Line item in the Consolidated Balance Sheet in which the hedged item is included (Dollars in Millions)	Carrying Amount of the Hedged Liability		Cumulative Amount of Fair Value Hedging Gain/ (Loss) Included in the Carrying Amount of the Hedged Liability	
	July 2, 2023	January 1, 2023	July 2, 2023	January 1, 2023
Long-term Debt	\$ 8,662	8,665	(1,443)	(1,435)

The following table is the effect of derivatives not designated as hedging instruments for the fiscal second quarters ended 2023 and 2022:

(Dollars in Millions)	Location of Gain/(Loss) Recognized in Income on Derivative	Gain/(Loss) Recognized In Income on Derivative		Gain/(Loss) Recognized In Income on Derivative	
		Fiscal Second Quarter Ended		Fiscal Six Months Ended	
Derivatives Not Designated as Hedging Instruments		July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022
Foreign Exchange Contracts	Other (income) expense	\$ 33	73	2	102

The following table is the effect of net investment hedges for the fiscal second quarters ended in 2023 and 2022:

(Dollars in Millions)	Gain/(Loss) Recognized In Accumulated OCI		Location of Gain or (Loss) Reclassified from Accumulated Other Comprehensive Income Into Income	Gain/(Loss) Reclassified From Accumulated OCI Into Income	
	July 2, 2023	July 3, 2022		July 2, 2023	July 3, 2022
Debt	\$ 11	202	Interest (income) expense	—	—
Cross Currency interest rate swaps	\$ (24)	313	Interest (income) expense	—	—

The following table is the effect of net investment hedges for the fiscal six months ended in 2023 and 2022

(Dollars in Millions)	Gain/(Loss) Recognized In Accumulated OCI		Location of Gain or (Loss) Reclassified from Accumulated Other Comprehensive Income Into Income	Gain/(Loss) Reclassified From Accumulated OCI Into Income	
	July 2, 2023	July 3, 2022		July 2, 2023	July 3, 2022
Debt	\$ (66)	270	Interest (income) expense	—	—
Cross Currency interest rate swaps	\$ 666	873	Interest (income) expense	—	—

The Company holds equity investments with readily determinable fair values and equity investments without readily determinable fair values. The Company has elected to measure equity investments that do not have readily determinable fair

values at cost minus impairment, if any, plus or minus changes resulting from observable price changes in orderly transactions for the identical or a similar investment of the same issuer.

The following table is a summary of the activity related to equity investments:

(Dollars in Millions)	January 1, 2023			July 2, 2023	
	Carrying Value	Changes in Fair Value Reflected in Net Income ⁽¹⁾	Sales/Purchases/Other ⁽²⁾	Carrying Value	Non Current Other Assets
Equity Investments with readily determinable value	\$ 576	(30)	(53)	493	493
Equity Investments without readily determinable value	\$ 698	(26)	75	747	747

⁽¹⁾ Recorded in Other Income/Expense

⁽²⁾ Other includes impact of currency

Fair value is the exit price that would be received to sell an asset or paid to transfer a liability. Fair value is a market-based measurement determined using assumptions that market participants would use in pricing an asset or liability. In accordance with ASC 820, a three-level hierarchy was established to prioritize the inputs used in measuring fair value. The levels within the hierarchy are described below with Level 1 inputs having the highest priority and Level 3 inputs having the lowest.

The fair value of a derivative financial instrument (i.e., forward foreign exchange contracts, interest rate contracts) is the aggregation by currency of all future cash flows discounted to its present value at the prevailing market interest rates and subsequently converted to the U.S. Dollar at the current spot foreign exchange rate. The Company does not believe that fair values of these derivative instruments materially differ from the amounts that could be realized upon settlement or maturity, or that the changes in fair value will have a material effect on the Company's results of operations, cash flows or financial position. The Company also holds equity investments which are classified as Level 1 and debt securities which are classified as Level 2. The Company holds acquisition related contingent liabilities based upon certain regulatory and commercial events, which are classified as Level 3, whose values are determined using discounted cash flow methodologies or similar techniques for which the determination of fair value requires significant judgment or estimations.

The following three levels of inputs are used to measure fair value:

Level 1 — Quoted prices in active markets for identical assets and liabilities.

Level 2 — Significant other observable inputs.

Level 3 — Significant unobservable inputs.

The Company's significant financial assets and liabilities measured at fair value as of July 2, 2023 and January 1, 2023 were as follows:

(Dollars in Millions)	July 2, 2023			January 1, 2023	
	Level 1	Level 2	Level 3	Total	Total ⁽¹⁾
Derivatives designated as hedging instruments:					
Assets:					
Forward foreign exchange contracts	\$ —	893	—	893	629
Interest rate contracts ⁽²⁾	—	1,310	—	1,310	1,534
Total	—	2,203	—	2,203	2,163
Liabilities:					
Forward foreign exchange contracts	—	446	—	446	511
Interest rate contracts ⁽²⁾	—	3,691	—	3,691	2,778
Total	—	4,137	—	4,137	3,289
Derivatives not designated as hedging instruments:					
Assets:					
Forward foreign exchange contracts	—	55	—	55	38
Liabilities:					
Forward foreign exchange contracts	—	22	—	22	68
Other Investments:					
Equity investments ⁽³⁾	493	—	—	493	576
Debt securities ⁽⁴⁾	—	9,831	—	9,831	10,487
Other Liabilities					
Contingent consideration ⁽⁵⁾	\$ —	—	1,142	1,142	1,120

Gross to Net Derivative Reconciliation	July 2, 2023		January 1, 2023
(Dollars in Millions)			
Total Gross Assets	\$	2,258	2,201
Credit Support Agreement (CSA)		(2,119)	(2,176)
Total Net Asset		139	25
Total Gross Liabilities		4,159	3,357
Credit Support Agreement (CSA)		(3,890)	(3,023)
Total Net Liabilities	\$	269	334

Summarized information about changes in liabilities for contingent consideration for the fiscal second quarters ended July 2, 2023 and July 3, 2022 is as follows:

(Dollars in Millions)	July 2, 2023	July 3, 2022
Beginning Balance	\$ 1,120	533
Changes in estimated fair value ⁽⁶⁾	25	(88)
Additions	—	91
Payments	(3)	—
Ending Balance	\$ 1,142	536

- (1) 2022 assets and liabilities are all classified as Level 2 with the exception of equity investments of \$576 million, which are classified as Level 1 and contingent consideration of \$1,120 million, classified as Level 3.
- (2) Includes cross currency interest rate swaps and interest rate swaps.
- (3) Classified as non-current other assets.
- (4) Classified within cash equivalents and current marketable securities.
- (5) Includes \$1,137 million and \$1,116 million, classified as non-current other liabilities as of July 2, 2023 and January 1, 2023, respectively.
- (6) Ongoing fair value adjustment amounts are primarily recorded in Research and Development expense.

The Company's cash, cash equivalents and current marketable securities as of July 2, 2023 comprised:

(Dollars in Millions)	Carrying Amount	Gain/(Loss)	Estimated Fair Value	Cash & Cash Equivalents	Current Marketable Securities
Cash	\$ 4,430	—	4,430	4,430	—
Non-U.S. sovereign securities	423	—	423	423	—
U.S. reverse repurchase agreements	8,991	—	8,991	8,991	—
Corporate debt securities ⁽¹⁾	641	—	641	368	273
Money market funds	3,570	—	3,570	3,570	—
Time deposits ⁽¹⁾	619	—	619	619	—
Subtotal	18,674	—	18,674	18,401	273
Unrealized Loss					
U.S. Gov't securities	9,416	(5)	9,411	2,729	6,682
U.S. Gov't Agencies	123	(2)	121	—	121
Other sovereign securities	9	—	9	7	2
Corporate debt securities	291	(1)	290	46	244
Subtotal available for sale debt ⁽²⁾	\$ 9,839	(8)	9,831	2,782	7,049
Total cash, cash equivalents and current marketable securities	\$ 28,513	(8)	28,505	21,183	7,322

⁽¹⁾ Held to maturity investments are reported at amortized cost and gains or losses are reported in earnings.

⁽²⁾ Available for sale debt securities are reported at fair value with unrealized gains and losses reported net of taxes in other comprehensive income.

As of the fiscal year ended January 1, 2023, the carrying amount was approximately the same as the estimated fair value.

Fair value of government securities and obligations and corporate debt securities was estimated using quoted broker prices and significant other observable inputs.

The Company classifies all highly liquid investments with stated maturities of three months or less from date of purchase as cash equivalents and all highly liquid investments with stated maturities of greater than three months from the date of purchase as current marketable securities. Available for sale securities with stated maturities of greater than one year from the date of purchase are available to fund current operations and are classified as either cash equivalents or current marketable securities.

The contractual maturities of the available for sale securities as of July 2, 2023 are as follows:

(Dollars in Millions)	Cost Basis	Fair Value
Due within one year	\$ 9,823	9,815
Due after one year through five years	16	16
Due after five years through ten years	—	—
Total debt securities	\$ 9,839	9,831

Financial Instruments not measured at Fair Value:

The following financial liabilities are held at carrying amount on the consolidated balance sheet as of July 2, 2023:

(Dollars in Millions)	Carrying Amount	Estimated Fair Value
Financial Liabilities		
Current Debt	\$ 11,701	11,676
Non-Current Debt		
5.50% Notes due 2024 (500MM GBP 1.2605)	629	627
2.625% Notes due 2025	749	726
0.55% Notes due 2025	926	914
2.45% Notes due 2026	1,997	1,890
2.95% Notes due 2027	875	954
0.95% Notes due 2027	1,394	1,313
2.90% Notes due 2028	1,497	1,411
1.150% Notes due 2028 (750MM Euro 1.0873)	811	728
6.95% Notes due 2029	298	347
1.30% Notes due 2030	1,605	1,435
4.95% Debentures due 2033	498	525
4.375% Notes due 2033	854	857
1.650% Notes due 2035 (1.5B Euro 1.0873)	1,619	1,400
3.55% Notes due 2036	840	901
5.95% Notes due 2037	993	1,126
3.625% Notes due 2037	1,334	1,355
3.40% Notes due 2038	992	889
5.85% Debentures due 2038	697	783
4.50% Debentures due 2040	541	539
2.10% Notes due 2040	826	704
4.85% Notes due 2041	297	299
4.50% Notes due 2043	496	499
3.70% Notes due 2046	1,977	1,758
3.75% Notes due 2047	811	871
3.50% Notes due 2048	743	640
2.25% Notes due 2050	807	666
2.45% Notes due 2060	1,053	811
5.50% Debentures due 2025*	748	752
5.35% Debentures due 2026*	747	756
5.05% Debentures due 2028*	994	1,006
5.00% Debentures due 2030*	992	1,006
4.90% Debentures due 2033*	1,240	1,264
5.10% Debentures due 2043*	741	763
5.05% Debentures due 2053*	1,476	1,531
5.20% Debentures due 2063*	738	765
Other	66	66
Total Non-Current Debt	\$ 33,901	32,877

*Tranches related to Kenvue

The weighted average effective interest rate on non-current debt is 3.54%.

The excess of the carrying value over the estimated fair value of debt was \$1.6 billion at January 1, 2023.

Fair value of the non-current debt was estimated using market prices, which were corroborated by quoted broker prices and significant other observable inputs.

In March 2023, Kenvue priced an offering of senior unsecured notes (the Notes) in an aggregate principal amount of \$7.75 billion (tranches with an * in the above table). The Notes were initially fully and unconditionally guaranteed (the Guarantees) on a senior unsecured basis by the Company. The Guarantees terminated on April 5, 2023, upon completion of the Consumer Health Business transfer.

The current debt balance as of July 2, 2023 includes \$9.8 billion of commercial paper (\$0.7 billion of which relates to Kenvue) which has a weighted average interest rate of 4.99% and a weighted average maturity of approximately forty days.

NOTE 5 — INCOME TAXES

The worldwide effective income tax rates for the fiscal six months of 2023 and 2022 were 15.8% and 14.9%, respectively. The increase in the consolidated tax rate is primarily due to the planned separation of the Company's Consumer Health business. The Company has recognized approximately \$0.6 billion in incremental international tax costs due to the reorganization of certain international subsidiaries which was recorded in the fiscal second quarter. The Company also had more income in lower tax jurisdictions relative to higher tax jurisdictions versus the prior year, due primarily to the approximately \$7 billion charge related to the talc settlement proposal in the United States at an effective tax rate of 23.5% in the fiscal six months of 2023 (for further information see Note 11 to the Consolidated Financial Statements). Additionally, the prior year's effective tax rate benefited from the impact of certain provisions of the Tax Cuts and Jobs Act of 2017 that became effective in 2022, which were partially offset by one-time tax costs in the fiscal six months of 2022.

On July 21, 2023, the IRS issued Notice 2023-55 which provides guidance to taxpayers in determining whether a foreign tax is eligible for a U.S. foreign tax credit for tax years 2022 and 2023, specifically delaying until 2024 the application of unfavorable foreign tax credit regulations that were originally issued late last year. As a result of this new guidance, the Company has concluded that it is applicable to certain of its tax positions and therefore will record a tax benefit of approximately \$0.5 billion in the fiscal third quarter of 2023.

The Company also received tax benefits from stock-based compensation that were either exercised or vested during each of the fiscal six months ended.

As of July 2, 2023, the Company had approximately \$3.0 billion of liabilities from unrecognized tax benefits. The Company conducts business and files tax returns in numerous countries and currently has tax audits in progress in a number of jurisdictions. With respect to the United States, the IRS has completed its audit for the tax years through 2012 and is currently auditing tax years 2013 through 2016. The Company currently expects completion of this audit and settlement of the related tax liabilities in the next 12 months. As a result, the Company has classified approximately \$0.5 billion of unrecognized tax benefits and associated interest as a current liability on the "Accrued taxes on Income" line of the Consolidated Balance Sheet as of the end of the second fiscal quarter of 2023 in anticipation of final settlement. The Company made a payment in the fiscal second quarter for approximately \$1.4 billion to the U.S. Treasury for the previously reserved estimated liability of the 2013-2016 IRS Audit. The completion of this tax audit may result in additional adjustments to the Company's unrecognized tax benefit liability. In other major jurisdictions where the Company conducts business, the years that remain open to tax audit go back to the year 2008. The Company believes it is possible that some tax audits may be completed over the next twelve months by taxing authorities in some jurisdictions. However, the Company is not able to provide a reasonably reliable estimate of the timing of any other future tax payments relating to uncertain tax positions.

NOTE 6 — PENSIONS AND OTHER BENEFIT PLANS

Components of Net Periodic Benefit Cost

Net periodic benefit costs for the Company's defined benefit retirement plans and other benefit plans include the following components:

(Dollars in Millions)	Fiscal Second Quarter Ended				Fiscal Six Months Ended			
	Retirement Plans		Other Benefit Plans		Retirement Plans		Other Benefit Plans	
	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022
Service cost	\$ 216	319	69	80	428	640	137	160
Interest cost	373	229	55	27	727	459	109	53
Expected return on plan assets	(694)	(693)	(2)	(2)	(1,362)	(1,392)	(3)	(4)
Amortization of prior service cost/(credit)	(46)	(46)	(1)	(2)	(92)	(92)	(1)	(3)
Recognized actuarial (gains) losses	(50)	167	7	31	(100)	329	13	61
Curtailments and settlements	—	—	—	—	—	1	—	—
Net periodic benefit cost/(credit)	\$ (201)	(24)	128	134	(399)	(55)	255	267

The service cost component of net periodic benefit cost is presented in the same line items on the Consolidated Statement of Earnings where other employee compensation costs are reported, including Cost of products sold, Research and development expense, and Selling, marketing and administrative expenses. All other components of net periodic benefit cost are presented as part of Other (income) expense, net on the Consolidated Statement of Earnings.

Company Contributions

For the fiscal six months ended July 2, 2023, the Company contributed \$58 million and \$12 million to its U.S. and international retirement plans, respectively. The Company plans to continue to fund its U.S. defined benefit plans to comply with the Pension Protection Act of 2006. International plans are funded in accordance with local regulations.

NOTE 7 — ACCUMULATED OTHER COMPREHENSIVE INCOME

Components of other comprehensive income (loss) consist of the following:

(Dollars in Millions)	Foreign Currency Translation	Gain/(Loss) On Securities	Employee Benefit Plans	Gain/(Loss) On Derivatives & Hedges	Total Accumulated Other Comprehensive Income (Loss)
January 1, 2023	\$ (11,813)	(27)	(897)	(230)	(12,967)
Net change	(896)	21	(138)	297	(716)
Amount attributable to non-controlling interest	548				548
July 2, 2023	\$ (12,161)	(6)	(1,035)	67	(13,135)

Amounts in accumulated other comprehensive income are presented net of the related tax impact. Foreign currency translation is not adjusted for income taxes where it relates to permanent investments in international subsidiaries. For additional details on comprehensive income see the Consolidated Statements of Comprehensive Income.

Details on reclassifications out of Accumulated Other Comprehensive Income:

Gain/(Loss) On Securities - reclassifications released to Other (income) expense, net.

Employee Benefit Plans - reclassifications are included in net periodic benefit cost. See Note 6 for additional details.

Gain/(Loss) On Derivatives & Hedges - reclassifications to earnings are recorded in the same account as the underlying transaction. See Note 4 for additional details.

NOTE 8 — EARNINGS PER SHARE

The following is a reconciliation of basic net earnings per share to diluted net earnings per share:

(Shares in Millions)	Fiscal Second Quarter Ended		Fiscal Six Months Ended	
	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022
Basic net earnings per share	\$ 1.98	1.83	1.95	3.79
Average shares outstanding — basic	2,598.4	2,629.6	2,601.9	2,629.4
Potential shares exercisable under stock option plans	95.2	143.7	96.9	141.8
Less: shares which could be repurchased under treasury stock method	(67.9)	(105.4)	(68.1)	(102.0)
Average shares outstanding — diluted	2,625.7	2,667.9	2,630.7	2,669.2
Diluted net earnings per share	\$ 1.96	1.80	1.93	3.73

The diluted net earnings per share calculation for the fiscal second quarter ended July 2, 2023 excluded 50.8 million shares related to stock options, as the exercise price of these options was greater than the average market value of the Company's stock. The diluted net earnings per share calculation for the fiscal second quarter ended July 3, 2022 included all shares related to stock options, as the exercise price of all options was less than the average market value of the Company's stock.

The diluted net earnings per share calculation for the fiscal six months ended July 2, 2023 excluded 46.8 million shares related to stock options, as the exercise price of these options was greater than the average market value of the Company's stock. The diluted net earnings per share calculation for the fiscal six months ended July 3, 2022 included all shares related to stock options, as the exercise price of all options was less than the average market value of the Company's stock.

NOTE 9 — SEGMENTS OF BUSINESS AND GEOGRAPHIC AREAS

SALES BY SEGMENT OF BUSINESS

(Dollars in Millions)	Fiscal Second Quarter Ended			Fiscal Six Months Ended		
	July 2, 2023	July 3, 2022	Percent Change	July 2, 2023	July 3, 2022	Percent Change
CONSUMER HEALTH						
OTC						
U.S.	\$ 712	663	7.5 %	\$ 1,457	1,333	9.3 %
International	947	818	15.7	1,844	1,609	14.6
Worldwide	1,659	1,482	12.0	3,301	2,943	12.2
Skin Health/Beauty						
U.S.	650	629	3.4	1,267	1,173	8.0
International	498	497	0.3	991	965	2.7
Worldwide	1,148	1,126	2.0	2,258	2,138	5.6
Oral Care						
U.S.	173	170	1.7	332	313	6.2
International	225	224	0.3	427	447	(4.7)
Worldwide	398	394	0.9	759	760	(0.2)
Baby Care						
U.S.	99	88	12.5	195	173	12.7
International	261	287	(9.1)	524	557	(6.0)
Worldwide	360	375	(4.0)	719	730	(1.6)
Women's Health						
U.S.	4	3	(2.1)	7	7	(0.1)
International	235	228	3.5	449	452	(0.7)
Worldwide	238	230	3.4	455	458	(0.6)
Wound Care/Other						
U.S.	149	133	12.3	264	245	7.9
International	58	65	(10.3)	107	117	(8.4)
Worldwide	207	197	4.9	371	361	2.6
TOTAL CONSUMER HEALTH						
U.S.	1,787	1,687	6.0	3,522	3,244	8.6
International	2,224	2,118	5.0	4,341	4,147	4.7
Worldwide	4,011	3,805	5.4	7,863	7,391	6.4

PHARMACEUTICAL

Immunology							
U.S.	2,865	2,853	0.4	5,313	5,354	(0.8)	
International	1,631	1,559	4.7	3,295	3,176	3.8	
Worldwide	4,496	4,411	1.9	8,608	8,530	0.9	
<u>REMICADE</u>							
U.S.	277	391	(29.3)	553	749	(26.2)	
U.S. Exports	33	44	(24.9)	74	124	(40.3)	
International	152	212	(28.2)	322	437	(26.2)	
Worldwide	462	647	(28.6)	949	1,310	(27.5)	
<u>SIMPONI / SIMPONIARIA</u>							
U.S.	285	301	(5.1)	556	588	(5.4)	
International	244	266	(8.2)	510	549	(7.1)	
Worldwide	529	566	(6.6)	1,066	1,137	(6.2)	
<u>STELARA</u>							
U.S.	1,817	1,731	4.9	3,268	3,110	5.1	
International	981	868	13.0	1,974	1,777	11.1	
Worldwide	2,797	2,599	7.6	5,241	4,887	7.2	
<u>TREMFYA</u>							
U.S.	450	382	17.8	856	773	10.7	
International	255	214	19.4	489	413	18.4	
Worldwide	706	597	18.3	1,346	1,187	13.4	
<u>OTHER IMMUNOLOGY</u>							
U.S.	4	3	17.8	7	9	(30.2)	
International	0	0	—	0	0	—	
Worldwide	4	3	17.8	7	9	(30.2)	
Infectious Diseases							
U.S.	395	415	(4.9)	787	876	(10.2)	
International	727	901	(19.4)	1,920	1,737	10.5	
Worldwide	1,121	1,316	(14.8)	2,707	2,613	3.6	
<u>COVID-19 VACCINE</u>							
U.S.	—	45	*	—	120	*	
International	285	499	(43.0)	1,032	881	17.1	
Worldwide	285	544	(47.7)	1,032	1,001	3.0	
<u>EDURANT / rilpivirine</u>							
U.S.	8	9	(9.0)	17	18	(5.3)	
International	258	215	19.9	529	454	16.4	
Worldwide	266	225	18.6	546	473	15.6	
<u>PREZISTA / PREZCOBIX / REZOLSTA / SYMTUZA</u>							
U.S.	382	355	7.6	760	724	5.0	
International	109	110	(0.2)	208	242	(13.9)	
Worldwide	491	464	5.8	968	965	0.3	

<u>OTHER INFECTIOUS DISEASES</u>						
U.S.	5	6	(27.2)	10	14	(30.6)
International	74	77	(3.5)	151	160	(5.6)
Worldwide	79	83	(5.2)	161	174	(7.6)
Neuroscience						
U.S.	1,029	896	14.9	2,007	1,739	15.4
International	764	837	(8.8)	1,590	1,735	(8.4)
Worldwide	1,793	1,734	3.5	3,597	3,475	3.5
<u>CONCERTA / methylphenidate</u>						
U.S.	64	38	68.2	134	73	84.0
International	143	123	16.3	279	245	13.9
Worldwide	208	161	28.6	414	318	30.0
<u>INVEGA SUSTENNA / XEPLION / INVEGA TRINZA / TREVICTA</u>						
U.S.	721	691	4.3	1,434	1,352	6.0
International	310	362	(14.4)	641	749	(14.5)
Worldwide	1,031	1,054	(2.1)	2,075	2,102	(1.3)
<u>SPRAVATO</u>						
U.S.	144	74	93.2	255	135	88.3
International	25	11	*	45	20	*
Worldwide	169	85	98.2	300	155	93.1
<u>OTHER NEUROSCIENCE⁽¹⁾</u>						
U.S.	100	93	9.3	184	179	3.1
International	286	341	(16.4)	625	721	(13.4)
Worldwide	386	433	(10.9)	809	900	(10.1)
Oncology						
U.S.	2,069	1,679	23.2	3,958	3,261	21.4
International	2,329	2,362	(1.4)	4,552	4,731	(3.8)
Worldwide	4,398	4,042	8.8	8,510	7,992	6.5
<u>CARVYKTI</u>						
U.S.	114	24	*	184	24	*
International	3	—	*	5	—	*
Worldwide	117	24	*	189	24	*
<u>DARZALEX</u>						
U.S.	1,322	1,021	29.5	2,513	1,974	27.3
International	1,110	965	15.0	2,182	1,868	16.8
Worldwide	2,431	1,986	22.4	4,695	3,842	22.2
<u>ERLEADA</u>						
U.S.	241	233	3.6	490	439	11.8
International	326	218	49.7	619	412	50.3
Worldwide	567	450	25.9	1,109	850	30.4
<u>IMBRUVICA</u>						
U.S.	262	349	(24.9)	532	719	(26.0)
International	579	620	(6.7)	1,136	1,288	(11.8)
Worldwide	841	970	(13.2)	1,668	2,008	(16.9)
<u>ZYTIGA / abiraterone acetate</u>						

U.S.	9	19	(55.2)	25	38	(34.7)
International	218	486	(55.1)	447	1,006	(55.6)
Worldwide	227	505	(55.1)	472	1,044	(54.8)
<u>OTHER ONCOLOGY</u>						
U.S.	122	33	*	214	67	*
International	92	72	27.8	162	156	3.6
Worldwide	214	106	*	376	224	68.0
<u>Pulmonary Hypertension</u>						
U.S.	684	560	22.0	1,284	1,132	13.4
International	289	284	2.0	561	563	(0.4)
Worldwide	972	843	15.3	1,844	1,695	8.8
<u>OPSUMIT</u>						
U.S.	328	265	23.7	601	538	11.6
International	179	173	3.4	346	343	0.9
Worldwide	507	438	15.7	947	881	7.5
<u>UPTRAVI</u>						
U.S.	338	272	24.2	642	541	18.7
International	61	56	10.0	119	112	6.6
Worldwide	399	328	21.8	761	653	16.6
<u>OTHER PULMONARY HYPERTENSION</u>						
U.S.	18	23	(23.8)	41	53	(23.0)
International	48	55	(10.9)	95	108	(11.7)
Worldwide	66	78	(14.7)	136	161	(15.5)
<u>Cardiovascular / Metabolism / Other</u>						
U.S.	776	757	2.6	1,491	1,429	4.3
International	174	215	(19.0)	386	453	(14.7)
Worldwide	950	972	(2.2)	1,877	1,882	(0.3)
<u>XARELTO</u>						
U.S.	637	609	4.7	1,215	1,117	8.8
International	—	—	—	—	—	—
Worldwide	637	609	4.7	1,215	1,117	8.8
<u>OTHER⁽²⁾</u>						
U.S.	138	148	(6.3)	275	312	(11.8)
International	174	215	(19.0)	386	453	(14.7)
Worldwide	313	363	(13.8)	662	765	(13.5)
<u>TOTAL PHARMACEUTICAL</u>						
U.S.	7,818	7,159	9.2	14,841	13,791	7.6
International	5,913	6,158	(4.0)	12,303	12,395	(0.7)
Worldwide	13,731	13,317	3.1	27,144	26,186	3.7

MEDTECH						
Interventional Solutions						
U.S.	908	525	73.1	1,771	1,019	73.8
International	712	525	35.7	1,352	1,123	20.5
Worldwide	1,620	1,049	54.4	3,123	2,141	45.8
<u>ELECTROPHYSIOLOGY</u>						
U.S.	609	499	22.0	1,180	969	21.7
International	587	469	25.1	1,109	1,001	10.8
Worldwide	1,196	968	23.5	2,288	1,970	16.2
<u>ABIOMED⁽³⁾</u>						
U.S.	272	—	*	536	—	*
International	59	—	*	119	—	*
Worldwide	331	—	*	655	—	*
<u>OTHER INTERVENTIONAL SOLUTIONS</u>						
U.S.	27	26	4.5	55	51	10.8
International	67	56	20.0	125	121	2.8
Worldwide	93	81	15.1	180	171	5.1
Orthopaedics						
U.S.	1,388	1,338	3.7	2,751	2,627	4.7
International	878	820	7.0	1,759	1,719	2.3
Worldwide	2,265	2,157	5.0	4,510	4,345	3.8
<u>HIPS</u>						
U.S.	250	240	4.1	491	465	5.6
International	147	148	(0.8)	296	312	(5.1)
Worldwide	397	388	2.2	787	777	1.3
<u>KNEES</u>						
U.S.	221	216	2.4	447	417	7.2
International	142	133	6.3	284	271	4.8
Worldwide	363	349	3.9	731	688	6.3
<u>TRAUMA</u>						
U.S.	483	464	4.3	974	939	3.7
International	255	232	9.9	522	505	3.2
Worldwide	739	696	6.1	1,496	1,444	3.6
<u>SPINE, SPORTS & OTHER</u>						
U.S.	433	418	3.5	839	805	4.1
International	334	306	9.0	657	630	4.2
Worldwide	766	724	5.8	1,495	1,436	4.2
Surgery						
U.S.	1,015	992	2.2	1,990	1,913	4.0
International	1,580	1,458	8.4	3,039	2,971	2.3
Worldwide	2,594	2,450	5.9	5,028	4,884	3.0
<u>ADVANCED</u>						
U.S.	466	454	2.7	910	871	4.5
International	757	702	7.8	1,430	1,431	0.0
Worldwide	1,222	1,156	5.8	2,340	2,302	1.7
<u>GENERAL</u>						
U.S.	548	538	1.9	1,079	1,042	3.6

International	823	756	8.9	1,608	1,540	4.5
Worldwide	1,372	1,294	6.0	2,688	2,582	4.1
Vision						
U.S.	529	496	6.6	1,087	1,017	6.9
International	778	745	4.6	1,521	1,481	2.7
Worldwide	1,308	1,241	5.4	2,608	2,498	4.4
CONTACT LENSES / OTHER						
U.S.	409	374	9.1	853	774	10.1
International	530	519	2.2	1,039	1,030	0.9
Worldwide	939	894	5.1	1,892	1,804	4.9
SURGICAL						
U.S.	120	122	(1.1)	234	243	(3.6)
International	249	225	10.1	482	451	6.7
Worldwide	369	347	6.2	716	694	3.1
TOTAL MEDTECH						
U.S.	3,839	3,351	14.6	7,598	6,576	15.5
International	3,949	3,547	11.3	7,671	7,293	5.2
Worldwide	7,788	6,898	12.9	15,269	13,869	10.1
WORLDWIDE						
U.S.	13,444	12,197	10.2	25,961	23,611	10.0
International	12,086	11,823	2.2	24,315	23,835	2.0
Worldwide	\$ 25,530	24,020	6.3 %	\$ 50,276	47,446	6.0 %

*Percentage greater than 100% or not meaningful

(1) Inclusive of RISPERDAL CONSTA which was previously disclosed separately

(2) Inclusive of INVOKANA which was previously disclosed separately

(3) Acquired on December 22, 2022

EARNINGS BEFORE PROVISION FOR TAXES BY SEGMENT

(Dollars in Millions)	Fiscal Second Quarter Ended			Fiscal Six Months Ended		
	July 2, 2023	July 3, 2022	Percent Change	July 2, 2023	July 3, 2022	Percent Change
Consumer Health ⁽¹⁾	\$ 860	784	9.7 %	\$ 1,636	1,470	11.3%
Pharmaceutical ⁽²⁾	4,862	4,420	10.0	9,306	8,344	11.5
MedTech ⁽³⁾	1,699	1,141	48.9	3,144	2,618	20.1
Segment earnings before provision for taxes	7,421	6,345	17.0	14,086	12,432	13.3
Less: Expense not allocated to segments ⁽⁴⁾	377	237		7,479	360	
Less: Consumer Health separation costs	282	268		582	370	
Worldwide income before tax	\$ 6,762	5,840	15.8%	\$ 6,025	11,702	(48.5)%

*Percentage greater than 100% or not meaningful

(1) Consumer Health includes:

- Intangible amortization expense of \$0.1 billion in both the fiscal second quarter of 2023 and 2022.
- Intangible amortization expense of \$0.2 billion in both the fiscal six months of 2023 and 2022.

(2) Pharmaceutical includes:

- Intangible amortization expense of \$0.7 billion in both the fiscal second quarter of 2023 and 2022.
- Intangible amortization expense of \$1.5 billion in both the fiscal six months of 2023 and 2022.

- One-time COVID-19 Vaccine related exit costs of \$0.2 billion and \$0.3 billion in the fiscal second quarter of 2023 and 2022, respectively. One-time COVID-19 Vaccine related exit costs of \$0.6 billion and \$0.3 billion in the fiscal six months of 2023 and 2022, respectively.
- A restructuring related charge of \$0.1 billion and \$0.3 billion in the fiscal second quarter and fiscal six months of 2023, respectively.
- In the fiscal six months of 2022, the Company recorded an intangible asset impairment charge of approximately \$0.6 billion related to an in-process research and development asset, bermekimab (JnJ-77474462), an investigational drug for the treatment of Atopic Dermatitis (AD) and Hidradenitis Suppurativa (HS).
- Unfavorable changes in the fair value of securities of \$0.1 billion in the fiscal second quarter of 2022. Unfavorable changes in the fair value of securities of \$0.1 billion and \$0.5 billion in the fiscal six months of 2023 and 2022, respectively.
- Favorable litigation related items of \$0.1 billion in both the fiscal second quarter and fiscal six months of 2023.

⁽³⁾ MedTech includes:

- Intangible amortization expense of \$0.4 billion and \$0.3 billion in the fiscal second quarter of 2023 and 2022, respectively. Intangible amortization expense of \$0.8 billion and \$0.5 billion in the fiscal six months of 2023 and 2022, respectively.
- Net favorable litigation matters of \$0.2 billion in both the fiscal second quarter and fiscal six months of 2023 and \$0.3 billion of expense in both the fiscal second quarter and fiscal six months of 2022.
- Acquisition and integration related expense of \$0.1 billion in the fiscal six months of 2023.
- A restructuring related charge of \$0.1 billion in the fiscal second quarter of 2022 and \$0.2 billion in the fiscal six months of 2022.

⁽⁴⁾ Amounts not allocated to segments include interest income/expense and general corporate income/expense. The fiscal six months of 2023 includes an approximately \$7 billion incremental charge primarily related to the talc settlement proposal. See Note 11, Legal Proceedings, for additional details.

SALES BY GEOGRAPHIC AREA

(Dollars in Millions)	Fiscal Second Quarter Ended			Fiscal Six Months Ended		
	July 2, 2023	July 3, 2022	Percent Change	July 2, 2023	July 3, 2022	Percent Change
United States	\$ 13,444	12,197	10.2 %	\$ 25,961	23,611	10.0 %
Europe	5,894	6,085	(3.1)	12,226	12,109	1.0
Western Hemisphere, excluding U.S.	1,713	1,536	11.5	3,300	3,018	9.3
Asia-Pacific, Africa	4,479	4,202	6.6	8,789	8,708	0.9
Total	\$ 25,530	24,020	6.3 %	\$ 50,276	47,446	6.0 %

NOTE 10— ACQUISITIONS AND DIVESTITURES

There were no material acquisitions or divestitures in the fiscal first quarter or fiscal second quarter of 2023.

On December 22, 2022, the Company completed the acquisition of Abiomed, a leading, first-to-market provider of cardiovascular medical technology with a first-in-kind portfolio for the treatment of coronary artery disease and heart failure which also has an extensive innovation pipeline of life-saving technologies. The transaction broadens the Company's position as a growing cardiovascular innovator, advancing the standard of care in heart failure and recovery, one of healthcare's largest areas of unmet need. The transaction was accounted for as a business combination and the results of operations were included in the MedTech segment as of the date of the acquisition. The acquisition was completed through a tender offer for all outstanding shares. The consideration paid in the acquisition consisted of an upfront payment of \$380.00 per share in cash, amounting to \$17.1 billion, net of cash acquired, as well as a non-tradeable contingent value right (CVR) entitling the holder to receive up to \$35.00 per share in cash (which with respect to the CVRs total approximately \$1.6 billion in the aggregate) if certain commercial and clinical milestones are achieved. The corresponding enterprise value (without taking into account the CVRs) of approximately \$16.5 billion includes cash, cash equivalents and marketable securities acquired.

The milestones of the CVR consist of:

- \$17.50 per share, payable if net sales for Abiomed products exceeds \$3.7 billion during Johnson & Johnson's fiscal second quarter of 2027 through fiscal first quarter of 2028, or if this threshold is not met during this period and is

- subsequently met during any rolling four quarter period up to the end of Johnson & Johnson's fiscal first quarter of 2029, \$8.75 per share;
- b. \$7.50 per share payable upon FDA premarket application approval of the use of Impella products in ST-elevated myocardial infarction (STEMI) patients without cardiogenic shock by January 1, 2028; and
 - c. \$10.00 per share payable upon the first publication of a Class I recommendation for the use of Impella products in high risk PCI or STEMI with or without cardiogenic shock within four years from their respective clinical endpoint publication dates, but in all cases no later than December 31, 2029.

The fair value of the acquisition was initially allocated to assets acquired of \$19.9 billion (net of \$0.3 billion cash acquired), primarily to goodwill for \$10.9 billion, amortizable intangible assets for \$6.6 billion, IPR&D for \$1.1 billion, marketable securities of \$0.6 billion and liabilities assumed of \$2.8 billion, which includes the fair value of the contingent consideration mentioned above for \$0.7 billion and deferred taxes of \$1.8 billion. The goodwill is primarily attributable to the commercial acceleration and expansion of the portfolio and is not expected to be deductible for tax purposes. The contingent consideration was recorded in Other Liabilities on the Consolidated Balance Sheet.

As the acquisition occurred in December 2022, the Company is still finalizing the allocation of the purchase price to the individual assets acquired and liabilities assumed. The allocation of the purchase price included in the current period balance sheet is based on the best estimate of management and is preliminary and subject to change. To assist management in the allocation, the Company engaged valuation specialists to prepare appraisals. The Company will finalize the amounts recognized as the information necessary to complete the analysis is obtained. The Company expects to finalize these amounts as soon as possible but no later than one year from the acquisition date. In the fiscal first quarter of 2023, there were purchase price allocation adjustments netting to approximately \$0.1 billion with an offsetting increase to goodwill. In the fiscal second quarter of 2023, there were no purchase price allocation adjustments.

The amortizable intangible assets were primarily comprised of already in-market products of the Impella platform with an average weighted life of 14 years. The IPR&D assets were valued for technology programs for unapproved products. The value of the IPR&D was calculated using probability-adjusted cash flow projections discounted for the risk inherent in such projects. The probability of success factor ranged from 52% to 70%. The discount rate applied was 9.5%.

In 2022, the Company recorded acquisition related costs before tax of approximately \$0.3 billion, which was recorded in Other (income)/expense. In the fiscal six months of 2023, the Company recorded acquisition related costs before tax of approximately \$0.1 billion, which was primarily recorded in Other (income)/expense.

There were no material acquisitions or divestitures in the fiscal first quarter or fiscal second quarter of 2022.

NOTE 11 — LEGAL PROCEEDINGS

Johnson & Johnson and certain of its subsidiaries are involved in various lawsuits and claims regarding product liability; intellectual property; commercial; indemnification and other matters; governmental investigations; and other legal proceedings that arise from time to time in the ordinary course of their business.

The Company records accruals for loss contingencies associated with these legal matters when it is probable that a liability will be incurred, and the amount of the loss can be reasonably estimated. As of July 2, 2023, the Company has determined that the liabilities associated with certain litigation matters are probable and can be reasonably estimated. The Company has accrued for these matters and will continue to monitor each related legal issue and adjust accruals as might be warranted based on new information and further developments in accordance with ASC 450-20-25. For these and other litigation and regulatory matters discussed below for which a loss is probable or reasonably possible, the Company is unable to estimate the possible loss or range of loss beyond the amounts accrued. Amounts accrued for legal contingencies often result from a complex series of judgments about future events and uncertainties that rely heavily on estimates and assumptions including timing of related payments. The ability to make such estimates and judgments can be affected by various factors including, among other things, whether damages sought in the proceedings are unsubstantiated or indeterminate; scientific and legal discovery has not commenced or is not complete; proceedings are in early stages; matters present legal uncertainties; there are significant facts in dispute; procedural or jurisdictional issues; the uncertainty and unpredictability of the number of potential claims; ability to achieve comprehensive multi-party settlements; complexity of related cross-claims and counterclaims; and/or there are numerous parties involved. To the extent adverse awards, judgments or verdicts have been rendered against the Company, the Company does not record an accrual until a loss is determined to be probable and can be reasonably estimated.

In the Company's opinion, based on its examination of these matters, its experience to date and discussions with counsel, the ultimate outcome of legal proceedings, net of liabilities accrued in the Company's balance sheet, is not expected to have a material adverse effect on the Company's financial position. However, the resolution of, or increase in accruals for, one or more of these matters in any reporting period may have a material adverse effect on the Company's results of operations and cash flows for that period.

MATTERS CONCERNING TALC

A significant number of personal injury claims alleging that talc causes cancer were made against Johnson & Johnson Consumer Inc. and the Company arising out of the use of body powders containing talc, primarily JOHNSON'S Baby Powder. The number of these personal injury lawsuits, filed in state and federal courts in the United States as well as outside of the United States, continued to increase.

In talc cases that previously have gone to trial, the Company has obtained a number of defense verdicts, but there also have been verdicts against the Company, many of which have been reversed on appeal. In June 2020, the Missouri Court of Appeals reversed in part and affirmed in part a July 2018 verdict of \$4.7 billion in *Ingham v. Johnson & Johnson, et al.*, No. ED 207476 (Mo. App.), reducing the overall award to \$2.1 billion. An application for transfer of the case to the Missouri Supreme Court was subsequently denied and in June 2021, a petition for certiorari, seeking a review of the Ingham decision by the United States Supreme Court, was denied. In June 2021, the Company paid the award, which, including interest, totaled approximately \$2.5 billion. The facts and circumstances, including the terms of the award, were unique to the Ingham decision and not representative of other claims brought against the Company. The Company continues to believe that it has strong legal grounds to contest the other talc verdicts that it has appealed. Notwithstanding the Company's confidence in the safety of its talc products, in certain circumstances the Company has settled cases.

In October 2021, Johnson & Johnson Consumer Inc. (Old JJCI) implemented a corporate restructuring (the 2021 Corporate Restructuring). As a result of that restructuring, Old JJCI ceased to exist and three new entities were created: (a) LTL Management LLC, a North Carolina limited liability company (LTL or Debtor); (b) Royalty A&M LLC, a North Carolina limited liability company and a direct subsidiary of LTL (RAM); and (c) the Debtor's direct parent, Johnson & Johnson Consumer Inc., a New Jersey company (New JJCI). The Debtor received certain of Old JJCI's assets and became solely responsible for the talc-related liabilities of Old JJCI, including all liabilities related in any way to injury or damage, or alleged injury or damage, sustained or incurred in the purchase or use of, or exposure to, talc, including talc contained in any product, or to the risk of, or responsibility for, any such damage or injury, except for any liabilities for which the exclusive remedy is provided under a workers' compensation statute or act (the Talc-Related Liabilities).

In October 2021, notwithstanding the Company's confidence in the safety of its talc products, the Debtor filed a voluntary petition with the United States Bankruptcy Court for the Western District of North Carolina, Charlotte Division, seeking relief under chapter 11 of the Bankruptcy Code (the LTL Bankruptcy Case). All litigation against LTL, Old JJCI, New JJCI, the Company, other of their corporate affiliates, identified retailers, insurance companies, and certain other parties (the Protected Parties) was stayed, although LTL did agree to lift the stay on a small number of appeals where appeal bonds had been filed. The LTL Bankruptcy Case was transferred to the United States Bankruptcy Court for the District of New Jersey. Claimants filed motions to dismiss the LTL Bankruptcy Case and, following a multiple day hearing, the New Jersey Bankruptcy Court denied those motions in March 2022.

The claimants subsequently filed notices of appeal as to the denial of the motions to dismiss the LTL Bankruptcy Case and the extension of the stay to the Protected Parties. On January 30, 2023, the Third Circuit reversed the Bankruptcy Court's ruling and remanded to the Bankruptcy Court to dismiss the LTL bankruptcy.

LTL filed a petition for rehearing of the Third Circuit's decision, which was denied in March 2023. LTL subsequently filed a motion in the Third Circuit to stay the mandate directing the New Jersey Bankruptcy Court to dismiss the LTL bankruptcy pending filing and disposition of a petition for writ of certiorari to the United States Supreme Court. The Third Circuit denied the motion to stay the mandate and issued the mandate.

In April 2023, the New Jersey Bankruptcy Court dismissed the LTL Bankruptcy Case, effectively lifting the stay as to all parties and returning the talc litigation to the tort system. LTL re-filed in the United States Bankruptcy Court for the District of New Jersey seeking relief under chapter 11 of the Bankruptcy Code (the LTL 2 Bankruptcy Case). As a result of the new filing, all talc claims against LTL were again automatically stayed pursuant to section 362 of the Bankruptcy Code. Additionally, the New Jersey Bankruptcy Court issued a temporary restraining order staying all litigation as to LTL, Old JJCI, New JJCI, the Company, identified retailers, and certain other parties (the New Protected Parties).

Also in April 2023, the New Jersey Bankruptcy Court issued a decision that granted limited injunctive relief to the Company and the New Protected Parties (the LTL 2 Preliminary Injunction). The LTL 2 Preliminary Injunction remains in force until late August 2023, following the Bankruptcy Court's extension of the initial LTL 2 Preliminary Injunction in June 2023. Under the LTL 2 Preliminary Injunction, except for in those cases filed in the federal court ovarian cancer multi-district litigation, discovery in all personal injury and wrongful death matters is permitted to proceed. No trials may occur in any of the personal injury and wrongful death matters except for the Valadez trial after the Bankruptcy Court partially lifted the stay for Valadez and allowed it to proceed to trial. In July 2023, the jury returned a verdict in favor of Valadez for \$18.8 million in

compensatory damages but declined to award punitive damages. The Company will appeal. The terms of the court's lifting of the automatic stay will prevent any collection of judgment.

Furthermore, in April 2023, the Talc Claimants' Committee filed a motion to dismiss the LTL 2 Bankruptcy followed by similar motions from other claimants. Hearings on the motions to dismiss occurred in June 2023. On July 28, 2023, the court dismissed the LTL 2 Bankruptcy case and, the same day, the Company stated its intent to appeal the decision and to continue its efforts to obtain a resolution of the talc claims.

In the original bankruptcy case, the Company agreed to provide funding to LTL for the payment of amounts the New Jersey Bankruptcy Court determines are owed by LTL and the establishment of a \$2 billion trust in furtherance of this purpose. The Company established a reserve for approximately \$2 billion in connection with the aforementioned trust. After and as a result of the filing of the LTL Bankruptcy Case, the Company de-consolidated LTL through the second fiscal quarter, which is a related party. The impact of the de-consolidation is not material to the Company. In the LTL 2 Bankruptcy Case, the Company had agreed to contribute an additional amount which, when added to the prior \$2 billion, would be a total reserve of present value of approximately \$9 billion payable over 25 years (nominal value approximately \$12 billion discounted at a rate of 4.41%), to resolve all the current and future talc claims. The approximate \$9 billion reserve remains the Company's estimate of probable loss after the dismissal.

The parties have not yet reached a resolution of all talc matters and the Company is unable to estimate the possible loss or range of loss beyond the amount accrued.

A class action advancing claims relating to industrial talc was filed against the Company and others in New Jersey state court in May 2022 (the Edley Class Action). The Edley Class Action asserts, among other things, that the Company fraudulently defended past asbestos personal injury lawsuits arising from exposure to industrial talc mined, milled, and manufactured before January 6, 1989 by the Company's then wholly owned subsidiary, Windsor Minerals, Inc., which is currently a debtor in the Imerys Bankruptcy described hereafter. The Company removed the Edley Class Action to federal court in the District of New Jersey. In October 2022, the Company filed motions to dismiss and to deny certification of a class to pursue the Edley Class Action in the New Jersey District Court.

In February 2019, the Company's talc supplier, Imerys Talc America, Inc. and two of its affiliates, Imerys Talc Vermont, Inc. and Imerys Talc Canada, Inc. (collectively, Imerys) filed a voluntary petition under chapter 11 of the United States Code (the Bankruptcy Code) in the United States Bankruptcy Court for the District of Delaware (Imerys Bankruptcy). The Imerys Bankruptcy relates to Imerys's potential liability for personal injury from exposure to talcum powder sold by Imerys. In its bankruptcy, Imerys alleges it has claims against the Company for indemnification and rights to joint insurance proceeds.

In June 2020, Cyprus Mines Corporation and its parent, Cyprus Amax Minerals Company (CAMC) (together, Cyprus), which had owned certain Imerys talc mines, filed an adversary proceeding against the Company and Imerys in the Imerys Bankruptcy seeking a declaration of indemnity rights under certain contractual agreements (the Cyprus Adversary Proceeding). The Company denies such indemnification is owed and filed a motion to dismiss the adversary complaint. In February 2021, Cyprus filed a voluntary petition for relief under chapter 11 of the Bankruptcy Code and filed its Disclosure Statement and Plan (the Cyprus Plan). The Cyprus Plan contemplates a settlement with Imerys and talc claimants where Cyprus would make a monetary contribution to a trust established under the Imerys Plan in exchange for an injunction against talc claims asserted against it and certain protected parties. Cyprus has not yet sought approval of its Disclosure Statement and Plan. Cyprus, along with the Tort Claimants' Committee (TCC) and Future Claimants' Representative (FCR) appointed in the Cyprus chapter 11 case, have agreed to participate in the mediation with the Mediation Parties. In October 2021, the Company filed a Notice of Bankruptcy Filing and Stay of Proceedings clarifying that the automatic stay arising upon the filing of the LTL Bankruptcy Case should apply to the Cyprus Adversary Proceeding. In June 2022, Cyprus commenced an Adversary Proceeding in its chapter 11 case seeking an order enforcing the automatic stay by enjoining parties from commencing or continuing "talc-related claims" against CAMC. In June 2022, the court entered a preliminary injunction order enjoining claimants from pursuing talc-related claims against CAMC through January 2023. The court subsequently extended the preliminary injunction through the end of July 2023. In June 2023, Cyprus filed a motion seeking to further extend the preliminary injunction through the end of February 2024.

Imerys, the TCC, the FCR, certain of Imerys's insurers, and certain parties in the Cyprus Mines chapter 11 case (collectively the Mediation Parties) have been engaged in mediation since October 2021. In June 2023, the bankruptcy court entered an order extending the term of the mediation through the end of July 2023.

In July 2021, Imerys commenced an adversary proceeding against the Company in the Imerys Bankruptcy (the Imerys Adversary Proceeding). The Imerys Adversary Proceeding sought, among other things, certain declarations with respect to the indemnification obligations allegedly owed by the Company to Imerys. The Company filed a motion to dismiss the adversary proceeding.

In February 2021, several of the Company's insurers involved in coverage litigation in New Jersey State Court (the Coverage Action) filed a motion in the Imerys Bankruptcy Court proceeding seeking a determination that the automatic stay does not apply to the Coverage Action and, in the alternative, seeking relief from the automatic stay to allow them to continue to litigate their claims in the Coverage Action. The Court entered an agreed order modifying the stay to allow the litigation in the Coverage Action to continue.

In February 2018, a securities class action lawsuit was filed against the Company and certain named officers in the United States District Court for the District of New Jersey, alleging that the Company violated the federal securities laws by failing to disclose alleged asbestos contamination in body powders containing talc, primarily JOHNSON'S Baby Powder, and that purchasers of the Company's shares suffered losses as a result. In April 2019, the Company moved to dismiss the complaint. In December 2019, the Court denied, in part, the motion to dismiss. In April 2021, briefing on Plaintiff's motion for class certification was completed. The case was stayed in May 2022 pursuant to the LTL Bankruptcy Case and was reopened in May 2023. Defendants have requested that further fact discovery be stayed until the Court rules on Plaintiff's motion for class certification.

A lawsuit was brought against the Company in the Superior Court of California for the County of San Diego alleging violations of California's Consumer Legal Remedies Act (CLRA) relating to JOHNSON'S Baby Powder. In that lawsuit, the plaintiffs allege that the Company violated the CLRA by failing to provide required Proposition 65 warnings. In July 2019, the Company filed a notice of removal to the United States District Court for the Southern District of California and plaintiffs filed a second amended complaint shortly thereafter. In October 2019, the Company moved to dismiss the second amended complaint for failure to state a claim upon which relief may be granted. In response to those motions, plaintiffs filed a third amended complaint. In December 2019, the Company moved to dismiss the third amended complaint for failure to state a claim upon which relief may be granted. In April 2020, the Court granted the motion to dismiss but granted leave to amend. In May 2020, plaintiffs filed a Fourth Amended Complaint but indicated that they would be filing a motion for leave to file a fifth amended complaint. Plaintiffs filed a Fifth Amended Complaint in August 2020. The Company moved to dismiss the Fifth Amended Complaint for failure to state a claim upon which relief may be granted. In January 2021, the Court issued an Order and opinion ruling in the Company's favor and granting the motion to dismiss with prejudice. In February 2021, Plaintiffs filed a Notice of Appeal with the Ninth Circuit. Plaintiffs filed their opening brief in July 2021. The company filed its responsive brief in October 2021. After the Notice of Suggestion of Bankruptcy was filed with the Ninth Circuit, a stay was imposed, and the Court held the reply deadline in abeyance. The appeal continues to be held in abeyance, with the Company being required to file periodic status updates.

In June 2014, the Mississippi Attorney General filed a complaint in Chancery Court of The First Judicial District of Hinds County, Mississippi against the Company and Johnson & Johnson Consumer Companies, Inc. (now known as Johnson & Johnson Consumer Inc.) (collectively, JJCI). The complaint alleges that JJCI violated the Mississippi Consumer Protection Act by failing to disclose alleged health risks associated with female consumers' use of talc contained in JOHNSON'S Baby Powder and JOHNSON'S Shower to Shower (a product divested in 2012) and seeks injunctive and monetary relief. In February 2022, the trial court set the case for trial to begin in February 2023. However, in October 2022, the LTL bankruptcy court issued an order staying the case. In March 2023, the Third Circuit issued the mandate to dismiss the LTL Bankruptcy Case and in April 2023, the New Jersey Bankruptcy Court dismissed the LTL Bankruptcy Case, effectively lifting the stay as to this matter. The State requested a new trial setting. Later in April 2023, the trial court set a new trial date in April 2024. The parties are currently engaged in expert work, extensive discovery, and preparations for the upcoming trial.

In January 2020, the State of New Mexico filed a consumer protection case alleging that the Company deceptively marketed and sold its talcum powder products by making misrepresentations about the safety of the products and the presence of carcinogens, including asbestos. In March 2022, the New Mexico court denied the Company's motion to compel the State of New Mexico to engage in discovery of state agencies and denied the Company's request for interlocutory appeal of that decision. The Company then filed a Petition for Writ of Superintending Control and a Request for a Stay to the New Mexico Supreme Court on the issue of the State of New Mexico's discovery obligations. In April 2022, in view of the efforts to resolve talc-related claims in the LTL Bankruptcy Case, the Company and the State agreed to a 60-day stay of all matters except for the pending writ before the New Mexico Supreme Court, which expired in June 2022. Thereafter, the Company moved to enjoin prosecution of the case in the LTL Bankruptcy Case. In October 2022, the bankruptcy court issued an order staying the case. In December 2022, the State filed an appeal to the Third Circuit concerning the stay order. Separately, in September 2022, the New Mexico Supreme Court granted the Company's request for a stay pending further briefing on the scope of the State of New Mexico's discovery obligations. In March 2023, the Third Circuit issued the mandate to dismiss the LTL Bankruptcy Case and in April 2023, the New Jersey Bankruptcy Court dismissed the LTL Bankruptcy Case, effectively lifting the stay as to this matter. However, this case remains stayed as a result of the New Mexico Supreme Court's stay until such time as the Supreme Court issues an order concerning the State of New Mexico's discovery obligations.

Forty-two states and the District of Columbia (including Mississippi and New Mexico) have commenced a joint investigation into the Company's marketing of its talcum powder products. At this time, the multi-state group has not asserted any claims against the Company. Five states have issued Civil Investigative Demands seeking documents and other information. The Company has produced documents to Arizona, North Carolina, Texas, and Washington and entered into confidentiality

agreements. The Company has not received any follow up requests from those states. In March 2022, each of the forty-two states agreed to mediation of their claims in the LTL Bankruptcy Case. In July 2022, New Mexico and Mississippi indicated they would no longer voluntarily submit to further mediation in the LTL Bankruptcy and would proceed with their respective cases in state court. In March 2023, the mediation was terminated. The procedural history and status of the New Mexico and Mississippi matters specifically have been discussed above.

In addition, the Company has received inquiries, subpoenas, and requests to produce documents regarding talc matters and the LTL Bankruptcy Case from various governmental authorities. The Company has produced documents and responded to inquiries, and will continue to cooperate with government inquiries.

MATTERS CONCERNING OPIOIDS

Beginning in 2014 and continuing to the present, the Company and Janssen Pharmaceuticals, Inc. (JPI), along with other pharmaceutical companies, have been named in close to 3,500 lawsuits related to the marketing of opioids, including DURAGESIC, NUCYNTA and NUCYNTA ER. The majority of the cases have been filed by state and local governments. Similar lawsuits have also been filed by private plaintiffs and organizations, including but not limited to the following: individual plaintiffs on behalf of children born with Neonatal Abstinence Syndrome (NAS); hospitals; and health insurers/payors.

To date, the Company and JPI have litigated two of the cases to judgment and have prevailed in both, either at trial or on appeal.

In October 2019, the Company announced a proposed agreement in principle with a negotiating committee of state Attorneys General to settle all remaining government opioid litigation claims nationwide. Under the final national settlement agreement, which was announced in July 2021, the Company agreed to pay up to \$5.0 billion to resolve all opioid lawsuits and future opioid claims by states, cities, counties, local school districts and other special districts, and tribal governments, contingent on sufficient participation by eligible government entities, and with credits back for entities that declined or were ineligible to participate. In July 2021, the Company announced that the terms of the agreement to settle the state and subdivision claims had been finalized and approximately 60% of the all-in settlement was paid by the second fiscal quarter of 2023. The expected payment schedule provides that approximately \$0.6 billion of payments are to be paid by the end of the second fiscal quarter of 2024. The agreement is not an admission of liability or wrongdoing, and it provides for the release of all opioid-related claims against the Company, JPI, and their affiliates (including the Company's former subsidiaries Tasmanian Alkaloids Pty, Ltd. and Noramco, Inc.). As of July 2023, the Company and JPI had settled or otherwise resolved the opioid claims advanced by all government entity claimants except the State of Washington and its subdivisions, the City of Baltimore, a number of school districts and other special district claimants, and a handful of others.

The Company and JPI continue to defend the cases brought by the remaining government entity litigants as well as the cases brought by private litigants, including NAS claimants, hospitals, and health insurers/payors. Counting the private litigant cases, there are approximately 35 remaining opioid cases against the Company and JPI in various state courts, 475 remaining cases in the Ohio MDL, and 2 additional cases in other federal courts. Some of these cases have been dismissed and are being appealed by the plaintiffs; several others, including a case brought by DCH Health Systems in Alabama state court, are scheduled for trial in 2023, 2024, or 2025.

In addition, the Province of British Columbia filed suit against the Company and its Canadian affiliate Janssen Inc., and many other industry members, in Canada, and is seeking to have that action certified as an opt in class action on behalf of other provincial/territorial and the federal governments in Canada. Additional proposed class actions have been filed in Canada against the Company and Janssen Inc., and many other industry members, by and on behalf of people who used opioids (for personal injuries), municipalities and First Nations bands. These actions allege a variety of claims related to opioid marketing practices, including false advertising, unfair competition, public nuisance, consumer fraud violations, deceptive acts and practices, false claims and unjust enrichment. An adverse judgment in any of these lawsuits could result in the imposition of large monetary penalties and significant damages including, punitive damages, cost of abatement, substantial fines, equitable remedies and other sanctions.

From June 2017 through December 2019, the Company's Board of Directors received a series of shareholder demand letters alleging breaches of fiduciary duties related to the marketing of opioids. The Board retained independent counsel to investigate the allegations in the demands, and in April 2020, independent counsel delivered a report to the Board recommending that the Company reject the shareholder demands and take the steps that are necessary or appropriate to secure dismissal of related derivative litigation. The Board unanimously adopted the recommendations of the independent counsel's report.

In November 2019, one of the shareholders who sent a demand filed a derivative complaint against the Company as the nominal defendant and certain current and former directors and officers as defendants in the Superior Court of New Jersey. The complaint alleges breaches of fiduciary duties related to the marketing of opioids, and that the Company has suffered damages as a result of those alleged breaches. A series of additional derivative complaints making similar allegations against the same

and similar defendants were filed in New Jersey state and federal courts in 2019 and 2020. By 2022, all but two state court cases had been voluntarily dismissed. In February 2022, the state court granted the Company's motion to dismiss one of the two cases, and the shareholder that brought the second case filed a notice of dismissal. The shareholder whose complaint was dismissed filed a motion for reconsideration. In May 2022, the state court held oral argument on the motion for reconsideration and subsequently denied the motion. The shareholder has appealed the state court's dismissal order.

PRODUCT LIABILITY

The Company and certain of its subsidiaries are involved in numerous product liability claims and lawsuits involving multiple products. Claimants in these cases seek substantial compensatory and, where available, punitive damages. While the Company believes it has substantial defenses, it is not feasible to predict the ultimate outcome of litigation. From time to time, even if it has substantial defenses, the Company considers isolated settlements based on a variety of circumstances. The Company has accrued for these matters and will continue to monitor each related legal issue and adjust accruals as might be warranted based on new information and further developments in accordance with ASC 450-20-25. The Company accrues an estimate of the legal defense costs needed to defend each matter when those costs are probable and can be reasonably estimated. For certain of these matters, the Company has accrued additional amounts such as estimated costs associated with settlements, damages and other losses. Product liability accruals can represent projected product liability for thousands of claims around the world, each in different litigation environments and with different fact patterns. Changes to the accruals may be required in the future as additional information becomes available.

The table below contains the most significant of these cases and provides the approximate number of plaintiffs in the United States with direct claims in pending lawsuits regarding injuries allegedly due to the relevant product or product category as of July 2, 2023:

Product or product category	Number of Plaintiffs
Body powders containing talc, primarily JOHNSON'S Baby Powder	40,480
DePuy ASR XL Acetabular System and DePuy ASR Hip Resurfacing System	160
PINNACLE Acetabular Cup System	940
Pelvic meshes	7,160
ETHICON PHYSIOMESH Flexible Composite Mesh	910
RISPERDAL	240
ELMIRON	2,130
TYLENOL	520

The number of pending lawsuits is expected to fluctuate as certain lawsuits are settled or dismissed and additional lawsuits are filed. There may be additional claims that have not yet been filed.

MedTech

DePuy ASR XL Acetabular System and ASR Hip Resurfacing System

In August 2010, DePuy Orthopaedics, Inc. (DePuy) announced a worldwide voluntary recall of its ASR XL Acetabular System and DePuy ASR Hip Resurfacing System (ASR Hip) used in hip replacement surgery. Claims for personal injury have been made against DePuy and the Company. Cases filed in federal courts in the United States have been organized as a multi-district litigation in the United States District Court for the Northern District of Ohio. Litigation has also been filed in countries outside of the United States, primarily in the United Kingdom, Canada, Australia, Ireland, Germany, India and Italy. In November 2013, DePuy reached an agreement with a Court-appointed committee of lawyers representing ASR Hip plaintiffs to establish a program to settle claims with eligible ASR Hip patients in the United States who had surgery to replace their ASR Hips, known as revision surgery, as of August 2013. DePuy reached additional agreements in February 2015 and March 2017, which further extended the settlement program to include ASR Hip patients who had revision surgeries after August 2013 and prior to February 15, 2017. This settlement program has resolved more than 10,000 claims, thereby bringing to resolution significant ASR Hip litigation activity in the United States. However, lawsuits in the United States remain, and the settlement program does not address litigation outside of the United States. In Australia, a class action settlement was reached that resolved the claims of the majority of ASR Hip patients in that country. In Canada, the Company has reached agreements to settle the class actions filed in that country. The Company continues to receive information with respect to potential additional costs associated with this recall on a worldwide basis. The Company has established accruals for the costs associated with the United States settlement program and ASR Hip-related product liability litigation.

DePuy PINNACLE Acetabular Cup System

Claims for personal injury have also been made against DePuy Orthopaedics, Inc. and the Company (collectively, DePuy) relating to the PINNACLE Acetabular Cup System used in hip replacement surgery. Product liability lawsuits continue to be filed, and the Company continues to receive information with respect to potential costs and the anticipated number of cases. Most cases filed in federal courts in the United States have been organized as a multi-district litigation in the United States District Court for the Northern District of Texas (Texas MDL). Beginning on June 1, 2022, the Judicial Panel on Multidistrict Litigation ceased transfer of new cases into the Texas MDL, and there are now cases pending in federal court outside the Texas MDL. Litigation also has been filed in state courts and in countries outside of the United States. During the first quarter of 2019, DePuy established a United States settlement program to resolve these cases. As part of the settlement program, adverse verdicts have been settled. The Company has established an accrual for product liability litigation associated with the PINNACLE Acetabular Cup System and the related settlement program.

Ethicon Pelvic Mesh

Claims for personal injury have been made against Ethicon, Inc. (Ethicon) and the Company arising out of Ethicon's pelvic mesh devices used to treat stress urinary incontinence and pelvic organ prolapse. The Company continues to receive information with respect to potential costs and additional cases. Cases filed in federal courts in the United States had been organized as a multi-district litigation (MDL) in the United States District Court for the Southern District of West Virginia. In March 2021, the MDL Court entered an order closing the MDL. The MDL Court has remanded cases for trial to the jurisdictions where the case was originally filed and additional pelvic mesh lawsuits have been filed, and remain, outside of the MDL. The Company has settled or otherwise resolved the majority of the United States cases and the estimated costs associated with these settlements and the remaining cases are reflected in the Company's accruals. In addition, class actions and individual personal injury cases or claims seeking damages for alleged injury resulting from Ethicon's pelvic mesh devices have been commenced in various countries outside of the United States, including claims and cases in the United Kingdom, the Netherlands, Belgium, France, Ireland, Italy, Spain and Slovenia and class actions in Israel, Australia, Canada and South Africa. In November 2019, the Federal Court of Australia issued a judgment regarding its findings with respect to liability in relation to the three Lead Applicants and generally in relation to the design, manufacture, pre- and post-market assessments and testing, and supply and promotion of the devices in Australia used to treat stress urinary incontinence and pelvic organ prolapse. In September 2022, after exhausting its appeals, the Company reached an in-principle agreement to resolve the two pelvic mesh class actions in Australia and in March 2023 the Federal Court approved the settlement. The class actions in Canada were discontinued in 2020 as a result of a settlement of a group of cases and an agreement to resolve the Israeli class action was reached in May 2021. The parties in the Israeli class action are currently finalizing the terms of the settlement. A motion to approve the settlement was filed with the Court. The Company has established accruals with respect to product liability litigation associated with Ethicon's pelvic mesh products.

Ethicon Physiomes

Following a June 2016 worldwide market withdrawal of Ethicon Physiomes Flexible Composite Mesh (Physiomes), claims for personal injury have been made against Ethicon, Inc. (Ethicon) and the Company alleging personal injury arising out of the use of this hernia mesh device. Cases filed in federal courts in the United States have been organized as a multi-district litigation (MDL) in the United States District Court for the Northern District of Georgia. A multi-county litigation (MCL) also has been formed in New Jersey state court and assigned to Atlantic County for cases pending in New Jersey. In addition to the matters in the MDL and MCL, there are additional lawsuits pending in the United States District Court for the Southern District of Ohio, which are part of the MDL for polypropylene mesh devices manufactured by C.R. Bard, Inc., and lawsuits pending in two New Jersey MCLs formed for Proceed/Proceed Ventral Patch and Prolene Hernia systems, and lawsuits pending outside the United States. In May 2021, Ethicon and lead counsel for the plaintiffs entered into a term sheet to resolve approximately 3,600 Physiomes cases (covering approximately 4,300 plaintiffs) pending in the MDL and MCL at that time. A master settlement agreement (MSA) was entered into in September 2021 and includes 3,729 cases in the MDL and MCL. All deadlines and trial settings in those proceedings are currently stayed pending the completion of the settlement agreement. Of the cases subject to the MSA, 3,244 have been dismissed with prejudice. Ethicon has received releases from 3,559 plaintiffs, and releases continue to be submitted as part of the settlement process. Post-settlement cases in the Physiomes MDL and MCL are subject to docket control orders requiring early expert reports and discovery requirements. In May 2023, Ethicon entered an additional settlement to resolve the claims of 292 Physiomes claimants. That settlement is proceeding, and releases are being returned. There are two cases in the MDL and two in the MCL which are not included in either settlement and which remain subject to the docket control orders.

Claims have also been filed against Ethicon and the Company alleging personal injuries arising from the PROCEED Mesh and PROCEED Ventral Patch hernia mesh products. In March 2019, the New Jersey Supreme Court entered an order consolidating these cases pending in New Jersey as an MCL in Atlantic County Superior Court. Additional cases have been filed in various federal and state courts in the United States, and in jurisdictions outside the United States.

Ethicon and the Company also have been subject to claims for personal injuries arising from the PROLENE Polypropylene Hernia System. In January 2020, the New Jersey Supreme Court created an MCL in Atlantic County Superior Court to handle such cases. Cases involving this product have also been filed in other federal and state courts in the United States.

In October 2022, an agreement in principle, subject to various conditions, was reached to settle the majority of the pending cases involving Proceed, Proceed Ventral Patch, Prolene Hernia System and related multi-layered mesh products, as well as a number of unfiled claims. All litigation activities in the two New Jersey MCLs are stayed pending effectuation of the proposed settlement. Future cases that are filed in the New Jersey MCLs will be subject to docket control orders requiring early expert reports and discovery requirements.

The Company has established accruals with respect to product liability litigation associated with Ethicon Physiomesher Flexible Composite Mesh, PROCEED Mesh and PROCEED Ventral Patch, and PROLENE Polypropylene Hernia System products.

Pharmaceuticals

RISPERDAL

Claims for personal injury have been made against Janssen Pharmaceuticals, Inc. and the Company arising out of the use of RISPERDAL, and related compounds, indicated for the treatment of schizophrenia, acute manic or mixed episodes associated with bipolar I disorder and irritability associated with autism. Lawsuits primarily have been filed in state courts in Pennsylvania, California, and Missouri. Other actions are pending in various courts in the United States and Canada. Product liability lawsuits continue to be filed, and the Company continues to receive information with respect to potential costs and the anticipated number of cases. The Company has successfully defended a number of these cases but there have been verdicts against the Company, including a verdict in October 2019 of \$8.0 billion of punitive damages related to one plaintiff, which the trial judge reduced to \$6.8 million in January 2020. In September 2021, the Company entered into a settlement in principle with the counsel representing plaintiffs in this matter and in substantially all of the outstanding cases in the United States. The costs associated with this and other settlements are reflected in the Company's accruals.

ELMIRON

Claims for personal injury have been made against a number of Johnson & Johnson companies, including Janssen Pharmaceuticals, Inc. and the Company, arising out of the use of ELMIRON, a prescription medication indicated for the relief of bladder pain or discomfort associated with interstitial cystitis. These lawsuits, which allege that ELMIRON contributes to the development of permanent retinal injury and vision loss, have been filed in both state and federal courts across the United States. In December 2020, lawsuits filed in federal courts in the United States, including putative class action cases seeking medical monitoring, were organized as a multi-district litigation in the United States District Court for the District of New Jersey. In addition, cases have been filed in various state courts of New Jersey, which have been coordinated in a multi-county litigation in Bergen County, as well as the Court of Common Pleas in Philadelphia, which have been coordinated and granted mass tort designation. In addition, three class action lawsuits have been filed in Canada. Product liability lawsuits continue to be filed, and the Company continues to receive information with respect to potential costs and the anticipated number of cases. The Company has established accruals for defense and indemnity costs associated with ELMIRON related product liability litigation.

Consumer Health

TYLENOL

Claims for personal injury have been made against Johnson and Johnson Consumer Inc. (JJCI) in federal court, arising out of the use of TYLENOL, an over-the-counter pain medication, alleging that prenatal exposure to acetaminophen is associated with the development of autism spectrum disorder and/or attention-deficit/hyperactivity disorder. In October 2022, lawsuits filed in federal courts in the United States were organized as a multi-district litigation in the United States District Court for the Southern District of New York. In addition, a lawsuit has been filed in state court against JJCI, Kenvue and the Company, and lawsuits have been filed in Canada against Johnson & Johnson Inc. and the Company. Product liability lawsuits continue to be filed, and the Company continues to receive information with respect to potential costs and the anticipated number of cases. The Company has established accruals for defense costs associated with TYLENOL related product liability litigation.

INTELLECTUAL PROPERTY

Certain subsidiaries of the Company are subject, from time to time, to legal proceedings and claims related to patent, trademark and other intellectual property matters arising out of their businesses. Many of these matters involve challenges to the coverage and/or validity of the patents on various products and allegations that certain of the Company's products infringe the patents of third parties. Although these subsidiaries believe that they have substantial defenses to these challenges and allegations with respect to all significant patents, there can be no assurance as to the outcome of these matters. A loss in any of these cases could adversely affect the ability of these subsidiaries to sell their products, result in loss of sales due to loss of market exclusivity, require the payment of past damages and future royalties, and may result in a non-cash impairment charge for any associated intangible asset.

Pharmaceuticals - Litigation Against Filers of Abbreviated New Drug Applications (ANDAs)

The Company's subsidiaries have brought lawsuits against generic companies that have filed ANDAs with the U.S. FDA (or similar lawsuits outside of the United States) seeking to market generic versions of products sold by various subsidiaries of the Company prior to expiration of the applicable patents covering those products. These lawsuits typically include allegations of non-infringement and/or invalidity of patents listed in FDA's publication "Approved Drug Products with Therapeutic Equivalence Evaluations" (commonly known as the Orange Book). In each of these lawsuits, the Company's subsidiaries are seeking an order enjoining the defendant from marketing a generic version of a product before the expiration of the relevant patents (Orange Book Listed Patents). In the event the Company's subsidiaries are not successful in an action, or any automatic statutory stay expires before the court rulings are obtained, the generic companies involved would have the ability, upon regulatory approval, to introduce generic versions of their products to the market, resulting in the potential for substantial market share and revenue losses for the applicable products, and which may result in a non-cash impairment charge in any associated intangible asset. In addition, from time to time, the Company's subsidiaries may settle these types of actions and such settlements can involve the introduction of generic versions of the products at issue to the market prior to the expiration of the relevant patents.

The Inter Partes Review (IPR) process with the United States Patent and Trademark Office (USPTO), created under the 2011 America Invents Act, is also being used at times by generic companies in conjunction with ANDAs and lawsuits to challenge the applicable patents.

XARELTO

Beginning in March 2021, Janssen Pharmaceuticals, Inc.; Bayer Pharma AG; Bayer AG; and Bayer Intellectual Property GmbH filed patent infringement lawsuits in United States district courts against generic manufacturers who have filed ANDAs seeking approval to market generic versions of XARELTO before expiration of certain Orange Book Listed Patents. The following entities are named defendants: Dr. Reddy's Laboratories, Inc.; Dr. Reddy's Laboratories, Ltd.; Lupin Limited; Lupin Pharmaceuticals, Inc.; Taro Pharmaceutical Industries Ltd.; Taro Pharmaceuticals U.S.A., Inc.; Teva Pharmaceuticals USA, Inc.; Mylan Pharmaceuticals Inc.; Mylan Inc.; USV Private Limited; Mankind Pharma Limited; Apotex Inc.; Apotex Corp.; Biocon Pharma Limited; Biocon Limited; Biocon Pharma, Inc.; ScieGen Pharmaceuticals, Inc.; Auson Pharmaceuticals Inc.; Macleods Pharmaceuticals Ltd; Macleods Pharma USA, Inc.; Indoco Remedies Limited; and FPP Holding Company LLC. The following U.S. patents are included in one or more cases: 9,539,218; and 10,828,310. In April 2023, the Company entered into a confidential settlement agreement with ScieGen Pharmaceuticals, Inc. In June 2023, the Company entered into a confidential settlement agreement with USV Private Limited.

U.S. Patent No. 10,828,310 is also under consideration by the USPTO in an IPR proceeding.

OPSUMIT

Beginning in January 2023 Actelion Pharmaceuticals Ltd and Actelion Pharmaceuticals US, Inc. filed patent infringement lawsuits in United States district courts against generic manufacturers who have filed ANDAs seeking approval to market generic versions of OPSUMIT before expiration of certain Orange Book Listed Patents. The following entities are named defendants: Sun Pharmaceutical Industries Limited; Sun Pharmaceutical Industries, Inc.; Alembic Pharmaceuticals Ltd.; Alembic Pharmaceuticals, Inc.; MSN Laboratories Private Limited; MSN Pharmaceuticals Inc.; Apotex Inc.; and Apotex Corp. The following U.S. patents are included in one or more cases: 7,094,781; and 10,946,015.

INVEGA SUSTENNA

Beginning in January 2018, Janssen Pharmaceutica NV and Janssen Pharmaceuticals, Inc. filed patent infringement lawsuits in United States district courts against generic manufacturers who have filed ANDAs seeking approval to market generic versions of INVEGA SUSTENNA before expiration of the Orange Book Listed Patent. The following entities are named defendants: Teva Pharmaceuticals USA, Inc.; Mylan Laboratories Limited; Pharmascience Inc.; Mallinckrodt PLC; Specgx LLC; Tolmar, Inc.; and Accord Healthcare, Inc. The following U.S. patent is included in one or more cases: 9,439,906.

Beginning in February 2018, Janssen Inc. and Janssen Pharmaceutica NV initiated a Statement of Claim under Section 6 of the Patented Medicines (Notice of Compliance) Regulations against generic manufacturers who have filed ANDAs seeking approval to market generic versions of INVEGA SUSTENNA before expiration of the listed patent. The following entities are named defendants: Teva Canada Limited; Pharmascience Inc.; and Apotex Inc. The following Canadian patent is included in one or more cases: 2,655,335. In March 2023, the Federal Appeals Court dismissed Teva Canada Limited's appeal of the Federal Court's decision finding that its proposed generic product infringes the asserted patent and that the patent is not invalid.

INVEGA TRINZA

Beginning in September 2020, Janssen Pharmaceuticals, Inc., Janssen Pharmaceutica NV, and Janssen Research & Development, LLC filed patent infringement lawsuits in United States district courts against generic manufacturers who have filed ANDAs seeking approval to market generic versions of INVEGA TRINZA before expiration of the Orange Book Listed Patent. The following entities are named defendants: Mylan Laboratories Limited; Mylan Pharmaceuticals Inc.; and Mylan Institutional LLC. The following U.S. patent is included in one or more cases: 10,143,693. In May 2023, the District Court

issued a decision finding that Mylan's proposed generic product infringes the asserted patent and that the patent is not invalid. Mylan has appealed the verdict.

IMBRUVICA

Beginning in September 2021, Pharmacyclics LLC and Janssen Inc. initiated a Statement of Claim under Section 6 of the Patented Medicines (Notice of Compliance) Regulations against generic manufacturers who have filed ANDSs seeking approval to market generic versions of IMBRUVICA before expiration of certain listed patents. The following entities are named defendants: Natco Pharma (Canada) Inc. (Natco); and Sandoz Canada Inc. (Sandoz). The following patents are included in one or more cases: 2,663,116; 2,928,721; 2,800,913; 3,007,787; 3,007,788; 2,875,986; and 3,022,256. In July 2023, the Company entered into confidential settlement agreements with Natco and Sandoz.

SYM TUZA

Beginning in November 2021, Janssen Products, L.P., Janssen Sciences Ireland Unlimited Company, Gilead Sciences, Inc. and Gilead Sciences Ireland UC filed patent infringement lawsuits in United States district courts against generic manufacturers who have filed ANDAs seeking approval to market generic versions of SYMTUZA before expiration of certain Orange Book Listed Patents. The following entities are named defendants: Lupin Limited; Lupin Pharmaceuticals, Inc.; MSN Laboratories Private Ltd.; MSN Life Sciences Private Ltd.; MSN Pharmaceuticals Inc.; Apotex Inc.; and Apotex Corp. The following U.S. patents are included in one or more cases: 10,039,718 and 10,786,518.

ERLEADA

Beginning in May 2022, Aragon Pharmaceuticals, Inc., Janssen Biotech, Inc., Sloan Kettering Institute for Cancer Research and The Regents of the University of California filed patent infringement lawsuits in United States district courts against generic manufacturers who have filed ANDAs seeking approval to market generic versions of ERLEADA before expiration of certain Orange Book Listed Patents. The following entities are named defendants: Lupin Limited; Lupin Pharmaceuticals, Inc.; Zydus Worldwide DMCC; Zydus Pharmaceuticals (USA), Inc.; Zydus Lifesciences Limited; Sandoz Inc.; Eugia Pharma Specialities Limited; Aurobindo Pharma USA, Inc.; Auromedics Pharma LLC; Hetero Labs Limited Unit V; and Hetero USA, Inc. The following U.S. patents are included in one or more cases: 9,481,663; 9,884,054; 10,052,314; 10,702,508; 10,849,888; 8,445,507; 8,802,689; 9,388,159; 9,987,261; and RE49,353.

UPTRAVI

Beginning in November 2022, Actelion Pharmaceuticals US Inc., Actelion Pharmaceuticals Ltd and Nippon Shinyaku Co., Ltd. filed patent infringement lawsuits in United States district courts against generic manufacturers who have filed ANDAs seeking approval to market generic versions of UPTRAVI before expiration of certain Orange Book Listed Patents. The following entities are named defendants: Alembic Pharmaceuticals Limited, Alembic Pharmaceuticals Inc.; Lupin Ltd.; Lupin Pharmaceuticals, Inc.; Cipla Limited; Cipla USA Inc.; MSN Laboratories Private Ltd.; and MSN Pharmaceuticals Inc. The following U.S. patents are included in one or more cases: 8,791,122; 9,284,280; and 7,205,302 ('302 patent). In June 2023, the Company reached a confidential settlement agreement with respect to the '302 patent with Alembic Pharmaceuticals Limited and Alembic Pharmaceuticals Inc.

SPRAVATO

Beginning in May 2023, Janssen Pharmaceuticals, Inc. and Janssen Pharmaceutica NV filed patent infringement lawsuits in United States district courts against generic manufacturers who have filed ANDAs seeking approval to market generic versions of SPRAVATO before expiration of certain Orange Book Listed Patents. The following entities are named defendants: Sandoz Inc.; Hikma Pharmaceuticals Inc. USA; Hikma Pharmaceuticals PLC; Westward Columbus Inc.; Alkem Laboratories Ltd; and Ascend Laboratories, LLC. The following U.S. patents are included in one or more cases: 10,869,844; 11,173,134; 11,311,500; and 11,446,260.

GOVERNMENT PROCEEDINGS

Like other companies in the pharmaceutical, consumer health and medical devices industries, the Company and certain of its subsidiaries are subject to extensive regulation by national, state and local government agencies in the United States and other countries in which they operate. Such regulation has been the basis of government investigations and litigations. The most significant litigation brought by, and investigations conducted by, government agencies are listed below. It is possible that criminal charges and substantial fines and/or civil penalties or damages could result from government investigations or litigation.

MedTech

In August 2012, DePuy Orthopaedics, Inc., DePuy, Inc. (now known as DePuy Synthes, Inc.), and Johnson & Johnson Services, Inc. (collectively DePuy) received an informal request from the United States Attorney's Office for the District of Massachusetts and the Civil Division of the United States Department of Justice (the United States) for the production of materials relating to the DePuy ASR XL Hip device. In July 2014, the United States notified the United States District Court for

the District of Massachusetts that it had declined to intervene in a *qui tam* case filed pursuant to the False Claims Act against the companies concerning the hip devices. In February 2016, the District Court granted the companies' motion to dismiss with prejudice, unsealed the *qui tam* complaint, and denied the *qui tam* relators' request for leave to file a further amended complaint. The *qui tam* relators appealed the case to the United States Court of Appeals for the First Circuit. In July 2017, the First Circuit affirmed the District Court's dismissal in part, reversed in part, and affirmed the decision to deny the relators' request to file a third amended complaint. In March 2021, DePuy filed its motion to strike and dismiss the relators' second amended complaint; the District Court denied DePuy's motion to strike and dismiss in July 2021. DePuy filed a motion for reconsideration of the District Court's July 2021 ruling. In November 2021, the District Court granted DePuy's motion for reconsideration and dismissed the case with prejudice. The District Court's order was unsealed in December 2021. The relators filed several post-dismissal motions, including a January 2022 omnibus motion for reconsideration, which the District Court denied. Following the District Court's order dismissing the case with prejudice, DePuy filed a December 2021 motion seeking the recovery of attorneys' fees and costs, which the District Court denied except as to costs. The Relators appealed the District Court's dismissal of the case to the First Circuit. On May 18, 2023, the First Circuit affirmed the District Court's dismissal of the case.

In October 2012, the Company was contacted by the California Attorney General's office regarding a multi-state Attorney General investigation of the marketing of surgical mesh products for hernia and urogynecological purposes by the Company's subsidiary, Ethicon, Inc. (Ethicon). In May 2016, California and Washington filed civil complaints against the Company, Ethicon and Ethicon US, LLC alleging violations of their consumer protection statutes. Similar complaints were filed against the companies by the following states: Kentucky, Mississippi, West Virginia and Oregon. In October 2019, the Company and Ethicon settled the multi-state investigation with 41 other states and the District of Columbia. Between April 2019 and February 2023, the Company settled with Washington, West Virginia, Oregon, Mississippi and Kentucky. After a trial in California and various appeals, the Company satisfied the judgment of penalties in California.

In July 2018, the Public Prosecution Service in Rio de Janeiro and representatives from the Brazilian antitrust authority CADE inspected the offices of more than 30 companies including Johnson & Johnson do Brasil Indústria e Comércio de Produtos para Saúde Ltda. The authorities appear to be investigating allegations of possible anti-competitive behavior and possible improper payments in the medical device industry. The Company continues to respond to inquiries regarding the Foreign Corrupt Practices Act from the United States Department of Justice and the United States Securities and Exchange Commission.

Pharmaceuticals

The Company and several of its pharmaceutical subsidiaries (the J&J AWP Defendants), along with numerous other pharmaceutical companies, were named as defendants in a series of lawsuits in state and federal courts involving allegations that the pricing and marketing of certain pharmaceutical products amounted to fraudulent and otherwise actionable conduct because, among other things, the companies allegedly reported an inflated Average Wholesale Price (AWP) for the drugs at issue. Payors alleged that they used those AWP's in calculating provider reimbursement levels. The plaintiffs in these cases included three classes of private persons or entities that paid for any portion of the purchase of the drugs at issue based on AWP, and state government entities that made Medicaid payments for the drugs at issue based on AWP. Many of these cases, both federal actions and state actions removed to federal court, were consolidated for pre-trial purposes in a multi-district litigation in the United States District Court for the District of Massachusetts, where all claims against the J&J AWP Defendants were ultimately dismissed. The J&J AWP Defendants also prevailed in a case brought by the Commonwealth of Pennsylvania. Other AWP cases have been resolved through court order or settlement. The case brought by Illinois was settled after trial. In New Jersey, a putative class action based upon AWP allegations is pending against Centocor, Inc. and Ortho Biotech Inc. (both now Janssen Biotech, Inc.), the Company and ALZA Corporation. In June 2023, an action alleging AWP-related claims was filed in a Pennsylvania state court against the same Company subsidiaries by a putative class of payors.

In July 2016, the Company and Janssen Products, LP were served with a *qui tam* complaint pursuant to the False Claims Act filed in the United States District Court for the District of New Jersey alleging the off-label promotion of two HIV products, PREZISTA and INTELENCE, and anti-kickback violations in connection with the promotion of these products. The complaint was filed under seal in December 2012. The federal and state governments have declined to intervene, and the lawsuit is being prosecuted by the relators. The Court denied summary judgment on all claims in December 2021. Daubert motions were granted in part and denied in part in January 2022, and the case is proceeding to trial.

In March 2017, Janssen Biotech, Inc. (JBI) received a Civil Investigative Demand from the United States Department of Justice regarding a False Claims Act investigation concerning management and advisory services provided to rheumatology and gastroenterology practices that purchased REMICADE or SIMPONI ARIA. In August 2019, the United States Department of Justice notified JBI that it was closing the investigation. Subsequently, the United States District Court for the District of Massachusetts unsealed a *qui tam* False Claims Act complaint, which was served on the Company. The Department of Justice had declined to intervene in the *qui tam* lawsuit in August 2019. The Company filed a motion to dismiss, which was granted in part and denied in part. Discovery is underway.

From time to time, the Company has received requests from a variety of United States Congressional Committees to produce information relevant to ongoing congressional inquiries. It is the policy of Johnson & Johnson to cooperate with these inquiries by producing the requested information.

GENERAL LITIGATION

The Company (subsequently substituted by Johnson & Johnson Consumer Inc. (JJCI)) along with more than 120 other companies, is a defendant in a cost recovery and contribution action brought by Occidental Chemical Corporation in June 2018 in the United States District Court for the District of New Jersey, related to the clean-up of a section of the Lower Passaic River in New Jersey.

The Company or its subsidiaries are also parties to various proceedings brought under the Comprehensive Environmental Response, Compensation, and Liability Act, commonly known as Superfund, and comparable state, local or foreign laws in which the primary relief sought is the Company's agreement to implement remediation activities at designated hazardous waste sites or to reimburse the government or third parties for the costs they have incurred in performing remediation as such sites.

In October 2017, certain United States service members and their families brought a complaint against a number of pharmaceutical and medical devices companies, including Johnson & Johnson and certain of its subsidiaries in United States District Court for the District of Columbia, alleging that the defendants violated the United States Anti-Terrorism Act. The complaint alleges that the defendants provided funding for terrorist organizations through their sales practices pursuant to pharmaceutical and medical device contracts with the Iraqi Ministry of Health. In July 2020, the District Court dismissed the complaint. In January 2022, the United States Court of Appeals for the District of Columbia Circuit reversed the District Court's decision. In June 2023, defendants filed a petition for a writ of certiorari to the United States Supreme Court.

MedTech

In October 2020, Fortis Advisors LLC (Fortis), in its capacity as representative of the former stockholders of Auris Health Inc. (Auris), filed a complaint against the Company, Ethicon Inc., and certain named officers and employees (collectively, Ethicon) in the Court of Chancery of the State of Delaware. The complaint alleges breach of contract, fraud, and other causes of action against Ethicon in connection with Ethicon's acquisition of Auris in 2019. The complaint seeks damages and other relief. In December 2021, the Court granted in part and denied in part defendants' motion to dismiss certain causes of action. All claims against the individual defendants were dismissed. The trial is scheduled for January 2024.

Pharmaceuticals

In June 2019, the United States Federal Trade Commission (FTC) issued a Civil Investigative Demand to the Company and Janssen Biotech, Inc. (collectively, Janssen) in connection with its investigation of whether Janssen's REMICADE contracting practices violate federal antitrust laws. The Company has produced documents and information responsive to the Civil Investigative Demand. Janssen is in ongoing discussions with the FTC staff regarding its inquiry.

In February 2022, the United States Federal Trade Commission (FTC) issued Civil Investigative Demands to Johnson & Johnson and Janssen Biotech, Inc. (collectively, Janssen) in connection with its investigation of whether advertising practices for REMICADE violate federal law. Janssen has produced documents and information responsive to the Civil Investigative Demands. Janssen is in ongoing discussions with the FTC staff regarding the inquiry.

In June 2022, Genmab A/S filed a Notice for Arbitration with International Institute for Conflict Prevention and Resolution (CPR) against Janssen Biotech, Inc. seeking milestones and an extended royalty term for Darzalex FASPRO. In April 2023, the Arbitration Panel ruled in Janssen's favor and dismissed Genmab's claims. In May 2023, Genmab appealed that award.

In October 2018, two separate putative class actions were filed against Actelion Pharmaceutical Ltd., Actelion Pharmaceuticals U.S., Inc., and Actelion Clinical Research, Inc. (collectively Actelion) in United States District Court for the District of Maryland and United States District Court for the District of Columbia. The complaints allege that Actelion violated state and federal antitrust and unfair competition laws by allegedly refusing to supply generic pharmaceutical manufacturers with samples of TRACLEER. TRACLEER is subject to a Risk Evaluation and Mitigation Strategy required by the U.S. Food and Drug Administration, which imposes restrictions on distribution of the product. In January 2019, the plaintiffs dismissed the District of Columbia case and filed a consolidated complaint in the United States District Court for the District of Maryland.

In June 2022, Janssen Pharmaceuticals, Inc. filed a Demand for Arbitration against Emergent Biosolutions Inc. et al (EBSI) with the American Arbitration Association, alleging that EBSI breached the parties' Manufacturing Services Agreement for the Company's COVID-19 vaccine. In July 2022, Emergent filed its answering statement and counterclaims. The hearing is scheduled for March 2024.

In October 2022, Janssen Pharmaceuticals, Inc. (Janssen) filed a Demand for Arbitration against Merck Sharp & Dohme Corp. (Merck) with the American Arbitration Association pursuant to the Parties' agreements relating to production of drug substance and drug product for the Company's COVID-19 vaccine. Janssen and Merck settled the matter in May 2023.

Consumer Health

In November 2019, the Company received a demand for indemnification from Pfizer Inc. (Pfizer), pursuant to the 2006 Stock and Asset Purchase Agreement between the Company and Pfizer. Also in November 2019, Johnson & Johnson Inc. received notice reserving rights to claim indemnification from Sanofi Consumer Health, Inc. (Sanofi), pursuant to the 2016 Asset Purchase Agreement between Johnson & Johnson Inc. and Sanofi. In January 2020, Johnson & Johnson received a demand for indemnification from Boehringer Ingelheim Pharmaceuticals, Inc. (Boehringer Ingelheim), pursuant to the 2006 Asset Purchase Agreement among the Company, Pfizer, and Boehringer Ingelheim. In November 2022, Johnson & Johnson received a demand for indemnification from GlaxoSmithKline LLC (GSK), pursuant to the 2006 Stock and Asset Purchase Agreement between the Company and Pfizer, and certain 1993, 1998, and 2002 agreements between Glaxo Wellcome and Warner-Lambert entities. The notices seek indemnification for legal claims related to over-the-counter ZANTAC (ranitidine) products. Plaintiffs in the underlying actions allege that ZANTAC and other over-the-counter ranitidine medications contain unsafe levels of NDMA (N-nitrosodimethylamine) and can cause and/or have caused various cancers in patients using the products, and seek injunctive and monetary relief. The Company and Johnson & Johnson Inc. have also been named in putative class actions filed in Canada with similar allegations regarding ZANTAC or ranitidine use. Johnson & Johnson Inc. was also named as a defendant along with other manufacturers in various personal injury actions in Canada related to ZANTAC products. Johnson & Johnson Inc. has provided Sanofi notice reserving rights to claim indemnification pursuant to the 2016 Asset Purchase Agreement related to the class actions and personal injury actions.

Beginning in May 2021, multiple putative class actions were filed in state and federal courts (California, Florida, New York, and New Jersey) against various Johnson & Johnson entities alleging violations of state consumer fraud statutes based on nondisclosure of alleged benzene contamination of certain Neutrogena and Aveeno sunscreen products and the affirmative promotion of those products as "safe"; and, in at least one case, alleging a strict liability manufacturing defect and failure to warn claims, asserting that the named plaintiffs suffered unspecified injuries as a result of alleged exposure to benzene. The Judicial Panel on Multi-District Litigation has consolidated all pending actions, except one product liability case and one case pending in New Jersey state court, in the United States District Court for the Southern District of Florida, Fort Lauderdale Division. In October 2021, the Company reached an agreement in principle for the settlement of a nationwide class, encompassing the claims of the consolidated actions, subject to approval by the Florida federal Court. In December 2021, plaintiffs in the consolidated actions filed a motion for preliminary approval of a nationwide class settlement. The court issued an order granting final approval of the settlement in February 2023. A Notice of Appeal was filed in April 2023.

NOTE 12— KENVUE SEPARATION

During the fiscal six months of 2023 the financial statements and accompanying footnotes continue to reflect the Company's consolidation of Kenvue.

On May 8, 2023, Kenvue, completed an initial public offering (the IPO) of 198,734,444 shares of its common stock, par value \$0.01 per share (the “Kenvue Common Stock”), including the underwriters’ full exercise of their option to purchase 25,921,884 shares to cover over-allotments, at an initial public offering price of \$22.00 per share for net proceeds of \$4.2 billion. Kenvue shares began trading on the New York Stock Exchange under the symbol “KVUE”. Additionally, on May 8, 2023, as partial consideration for the transfer of the Consumer Health business, Kenvue paid \$13.2 billion to Johnson & Johnson from the net proceeds of the initial public offering and debt financing transactions (approximately \$7.7 billion of unsecured notes and \$1.2 billion of commercial paper) in connection with the separation (See Note 4 to the Consolidated Financial Statements for additional details). The excess of the net proceeds from the IPO over the net book value of the Johnson & Johnson divested interest was \$2.5 billion and was recorded in additional paid-in capital.

As of the closing of the IPO, Johnson & Johnson owns 1,716,160,000 shares of Kenvue Common Stock, or approximately 89.6% of the total outstanding shares of Kenvue Common Stock, and continues to consolidate the financial results of Kenvue. As of July 2, 2023, the non-controlling interest of \$1.3 billion associated with Kenvue is reflected in equity attributable to non-controlling interests in the consolidated balance sheet. Other (income) expense, net on the consolidated statement of earnings for the fiscal second quarter and fiscal six months ended July 2, 2023, includes \$37 million related to the 10.4% non-controlling interest in Kenvue from the time of the initial public offering on May 8, 2023 through the end of the fiscal second quarter.

Subsequent to the fiscal second quarter on July 24, 2023, the Company announced its intention to split-off at least 80.1% of the shares of Kenvue through an exchange offer. Through the planned exchange offer, Johnson & Johnson shareholders can exchange all, some or none of their shares of Johnson & Johnson Common Stock for shares of Kenvue Common Stock, subject to the terms of the offer. The exchange offer will permit Johnson & Johnson shareholders to exchange some, all or none of their shares of Johnson & Johnson Common Stock for shares of Kenvue Common Stock at a 7% discount, subject to an upper limit of 8.0549 shares of Kenvue Common Stock per share of Johnson & Johnson Common Stock tendered and accepted in the exchange offer. If the upper limit is not in effect, tendering shareholders are expected to receive approximately \$107.53 of Kenvue Common Stock for every \$100 of Johnson & Johnson Common Stock tendered. Once the exchange offer is consummated (including all related transactions), the Company will no longer be the majority shareholder of Kenvue, and Kenvue will be a separate and independent company. The shares of Johnson & Johnson Common Stock acquired by the Company in the exchange offer will be recorded as an acquisition of treasury stock at a cost equal to the market value of the shares of Johnson & Johnson Common Stock accepted in the exchange offer at its expiration. Any difference between the net book value of Kenvue attributable to Johnson & Johnson and the market value of the shares of Johnson & Johnson Common Stock acquired at that date will be recognized by the Company as a gain on disposal of discontinued operations net of any direct and incremental expenses of the exchange offer on the disposal of its Kenvue Common Stock. Upon completion of the exchange offer, and assuming the Company no longer has a controlling financial interest in Kenvue, Kenvue’s historical results will be shown in the Company’s financial statements as discontinued operations, and, in subsequent periods, the Company’s financial statements will no longer reflect the assets, liabilities, results of operations or cash flows attributable to Kenvue.

Item 2 — MANAGEMENT’S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

RESULTS OF OPERATIONS

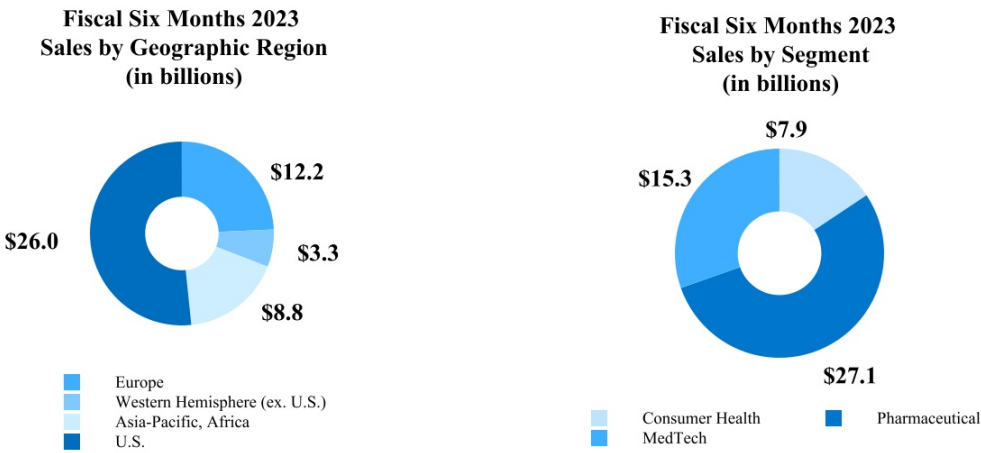
Sales to Customers

Analysis of Consolidated Sales

For the fiscal six months of 2023, worldwide sales were \$50.3 billion, a total increase of 6.0%, including an operational increase of 8.2% as compared to 2022 fiscal six months sales of \$47.4 billion. Currency fluctuations had a negative impact of 2.2% for the fiscal six months of 2023. In the fiscal six months of 2023, the net impact of acquisitions and divestitures on worldwide operational sales growth was a positive 1.3%.

Sales by U.S. companies were \$26.0 billion in the fiscal six months of 2023, which represented an increase of 10.0% as compared to the prior year. In the fiscal six months of 2023, the net impact of acquisitions and divestitures on the U.S. operational sales growth was a positive 2.3%. Sales by international companies were \$24.3 billion, an increase of 2.0%, including an operational increase of 6.5%, and a negative currency impact of 4.5% as compared to the fiscal six months sales of 2022. In the fiscal six months of 2023, the net impact of acquisitions and divestitures on the international operational sales growth was a positive 0.3%.

In the fiscal six months of 2023, sales by companies in Europe achieved growth of 1.0%, which included an operational increase of 3.0% and a negative currency impact of 2.0%. Sales by companies in the Western Hemisphere, excluding the U.S., achieved growth of 9.3%, which included an operational increase of 16.1%, and a negative currency impact of 6.8%. Sales by companies in the Asia-Pacific, Africa region achieved growth of 0.9%, including an operational increase of 8.2% and a negative currency impact of 7.3%.



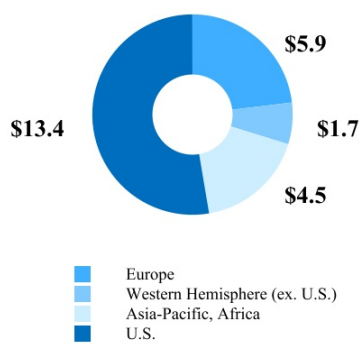
Note: values may have been rounded

For the fiscal second quarter of 2023, worldwide sales were \$25.5 billion, a total increase of 6.3%, which included operational growth of 7.5% and a negative currency impact of 1.2% as compared to 2022 fiscal second quarter sales of \$24.0 billion. In the fiscal second quarter of 2023, the net impact of acquisitions and divestitures on worldwide operational sales growth was a positive 1.3%.

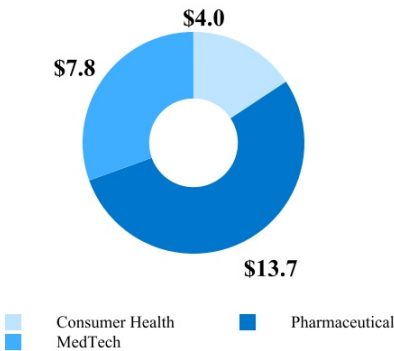
Sales by U.S. companies were \$13.4 billion in the fiscal second quarter of 2023, which represented an increase of 10.2% as compared to the prior year. In the fiscal second quarter of 2023, the net impact of acquisitions and divestitures on the U.S. operational sales growth was a positive 2.2%. Sales by international companies were \$12.1 billion, a total increase of 2.2%, which included operational growth of 4.7% and a negative currency impact of 2.5%. In the fiscal second quarter of 2023, the net impact of acquisitions and divestitures on the international operational sales growth was a positive 0.3%.

In the fiscal second quarter of 2023, sales by companies in Europe experienced a decline of 3.1%, which included an operational decline of 3.9% and a positive currency impact of 0.8%. Sales by companies in the Western Hemisphere, excluding the U.S., achieved growth of 11.5%, including operational growth of 17.7% and a negative currency impact of 6.2%. Sales by companies in the Asia-Pacific, Africa region achieved growth of 6.6%, including operational growth of 12.5% and a negative currency impact of 5.9%.

Q2 2023 Sales by Geographic Region
(in billions)



Q2 2023 Sales by Segment
(in billions)



Note: values may have been rounded

Analysis of Sales by Business Segments

Consumer Health

Consumer Health segment sales in the fiscal six months of 2023 were \$7.9 billion, an increase of 6.4% as compared to the same period a year ago, including operational growth of 9.5% and a negative currency impact of 3.1%. U.S. Consumer Health segment sales increased by 8.6%. International Consumer Health segment sales increased by 4.7%, including operational growth of 10.1% and a negative currency impact of 5.4%. In the fiscal six months of 2023, the net impact of acquisitions and divestitures on the Consumer Health segment operational sales growth was negligible.

Major Consumer Health Franchise Sales — Fiscal Six Months Ended

(Dollars in Millions)	July 2, 2023	July 3, 2022	Total Change	Operations Change	Currency Change
OTC	\$ 3,301	\$ 2,943	12.2 %	14.8 %	(2.6) %
Skin Health/Beauty	2,258	2,138	5.6	8.0	(2.4)
Oral Care	759	760	(0.2)	2.4	(2.6)
Baby Care	719	730	(1.6)	3.1	(4.7)
Women's Health	455	458	(0.6)	7.1	(7.7)
Wound Care/Other	371	361	2.6	4.9	(2.3)
Total Consumer Health Sales	\$ 7,863	\$ 7,391	6.4 %	9.5 %	(3.1) %

Consumer Health segment sales in the fiscal second quarter of 2023 were \$4.0 billion, an increase of 5.4% as compared to the same period a year ago, including operational growth of 7.7% and a negative currency impact of 2.3%. U.S. Consumer Health segment sales increased by 6.0%. International Consumer Health segment sales increased by 5.0% including operational growth of 9.0% and a negative currency impact of 4.0%. In the fiscal second quarter of 2023, the net impact of acquisitions and divestitures on the Consumer Health segment operational sales growth was negligible.

Major Consumer Health Franchise Sales — Fiscal Second Quarter Ended

(Dollars in Millions)	July 2, 2023	July 3, 2022	Total Change	Operations Change	Currency Change
OTC	\$ 1,659	\$ 1,482	12.0 %	13.9 %	(1.9) %
Skin Health/Beauty	1,148	1,126	2.0	3.5	(1.5)
Oral Care	398	394	0.9	2.7	(1.8)
Baby Care	360	375	(4.0)	(0.2)	(3.8)
Women's Health	238	230	3.4	10.0	(6.6)
Wound Care/Other	207	197	4.9	6.8	(1.9)
Total Consumer Health Sales	\$ 4,011	\$ 3,805	5.4 %	7.7 %	(2.3) %

The OTC franchise achieved operational growth of 13.9% as compared to the prior year fiscal second quarter. The growth was driven by price actions and strong TYLENOL and MOTRIN performance due to increased global Cough/Cold/Flu incidences.

The Skin Health/Beauty franchise achieved operational growth of 3.5% as compared to the prior year fiscal second quarter. The growth was driven by price actions and strength in NEUTROGENA particularly in Sun Care, driven by innovation and supply repositioning.

The Oral Care franchise achieved operational growth of 2.7% as compared to the prior year fiscal second quarter. The growth was driven by price actions, partially offset by category deceleration in Asia and the negative impact from the suspension of personal care sales in Russia.

The Baby Care franchise experienced an operational decline of 0.2% as compared to the prior year fiscal second quarter. The decline was driven by competitive pressures in Asia and the negative impact from the suspension of personal care sales in Russia.

The Women's Health franchise achieved operational growth of 10.0% as compared to the prior year fiscal second quarter primarily driven by price actions and continued strong growth in India partially offset by the negative impact from the suspension of personal care sales in Russia.

The Wound Care/Other franchise achieved operational growth of 6.8% as compared to the prior year fiscal second quarter primarily driven by price actions and innovation.

In November 2021, the Company announced its intention to separate the Company's Consumer Health business (Kenvue, the New Consumer Health Company), with the intention to create a new, publicly traded company by the end of the fiscal year 2023, pending market conditions. On May 8, 2023, Kenvue completed an initial public offering (the IPO). As of the closing of the IPO, the Company owns approximately 89.6% of the total outstanding shares of Kenvue Common Stock, and has continued to consolidate the financial results of Kenvue (See Note 12 to the Consolidated Financial Statements for additional details).

Pharmaceutical

Pharmaceutical segment sales in the fiscal six months of 2023 were \$27.1 billion, an increase of 3.7% as compared to the same period a year ago, with an operational increase of 5.5% and a negative currency impact of 1.8%. U.S. Pharmaceutical sales increased 7.6% as compared to the same period a year ago. International Pharmaceutical sales decreased by 0.7%, including operational growth of 3.1% offset by a negative currency impact of 3.8%. In the fiscal six months of 2023, the net impact of acquisitions and divestitures on the Pharmaceutical segment operational sales growth was a negative 0.1%.

Major Pharmaceutical Therapeutic Area Sales** — Fiscal Six Months Ended

(Dollars in Millions)	July 2, 2023	July 3, 2022	Total Change	Operations Change	Currency Change
Immunology	\$ 8,608	\$ 8,530	0.9 %	2.6 %	(1.7)%
REMICADE	949	1,310	(27.5)	(26.3)	(1.2)
SIMPONI/ SIMPONI ARIA	1,066	1,137	(6.2)	(3.3)	(2.9)
STELARA	5,241	4,887	7.2	8.7	(1.5)
TREMFYA	1,346	1,187	13.4	15.0	(1.6)
Other Immunology	7	9	(30.2)	(30.2)	—
Infectious Diseases	2,707	2,613	3.6	4.7	(1.1)
COVID-19 VACCINE	1,032	1,001	3.0	4.2	(1.2)
EDURANT/rilpivirine	546	473	15.6	17.1	(1.5)
PREZISTA/ PREZCOBIX/ REZOLSTA/ SYMTUZA	968	965	0.3	0.7	(0.4)
Other Infectious Diseases	161	174	(7.6)	(3.6)	(4.0)
Neuroscience	3,597	3,475	3.5	5.8	(2.3)
CONCERTA/methylphenidate	414	318	30.0	35.3	(5.3)
INVEGA SUSTENNA/ XEPLION/ INVEGA TRINZA/ TREVICTA	2,075	2,102	(1.3)	0.1	(1.4)
SPRAVATO	300	155	93.1	93.8	(0.7)
Other Neuroscience ⁽¹⁾	809	900	(10.1)	(6.4)	(3.7)
Oncology	8,510	7,992	6.5	8.8	(2.3)
CARVYKTI	189	24	*	*	*
DARZALEX	4,695	3,842	22.2	24.5	(2.3)
ERLEADA	1,109	850	30.4	33.2	(2.8)
IMBRUVICA	1,668	2,008	(16.9)	(15.0)	(1.9)
ZYTIGA/ abiraterone acetate	472	1,044	(54.8)	(52.3)	(2.5)
Other Oncology	376	224	68.0	70.4	(2.4)
Pulmonary Hypertension	1,844	1,695	8.8	10.8	(2.0)
OPSUMIT	947	881	7.5	9.5	(2.0)
UPTRAVI	761	653	16.6	17.5	(0.9)
Other Pulmonary Hypertension	136	161	(15.5)	(9.5)	(6.0)
Cardiovascular / Metabolism / Other	1,877	1,882	(0.3)	0.4	(0.7)
XARELTO	1,215	1,117	8.8	8.8	—
Other ⁽²⁾	662	765	(13.5)	(11.9)	(1.6)
Total Pharmaceutical Sales	\$ 27,144	\$ 26,186	3.7 %	5.5 %	(1.8)%

* Percentage greater than 100% or not meaningful

**Certain prior year amounts have been reclassified to conform to current year presentation

⁽¹⁾Inclusive of RISPERDAL CONSTA which was previously disclosed separately

⁽²⁾Inclusive of INVOKANA which was previously disclosed separately

Pharmaceutical segment sales in the fiscal second quarter of 2023 were \$13.7 billion, an increase of 3.1% as compared to the same period a year ago, including an operational increase of 3.8% and a negative currency impact of 0.7%. U.S. Pharmaceutical sales increased 9.2% as compared to the same period a year ago. International Pharmaceutical sales decreased by 4.0%, including an operational decline of 2.5% and a negative currency impact of 1.5%. In the fiscal second quarter of 2023, the net impact of acquisitions and divestitures on the Pharmaceutical segment operational sales growth was a negative 0.1%.

Major Pharmaceutical Therapeutic Area Sales** — Fiscal Second Quarter Ended

(Dollars in Millions)	July 2, 2023	July 3, 2022	Total Change	Operations Change	Currency Change
Immunology	\$ 4,496	\$ 4,411	1.9 %	2.6 %	(0.7)%
REMICADE	462	647	(28.6)	(27.6)	(1.0)
SIMPONI/ SIMPONI ARIA	529	566	(6.6)	(4.7)	(1.9)
STELARA	2,797	2,599	7.6	8.0	(0.4)
TREMFYA	706	597	18.3	18.9	(0.6)
Other Immunology	4	3	17.8	17.8	—
Infectious Diseases	1,121	1,316	(14.8)	(16.6)	1.8
COVID-19 VACCINE	285	544	(47.7)	(51.8)	4.1
EDURANT/tilpivirine	266	225	18.6	16.2	2.4
PREZISTA/ PREZCOBIX/ REZOLSTA/ SYMTUZA	491	464	5.8	5.6	0.2
Other Infectious Diseases	79	83	(5.2)	1.3	(6.5)
Neuroscience	1,793	1,734	3.5	5.5	(2.0)
CONCERTA/ methylphenidate	208	161	28.6	32.4	(3.8)
INVEGA SUSTENNA/ XEPLION/ INVEGA TRINZA/ TREVICTA	1,031	1,054	(2.1)	(1.5)	(0.6)
SPRAVATO	169	85	98.2	98.5	(0.3)
Other Neuroscience ⁽¹⁾	386	433	(10.9)	(5.8)	(5.1)
Oncology	4,398	4,042	8.8	9.7	(0.9)
CARVYKTI	117	24	*	*	*
DARZALEX	2,431	1,986	22.4	23.4	(1.0)
ERLEADA	567	450	25.9	26.9	(1.0)
IMBRUVICA	841	970	(13.2)	(12.6)	(0.6)
ZYTIGA/ abiraterone acetate	227	505	(55.1)	(53.8)	(1.3)
Other Oncology	214	106	*	*	*
Pulmonary Hypertension	972	843	15.3	16.5	(1.2)
OPSUMIT	507	438	15.7	16.7	(1.0)
UPTRAVI	399	328	21.8	22.5	(0.7)
Other Pulmonary Hypertension	66	78	(14.7)	(9.7)	(5.0)
Cardiovascular / Metabolism / Other	950	972	(2.2)	(2.0)	(0.2)
XARELTO	637	609	4.7	4.7	—
Other ⁽²⁾	313	363	(13.8)	(13.4)	(0.4)
Total Pharmaceutical Sales	\$ 13,731	\$ 13,317	3.1 %	3.8 %	(0.7)%

* Percentage greater than 100% or not meaningful

**Certain prior year amounts have been reclassified to conform to current year presentation

⁽¹⁾ Inclusive of RISPERDAL CONSTA which was previously disclosed separately

⁽²⁾ Inclusive of INVOKANA which was previously disclosed separately

Immunology products achieved operational growth of 2.6% as compared to the same period a year ago driven by market growth of STELARA (ustekinumab) partially offset by unfavorable patient mix and rebates. Additionally, growth of TREMFYA (guselkumab) was due to share gains and market growth partially offset by unfavorable patient mix. Lower sales of REMICADE (infliximab) were due to biosimilar competition.

Biosimilar versions of REMICADE have been introduced in the United States and certain markets outside the United States and additional competitors continue to enter the market. Continued infliximab biosimilar competition will result in a further reduction in sales of REMICADE.

The latest expiring United States composition of matter patent for STELARA (ustekinumab) expires in September 2023. STELARA (ustekinumab) U.S. sales in fiscal 2022 were approximately \$6.4 billion. Third parties have filed abbreviated Biologics License Applications with the FDA seeking approval to market biosimilar versions of STELARA. In May 2023, the Company settled litigation with Amgen under the Biosimilar Price Competition and Innovation Act of 2009. As a result of the settlement and other agreements with separate third parties, the Company does not anticipate the launch of a biosimilar version of STELARA until January 1, 2025.

Infectious disease products experienced an operational decline of 16.6% as compared to the same period a year ago primarily driven by a decline in COVID-19 vaccine revenue.

Neuroscience products achieved operational sales growth of 5.5% as compared to the same period a year ago. The growth of SPRAVATO (esketamine) was driven by ongoing launches in the U.S. and Europe. Growth was partially offset by declines from the paliperidone long-acting injectables, due to the XEPLION loss of exclusivity in the European Union.

Oncology products achieved operational sales growth of 9.7% as compared to the same period a year ago. Strong sales of DARZALEX (daratumumab) were driven by share gains in all regions and continued market growth. Growth of ERLEADA (apalutamide) was due to continued strong share gains, market growth, and increased penetration from new launches. Sales of CARVYKTI (ciltacabtagene autoleucel) were driven by continued market share gains from the ongoing launch. Additionally, sales from the launch of TECVAYLI (teclistamab-cqyv), included in Other Oncology, contributed to the growth. Growth was partially offset by ZYTIGA (abiraterone acetate) due to loss of exclusivity and IMBRUVICA (ibrutinib) due to global competitive pressures.

Pulmonary Hypertension achieved operational sales growth of 16.5% as compared to the same period a year ago. Sales growth was due to favorable patient mix, share gains and market growth from UPTRAVI (selexipag) and OPSUMIT (macitentan) partially offset by declines in Other Pulmonary Hypertension.

Cardiovascular / Metabolism / Other products experienced an operational decline of 2.0% as compared to the same period a year ago. The growth of XARELTO (rivaroxaban) was primarily driven by favorable patient mix and market growth partially offset by share loss.

The Company maintains a policy that no end customer will be permitted direct delivery of product to a location other than the billing location. This policy impacts contract pharmacy transactions involving non-grantee 340B covered entities for most of the Company's drugs, subject to multiple exceptions. Both grantee and non-grantee covered entities can maintain certain contract pharmacy arrangements under policy exceptions. The Company has been and will continue to offer 340B discounts to covered entities on all of its covered outpatient drugs, and it believes its policy will improve its ability to identify inappropriate duplicate discounts and diversion prohibited by the 340B statute. The 340B Drug Pricing Program is a U.S. federal government program requiring drug manufacturers to provide significant discounts on covered outpatient drugs to covered entities. This policy had discount implications which positively impacted sales to customers in fiscal 2023.

MedTech

The MedTech segment sales in the fiscal six months of 2023 were \$15.3 billion, an increase of 10.1% as compared to the same period a year ago, with an operational increase of 12.8% and a negative currency impact of 2.7%. U.S. MedTech sales increased 15.5%. International MedTech sales increased by 5.2%, including an operational increase of 10.3% and a negative currency impact of 5.1%. In the fiscal six months of 2023, the net impact of acquisitions and divestitures on the MedTech segment operational sales growth was a positive 4.7% related to the Abiomed acquisition on December 22, 2022.

Major MedTech Franchise Sales** — Fiscal Six Months Ended

(Dollars in Millions)	July 2, 2023	July 3, 2022	Total Change	Operations Change	Currency Change
Surgery	\$ 5,028	\$ 4,884	3.0 %	6.3 %	(3.3) %
Advanced	2,340	2,302	1.7	5.0	(3.3)
General	2,688	2,582	4.1	7.4	(3.3)
Orthopaedics	4,510	4,345	3.8	5.4	(1.6)
Hips	787	777	1.3	2.8	(1.5)
Knees	731	688	6.3	7.9	(1.6)
Trauma	1,496	1,444	3.6	4.9	(1.3)
Spine, Sports & Other	1,495	1,436	4.2	6.1	(1.9)
Interventional Solutions	3,123	2,141	45.8	49.3	(3.5)
Electrophysiology	2,288	1,970	16.2	19.5	(3.3)
Abiomed	655	—	*	*	*
Other Interventional Solutions	180	171	5.1	9.5	(4.4)
Vision	2,608	2,498	4.4	7.2	(2.8)
Contact Lenses/Other	1,892	1,804	4.9	8.0	(3.1)
Surgical	716	694	3.1	5.4	(2.3)
Total MedTech Sales	\$ 15,269	\$ 13,869	10.1 %	12.8 %	(2.7) %

* Percentage greater than 100% or not meaningful

**Certain prior year amounts have been reclassified to conform to current year presentation

The MedTech segment sales in the fiscal second quarter of 2023 were \$7.8 billion, an increase of 12.9% as compared to the same period a year ago, which included operational growth of 14.7% and a negative currency impact of 1.8%. U.S. MedTech sales increased 14.6%. International MedTech sales increased by 11.3%, including operational growth of 14.7% partially offset by a negative currency impact of 3.4%. In the fiscal second quarter of 2023, the net impact of acquisitions and divestitures on the MedTech segment operational sales growth was a positive 4.8%, related to the Abiomed acquisition.

Major MedTech Franchise Sales** — Fiscal Second Quarter Ended

(Dollars in Millions)	July 2, 2023	July 3, 2022	Total Change	Operations Change	Currency Change
Surgery	\$ 2,594	\$ 2,450	5.9 %	8.4 %	(2.5)%
Advanced	1,222	1,156	5.8	8.4	(2.6)
General	1,372	1,294	6.0	8.4	(2.4)
Orthopaedics	2,265	2,157	5.0	5.7	(0.7)
Hips	397	388	2.2	3.0	(0.8)
Knees	363	349	3.9	4.5	(0.6)
Trauma	739	696	6.1	6.5	(0.4)
Spine, Sports & Other	766	724	5.8	7.0	(1.2)
Interventional Solutions	1,620	1,049	54.4	56.9	(2.5)
Electrophysiology	1,196	968	23.5	25.9	(2.4)
Abiomed	331	—	*	*	*
Other Interventional Solutions	93	81	15.1	18.8	(3.7)
Vision	1,308	1,241	5.4	6.9	(1.5)
Contact Lenses/Other	939	894	5.1	6.6	(1.5)
Surgical	369	347	6.2	7.6	(1.4)
Total MedTech Sales	\$ 7,788	\$ 6,898	12.9 %	14.7 %	(1.8)%

* Percentage greater than 100% or not meaningful

**Certain prior year amounts have been reclassified to conform to current year presentation

The Surgery franchise achieved operational sales growth of 8.4% as compared to the prior year fiscal second quarter. The operational growth in Advanced Surgery was primarily driven by the following: Biosurgery global procedure growth and strength of the portfolio; Endocutter procedure recovery outside the U.S. and uptake of recently launched products partially offset by volume-based procurement impacts in China, competitive pressures predominantly in the U.S. and supply challenges; and Energy products procedure recovery outside the U.S. and strength of new products partially offset by volume-based procurement impacts in China and supply challenges. The operational growth in General Surgery was primarily driven by procedure recovery outside the U.S. coupled with technology penetration and benefits from the differentiated Wound Closure portfolio.

The Orthopaedics franchise achieved operational sales growth of 5.7% as compared to the prior year fiscal second quarter. The operational growth in hips reflects global procedure growth and continued strength of the portfolio. This was partially offset by supply challenges. The operational growth in knees was primarily driven by global procedure recovery, benefits from launches in the ATTUNE portfolio and pull through related to the VELYS Robotic assisted solution. This was partially offset by supply constraints. The operational growth in Trauma was driven by procedure recovery and the adoption of recently launched products. This was partially offset by volume-based procurement impacts in China. The operational growth in Spine, Sports & Other was primarily driven by procedure growth, positive uptake from new products and growth in VELYS Digital Solutions. This was partially offset by volume-based procurement impacts in China and continued competitive pressures in Spine.

The Interventional Solutions franchise achieved operational sales growth of 56.9% as compared to the prior year fiscal second quarter which includes sales from Abiomed acquired on December 22, 2022. Electrophysiology grew by double digits due to global procedure growth, new product performance and commercial execution. This was partially offset by the impacts of volume-based procurement in China. Abiomed sales reflect the strength of all commercialized regions and continued adoption of Impella 5.5 and Impella RP.

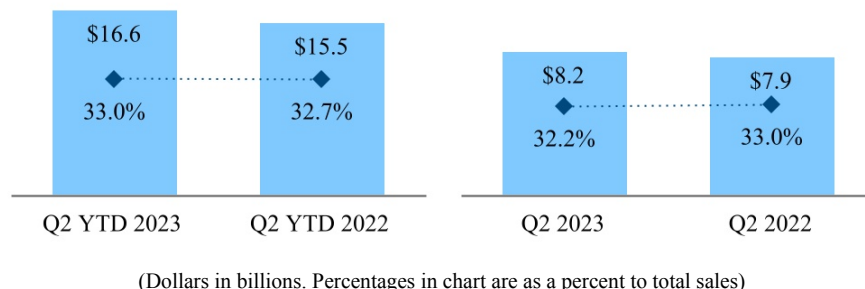
The Vision franchise achieved operational sales growth of 6.9% as compared to the prior year fiscal second quarter. The Contact Lenses/Other operational growth was primarily driven by the continued strong performance in the ACUVUE OASYS 1-Day family (including recent launches) and commercial execution. This was partially offset by impacts from strategic portfolio decisions and supply challenges. The Surgical operational growth was primarily driven by cataract procedure growth and continued strength of recent innovations. This was partially offset by softer Refractive and premium IOL markets challenges.

ANALYSIS OF CONSOLIDATED EARNINGS BEFORE PROVISION FOR TAXES ON INCOME

Consolidated earnings before provision for taxes on income for the fiscal six months of 2023 was \$6.0 billion representing 12.0% of sales as compared to \$11.7 billion in the fiscal six months of 2022, representing 24.7% of sales. The decline was primarily related to the talc settlement proposal in the fiscal first quarter.

Consolidated earnings before provision for taxes on income for the fiscal second quarter of 2023 was \$6.8 billion representing 26.5% of sales as compared to \$5.8 billion in the fiscal second quarter of 2022, representing 24.3% of sales.

Cost of Products Sold



Fiscal six months Q2 2023 versus Fiscal six months Q2 2022

Cost of products sold increased as a percent to sales driven by:

- Commodity inflation and Abiomed amortization in the MedTech business
- Commodity inflation in the Consumer Health business

The intangible asset amortization expense included in cost of products sold for the fiscal six months of 2023 and 2022 was \$2.4 billion and \$2.2 billion, respectively.

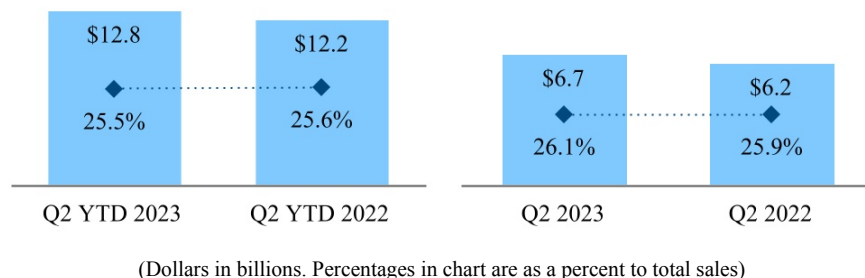
Q2 2023 versus Q2 2022

Cost of products sold decreased as a percent to sales primarily driven by:

- Favorable mix and lower one-time COVID-19 vaccine manufacturing related exit costs in 2023 in the Pharmaceutical business partially offset by
- Commodity inflation and Abiomed amortization in the MedTech business
- Commodity inflation in the Consumer Health business

The intangible asset amortization expense included in cost of products sold for the fiscal second quarters of 2023 and 2022 was \$1.2 billion and \$1.1 billion, respectively.

Selling, Marketing and Administrative Expenses



Fiscal six months Q2 2023 versus Fiscal six months Q2 2022

Selling, Marketing and Administrative Expenses decreased slightly as a percent to sales driven by:

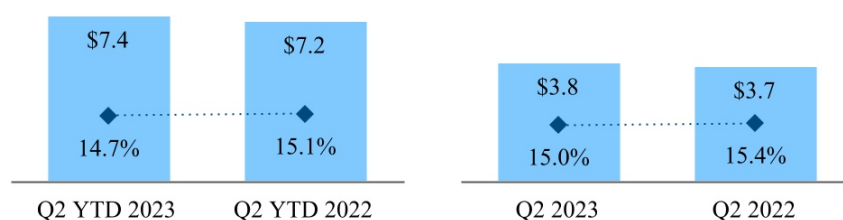
- Leveraging in the Pharmaceutical and MedTech businesses partially offset by
- An increase in Kenvue standalone start-up costs

Q2 2023 versus Q2 2022

Selling, Marketing and Administrative Expenses increased as a percent to sales primarily driven by:

- An increase in Kenvue standalone start-up costs partially offset by
- Leveraging in the Pharmaceutical and MedTech businesses

Research and Development Expense



(Dollars in billions. Percentages in chart are as a percent to total sales)

Fiscal six months Q2 2023 versus Fiscal six months Q2 2022

Research and Development decreased as a percent to sales driven by:

- Cost management initiatives in the MedTech business
- Portfolio prioritization in the Pharmaceutical business partially offset by
- acquired in-process research & development costs in the Pharmaceutical business

Q2 2023 versus Q2 2022

Research and Development decreased as a percent to sales driven by:

- Cost management initiatives in the MedTech business
- Portfolio prioritization in the Pharmaceutical business partially offset by
- acquired in-process research & development costs in the Pharmaceutical business

In-Process Research and Development (IPR&D) Impairments

In the fiscal first quarter of 2023, the Company recorded a charge of approximately \$0.1 billion associated with the IPR&D acquired with Pulsar Vascular in 2016. In the first fiscal six months of 2022, the Company recorded an intangible asset impairment charge of approximately \$0.6 billion related to an in-process research and development asset, bermekimab (JnJ-77474462), an investigational drug for the treatment of Atopic Dermatitis (AD) and Hidradenitis Suppurativa (HS). The Company acquired all rights to bermekimab from XBiotech, Inc. in the fiscal year 2020.

Interest (Income) Expense

Interest income in the first fiscal six months of 2023 was \$604 million as compared to \$86 million in the same period a year ago primarily due to higher rates of interest earned on cash balances. Interest income in the fiscal second quarter of 2023 was \$369 million as compared to \$64 million in the fiscal second quarter of 2022 primarily due to higher rates of interest earned on cash balances. Interest expense in the first fiscal six months of 2023 was \$561 million as compared to \$48 million in the same period a year ago primarily due to a higher debt balance at higher interest rates. Interest expense in the fiscal second quarter of 2023 was \$346 million as compared to \$38 million in the same period a year ago primarily due to a higher debt balance at higher

interest rates. The balance of cash, cash equivalents and current marketable securities was \$28.5 billion at the end of the fiscal second quarter of 2023 as compared to \$32.6 billion at the end of the fiscal second quarter of 2022. The Company's debt position was \$45.6 billion as of July 2, 2023, as compared to \$32.6 billion the same period a year ago.

Other (Income) Expense, Net*

Fiscal six months Q2 2023 versus Fiscal six months Q2 2022

Other (income) expense, net for the fiscal six months of 2023 was unfavorable by \$7.0 billion as compared to the prior year primarily due to the following:

Fiscal Six Months

(Dollars in Billions)(Income)/Expense

	2023	2022	Change
Litigation related ⁽¹⁾	\$ 6.8	0.4	6.4
Consumer Health separation costs	0.5	0.4	0.1
COVID-19 Vaccine related exit costs	0.4	0.0	0.4
Changes in the fair value of securities	0.1	0.5	(0.4)
Employee benefit plan related	(0.7)	(0.6)	(0.1)
Other ⁽²⁾	0.1	(0.5)	0.6
Total Other (Income) Expense, Net	\$ 7.2	0.2	7.0

⁽¹⁾ Primarily related to the talc settlement proposal. The fiscal six months of 2023 includes favorable intellectual property related litigation settlements of approximately \$0.3 billion.

⁽²⁾ Fiscal 2023 includes \$37 million related to the 10.4% non-controlling interest in Kenvue from the time of the initial public offering on May 8, 2023 through the end of the fiscal second quarter

Q2 2023 versus Q2 2022

Other (income) expense, net for the fiscal second quarter of 2023 was favorable by \$0.3 billion as compared to the prior year primarily due to the following:

Fiscal Second Quarter

(Dollars in Billions)(Income)/Expense

	2023	2022	Change
Consumer Health separation costs	\$ 0.2	0.3	(0.1)
COVID-19 Vaccine related exit costs	0.2	0.0	0.2
Employee benefit plan related	(0.4)	(0.3)	(0.1)
Litigation related ⁽¹⁾	(0.1)	0.4	(0.5)
Changes in the fair value of securities	0.0	0.1	(0.1)
Other ⁽²⁾	0.0	(0.2)	0.2
Total Other (Income) Expense, Net	\$ (0.1)	0.3	(0.4)

⁽¹⁾ The fiscal second quarter of 2023 includes favorable intellectual property related litigation settlements of approximately \$0.3 billion.

⁽²⁾ Fiscal 2023 includes \$37 million related to the 10.4% non-controlling interest in Kenvue from the time of the initial public offering on May 8, 2023 through the end of the fiscal second quarter

*Other (income) expense, net is the account where the Company records gains and losses related to the sale and write-down of certain investments in equity securities held by Johnson & Johnson Innovation - JJDC, Inc. (JJDC), changes in the fair value of securities, gains and losses on divestitures, gains and losses on sale of assets, certain transactional currency gains and losses, acquisition-related costs, litigation accruals and settlements, investment (income)/loss related to employee benefit plans, as well as royalty income.

EARNINGS BEFORE PROVISION FOR TAXES BY SEGMENT

Income before tax by segment of business for the fiscal six months were as follows:

(Dollars in Millions)	Income Before Tax		Segment Sales		Percent of Segment Sales	
	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022
Consumer Health	\$ 1,636	\$ 1,470	\$ 7,863	\$ 7,391	20.8 %	19.9 %
Pharmaceutical	9,306	8,344	27,144	26,186	34.3	31.9
MedTech	3,144	2,618	15,269	13,869	20.6	18.9
Segment earnings before tax	14,086	12,432	50,276	47,446	28.0	26.2
Less: Expenses not allocated to segments ⁽¹⁾	7,479	360				
Less: Consumer Health separation costs	582	370				
Worldwide income before tax	<u>\$ 6,025</u>	<u>\$ 11,702</u>	<u>\$ 50,276</u>	<u>\$ 47,446</u>	<u>12.0 %</u>	<u>24.7 %</u>

⁽¹⁾ Amounts not allocated to segments include interest (income) expense and general corporate (income) expense. The fiscal six months of 2023 includes an approximately \$7 billion incremental charge primarily related to the talc settlement proposal.

Consumer Health Segment

The Consumer Health segment income before tax as a percent of sales in the fiscal six months of 2023 was 20.8% versus 19.9% for the same period a year ago. The increase in the income before tax as a percent of sales in the fiscal six months of 2023 as compared to the prior year was primarily driven by the following:

- Pricing actions partially offset by
- Commodity inflation
- Increased Kenvue standalone start-up costs

Pharmaceutical Segment

The Pharmaceutical segment income before tax as a percent of sales in the fiscal six months of 2023 was 34.3% versus 31.9% for the same period a year ago. The increase in the income before tax as a percent of sales for the fiscal six months of 2023 as compared to the prior year was primarily driven by the following:

- An IPR&D charge of \$0.6 billion in 2022 related to bermekimab (JnJ-77474462), an investigational drug for the treatment of AD and Hidradenitis Suppurativa (HS)
- Favorable changes in the fair value of securities in 2023 of \$0.4 billion as compared to 2022
- Leveraging in selling, marketing and administrative expenses
- Portfolio prioritization partially offset by
- One-time COVID-19 Vaccine related exit costs of \$0.6 billion in 2023 as compared to \$0.3 billion in 2022
- Restructuring charges of \$0.3 billion in 2023
- Acquired in-process research & development costs of \$0.2 billion

MedTech Segment

The MedTech segment income before tax as a percent of sales in the fiscal six months of 2023 was 20.6% versus 18.9% for the same period a year ago. The increase in the income before tax as a percent of sales for the fiscal six months of 2023 was primarily driven by the following:

- Net favorable litigation matters of \$0.2 billion in 2023 as compared to expense of \$0.3 billion in 2022
- No Restructuring charges in 2023 versus \$0.2 billion in 2022
- Leveraging in selling and marketing expenses partially offset by
- Higher amortization expense of \$0.3 billion in 2023 related to Abiomed
- An IPR&D charge in 2023 of approximately \$0.1 billion related to the Pulsar Vascular acquisition
- Acquisition costs related to Abiomed of \$0.1 billion
- Commodity inflation in 2023

Income before tax by segment of business for the fiscal second quarters were as follows:

(Dollars in Millions)	Income Before Tax		Segment Sales		Percent of Segment Sales	
	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022	July 2, 2023	July 3, 2022
Consumer Health	\$ 860	\$ 784	\$ 4,011	\$ 3,805	21.4 %	20.6 %
Pharmaceutical	4,862	4,420	13,731	13,317	35.4	33.2
MedTech	1,699	1,141	7,788	6,898	21.8	16.5
Segment earnings before tax	7,421	6,345	25,530	24,020	29.1	26.4
Less: Expenses not allocated to segments ⁽¹⁾	377	237				
Less: Consumer Health separation costs	282	268				
Worldwide income before tax	\$ 6,762	\$ 5,840	\$ 25,530	\$ 24,020	26.5 %	24.3 %

⁽¹⁾Amounts not allocated to segments include interest (income) expense, certain litigation expenses and general corporate (income) expense.

Consumer Health Segment

The Consumer Health segment income before tax as a percent of sales in the fiscal second quarter of 2023 was 21.4% versus 20.6% for the same period a year ago. The increase in the income before tax as a percent of sales in the fiscal second quarter of 2023 as compared to the prior year was primarily driven by the following:

- Pricing actions
- Supply chain efficiencies partially offset by
- Commodity inflation
- Increased Kenvue standalone start-up costs

Pharmaceutical Segment

The Pharmaceutical segment income before tax as a percent of sales in the fiscal second quarter of 2023 was 35.4% versus 33.2% for the same period a year ago. The increase in the income before tax as a percent of sales for the fiscal second quarter as compared to the prior year was primarily driven by the following:

- Favorable litigation related items of \$0.1 billion in 2023
- Favorable changes in the fair value of securities in 2023 of \$0.1 billion as compared to 2022
- One-time COVID-19 Vaccine related exit costs of \$0.2 billion in 2023 as compared to \$0.3 billion in 2022
- Favorable mix
- Leveraging in selling, marketing and administrative expenses
- Portfolio prioritization partially offset by
- Restructuring charges of \$0.1 billion in 2023
- Acquired in-process research & development costs of \$0.2 billion

MedTech Segment

The MedTech segment income before tax as a percent of sales in the fiscal second quarter of 2023 was 21.8% versus 16.5% for the same period a year ago. The increase in the income before tax as a percent of sales for the fiscal second quarter was primarily driven by the following:

- Net favorable litigation matters of \$0.2 billion in 2023 as compared to expense of \$0.3 billion in 2022
- No Restructuring charges in 2023 versus \$0.1 billion in 2022
- Leveraging in selling, marketing and administrative expenses partially offset by
- Higher amortization expense of \$0.1 billion in 2023 related to Abiomed
- Commodity inflation in 2023

Restructuring

In the fiscal first and second quarters of 2023, the Company completed a prioritization of its research and development (R&D) investment within the Pharmaceutical segment to focus on the most promising medicines with the greatest benefit to patients. This resulted in the exit of certain programs within therapeutic areas. The R&D program exits are primarily in infectious diseases and vaccines including the discontinuation of its respiratory syncytial virus (RSV) adult vaccine program, hepatitis and

HIV development. The pre-tax restructuring charge of approximately \$0.1 billion and \$0.3 billion in the fiscal second quarter and fiscal six months of 2023, respectively, includes the termination of partnered and non-partnered program costs and asset impairments. The Company recorded a pre-tax charge of \$0.1 billion in the fiscal second quarter and \$0.2 billion in the first fiscal six months of 2022, related to a restructuring program of its Global Supply Chain. The Global Supply Chain program was announced in the second quarter of 2018 and was completed in the fiscal fourth quarter of 2022.

Provision for Taxes on Income

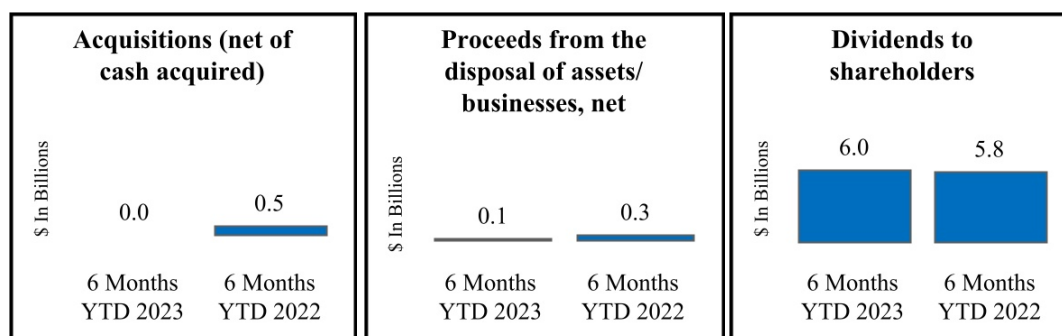
The worldwide effective income tax rate for the first fiscal six months of 2023 was 15.8% in 2023 and 14.9% in 2022.

On December 15, 2022, the European Union (EU) Member States formally adopted the EU's Pillar Two Directive, which generally provides for a minimum effective tax rate of 15%, as established by the Organization for Economic Co-operation and Development (OECD) Pillar Two Framework that was supported by over 130 countries worldwide. The EU's Pillar Two Directive effective dates are January 1, 2024, and January 1, 2025, for different aspects of the directive. On July 17, 2023, the OECD published Administrative Guidance proposing certain safe harbor rules that effectively extend certain effective dates to January 1, 2027. EU Member States will still need to adopt the OECD Administrative Guidance in their local Pillar Two legislation for such safe harbor rules to apply. A significant number of other countries are also considering implementing similar legislation. The Company is continuing to evaluate the potential impact on future periods of the Pillar Two Framework, pending legislative adoption by additional individual countries, including those within the European Union.

On July 21, 2023, the IRS issued Notice 2023-55 which provides guidance to taxpayers in determining whether a foreign tax is eligible for a U.S. foreign tax credit for tax years 2022 and 2023, specifically delaying until 2024 the application of unfavorable foreign tax credit regulations that were originally issued late last year. As a result of this new guidance, the Company has concluded that it is applicable to certain of its tax positions and therefore will record a tax benefit of approximately \$0.5 billion in the fiscal third quarter of 2023.

For discussion related to the 2023 provision for taxes refer to Note 5 to the Consolidated Financial Statements.

LIQUIDITY AND CAPITAL RESOURCES



Cash Flows

Cash and cash equivalents were \$21.2 billion at the end of the fiscal second quarter of 2023 as compared with \$14.1 billion at the end of fiscal year 2022. The primary sources and uses of cash that contributed to the \$7.1 billion increase were:

(Dollars In Billions)	
\$	14.1 Q4 2022 Cash and cash equivalents balance
	7.4 net cash generated from operating activities
	(0.5) net cash used by investing activities
	0.1 net cash generated from financing activities
	0.1 rounding
\$	21.2 Q2 2023 Cash and cash equivalents

In addition, the Company had \$7.3 billion in marketable securities at the end of the fiscal second quarter of 2023 and \$9.4 billion at the end of fiscal year 2022.

Cash flow from operations of \$7.4 billion was the result of:

(Dollars In Billions)	
\$	5.1 Net earnings
	non-cash expenses and other adjustments primarily for depreciation and amortization, stock-based compensation and
	2.5 asset write-downs partially offset by the deferred tax provision and net gain on sale of assets/businesses
	(1.3) an increase in accounts receivable and inventories
	(1.1) a decrease in accounts payable and accrued liabilities
	(1.1) an increase in other current and non-current assets
	3.4 an increase in other current and non-current liabilities
	(0.1) Rounding
\$	7.4 Cash Flow from operations

Cash flow used by investing activities of \$0.5 billion was primarily from:

(Dollars In Billions)	
	(2.0) additions to property, plant and equipment
	0.1 proceeds from the disposal of assets/businesses, net
	2.2 net sales of investments
	(0.8) credit support agreements activity, net and other
\$	(0.5) Net cash used by investing activities

Cash flow from financing activities of \$0.1 billion was primarily from:

(Dollars In Billions)	
\$	(6.0) dividends to shareholders
	(3.9) repurchase of common stock
	(1.9) net repayment of short and long term debt and other
	7.7 proceeds from Kenvue long term debt, net of issuance cost
	4.2 proceeds from Kenvue initial public offering
\$	0.1 Net cash from financing activities

The Company has access to substantial sources of funds at numerous banks worldwide. In September 2022, the Company secured a new 364-day Credit Facility of \$10 billion, which expires on September 7, 2023. In November 2022, the Company secured an additional 364-day revolving Credit Facility of \$10 billion, which has an expiration of November 21, 2023. Interest charged on borrowings under the credit line agreement is based on either Secured Overnight Financing Rate (SOFR) Reference Rate or other applicable market rate as allowed plus applicable margins. Commitment fees under the agreement are not material.

In March 2023, Kenvue, priced an offering of senior unsecured notes (the Notes) in an aggregate principal amount of \$7.75 billion (See Note 4 to the Consolidated Financial Statements for additional details). The Notes were initially fully and unconditionally guaranteed (the Guarantees) on a senior unsecured basis by the Company. The Guarantees terminated on April 5, 2023, upon completion of the Consumer Health Business transfer.

Further, in March 2023, Kenvue entered into a credit agreement providing for a five-year senior unsecured revolving credit facility (the Revolving Credit Facility) in an aggregate principal amount of \$4.0 billion to be made available in U.S. dollars and Euros. The Revolving Credit Facility contains representations and warranties, covenants and events of default that are customary for this type of financing, including covenants restricting the incurrence of liens and the entry into certain merger transactions. In addition, Kenvue entered into a commercial paper program (the Commercial Paper Program) of up to \$4.0

billion in aggregate principal amount of commercial paper under the Commercial Paper Program. The Commercial Paper Program contains representations and warranties, covenants and default that are customary for this type of financing.

On May 8, 2023, Kenvue completed an initial public offering (the IPO) of 198,734,444 shares of its common stock, par value \$0.01 per share (the “Kenvue Common Stock”), including the underwriters’ full exercise of their option to purchase 25,921,884 shares to cover over-allotments, at an initial public offering price of \$22.00 per share for net proceeds of \$4.2 billion. Kenvue shares began trading on the New York Stock Exchange under the symbol “KVUE”. Additionally, on May 8, 2023, as partial consideration for the transfer of the Consumer Health business, Kenvue paid \$13.2 billion to Johnson & Johnson from the net proceeds of the initial public offering and debt financing transactions (approximately \$7.7 billion of unsecured notes and \$1.2 billion of commercial paper) in connection with the separation. The excess of the net proceeds from the IPO over the net book value of the Johnson & Johnson divested interest was 2.5 billion and was recorded in additional paid-in capital.

As of the closing of the IPO, Johnson & Johnson owns 1,716,160,000 shares of Kenvue Common Stock, or approximately 89.6% of the total outstanding shares of Kenvue Common Stock, and continues to consolidate the financial results of Kenvue. As of July 2, 2023, the non-controlling interest of \$1.3 billion associated with Kenvue is reflected in equity attributable to non-controlling interests in the consolidated balance sheet. Other (income) expense, net on the consolidated statement of earnings for the fiscal second quarter and fiscal six months ended July 2, 2023, includes \$37 million related to the 10.4% non-controlling interest in Kenvue from the time of the initial public offering on May 8, 2023 through the end of the fiscal second quarter.

Subsequent to the fiscal second quarter on July 24, 2023, the Company announced its intention to split-off at least 80.1% of the shares of Kenvue through an exchange offer. Through the planned exchange offer, Johnson & Johnson shareholders can exchange all, some or none of their shares of Johnson & Johnson Common Stock for shares of Kenvue Common Stock, subject to the terms of the offer. (See Note 12 to the Consolidated Financial Statements for additional details). The shares of Johnson & Johnson Common Stock acquired by the Company in the exchange offer will be recorded as an acquisition of treasury stock at a cost equal to the market value of the shares of Johnson & Johnson Common Stock accepted in the exchange offer at its expiration. Any difference between the net book value of Kenvue attributable to Johnson & Johnson and the market value of the shares of Johnson & Johnson Common Stock acquired at that date will be recognized by the Company as a gain on disposal of discontinued operations net of any direct and incremental expenses of the exchange offer on the disposal of its Kenvue Common Stock. Upon completion of the exchange offer, and assuming the Company no longer has a controlling financial interest in Kenvue, Kenvue’s historical results will be shown in the Company’s financial statements as discontinued operations, and, in subsequent periods, the Company’s financial statements will no longer reflect the assets, liabilities, results of operations or cash flows attributable to Kenvue.

As of July 2, 2023, the Company’s cash, cash equivalents and marketable securities was approximately \$28.5 billion and had approximately \$45.6 billion of notes payable and long-term debt for a net debt position of \$17.1 billion as compared to the prior year net neutral position. The Company anticipates that operating cash flows, the ability to raise funds from external sources, borrowing capacity from existing committed credit facilities and access to the commercial paper markets will continue to provide sufficient resources to fund operating needs, including the Company’s remaining balance to be paid on the agreement to settle opioid litigation for approximately \$2.2 billion and the establishment of the approximately \$9 billion reserve (present value) for the talc settlement proposal (See Note 11 to the Consolidated Financial Statements for additional details). In addition, the Company monitors the global capital markets on an ongoing basis and from time to time may raise capital when market conditions are favorable.

In the fiscal second quarter of 2023, the Company paid approximately \$4.3 billion to the U.S. Treasury including \$1.5 billion related to the current installment due on foreign undistributed earnings as part of the TCJA charge (see Note 1 to the Consolidated Financial Statements in the Company’s Annual Report on Form 10-K for the fiscal year ended January 1, 2023), \$1.4 billion in pre-payments to resolve certain items previously reserved for and under examination in its 2013 through 2016 U.S. IRS audit, and \$1.4 billion primarily related to the normal estimated payments for the first six months of fiscal 2023.

On September 14, 2022, the Company announced that its Board of Directors approved a share repurchase program, authorizing the Company to purchase up to \$5.0 billion of the Company’s Common Stock. As of April 2, 2023, \$5.0 billion has been repurchased and the repurchase program was completed.

Dividends

On April 18, 2023, the Board of Directors declared a regular cash dividend of \$1.19 per share, payable on June 6, 2023, to shareholders of record as of May 23, 2023.

On July 20, 2023, the Board of Directors declared a regular cash dividend of \$1.19 per share, payable on September 7, 2023, to shareholders of record as of August 28, 2023. The Company expects to continue the practice of paying regular quarterly cash dividends.

OTHER INFORMATION

New Accounting Pronouncements

Refer to Note 1 to the Consolidated Financial Statements for new accounting pronouncements.

Economic and Market Factors

In July 2023, Janssen Pharmaceuticals, Inc. (Janssen) filed litigation against the U.S. Department of Health and Human Services as well as the Centers for Medicare and Medicaid Services challenging the constitutionality of the Inflation Reduction Act's (IRA) Medicare Drug Price Negotiation Program. The litigation requests a declaration that the IRA violates Janssen's rights under the First Amendment and the Fifth Amendment to the Constitution and therefore that Janssen is not subject to the IRA's mandatory pricing scheme.

Russia-Ukraine War

Although the long-term implications of Russia's invasion of Ukraine are difficult to predict at this time, the financial impact of the conflict in the fiscal second quarter and fiscal six months of 2023, including accounts receivable or inventory reserves, was not material. As of both the fiscal second quarter ending July 2, 2023, and the 2022 fiscal year ending January 1, 2023, the business of the Company's Ukraine subsidiaries represented less than 1% of the Company's consolidated assets and revenues. As of both the fiscal second quarter ending July 2, 2023, and the 2022 fiscal year ending January 1, 2023, the business of the Company's Russian subsidiaries represented less than 1% of the Company's consolidated assets and represented 1% of revenues.

In early March of 2022, the Company took steps to suspend all advertising, enrollment in clinical trials, and any additional investment in Russia. Additionally, at the end of March 2022, the Company made the decision to suspend supply of personal care products in Russia. The Company continues to supply its other products as patients rely on many of the products for healthcare purposes.

The Company operates in certain countries where the economic conditions continue to present significant challenges. The Company continues to monitor these situations and take appropriate actions. Inflation rates and currency exchange rates continue to have an effect on worldwide economies and, consequently, on the way the Company operates. The Company has accounted for operations in Venezuela, Argentina and Turkey (beginning in the fiscal second quarter of 2022) as highly inflationary, as the prior three-year cumulative inflation rate surpassed 100%. In the face of increasing costs, the Company strives to maintain its profit margins through cost reduction programs, productivity improvements and periodic price increases.

Governments around the world consider various proposals to make changes to tax laws, which may include increasing or decreasing existing statutory tax rates. In connection with various government initiatives, companies are required to disclose more information to tax authorities on operations around the world, which may lead to greater audit scrutiny of profits earned in other countries. A change in statutory tax rate in any country would result in the revaluation of the Company's deferred tax assets and liabilities related to that particular jurisdiction in the period in which the new tax law is enacted. This change would result in an expense or benefit recorded to the Company's Consolidated Statement of Earnings. The Company closely monitors these proposals as they arise in the countries where it operates. Changes to the statutory tax rate may occur at any time, and any related expense or benefit recorded may be material to the fiscal quarter and year in which the law change is enacted.

The Company faces various worldwide health care changes that may continue to result in pricing pressures that include health care cost containment and government legislation relating to sales, promotions and reimbursement of health care products.

Changes in the behavior and spending patterns of purchasers of healthcare products and services, including delaying medical procedures, rationing prescription medications, reducing the frequency of physician visits and foregoing healthcare insurance coverage, as a result of the current global economic downturn, may continue to impact the Company's businesses.

The Company faces regular intellectual property challenges from third parties, including generic and biosimilar manufacturers, seeking to manufacture and market generic and biosimilar versions of key pharmaceutical products prior to the expiration of the applicable patents. These challengers file Abbreviated New Drug Applications or abbreviated Biologics License Applications with the FDA or otherwise challenged the coverage and/or validity of the Company's patents. In the event the Company is not successful in defending the patent claims challenged in the resulting lawsuits, generic or biosimilar versions of the products at issue may be introduced to the market, resulting in the potential for substantial market share and revenue losses for those products, and which may result in a non-cash impairment charge in any associated intangible asset. There is also risk that one or more competitors could launch a generic or biosimilar version of the product at issue following regulatory approval even though one or more valid patents are in place.

Item 3 — QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK

There has been no material change in the Company’s assessment of its sensitivity to market risk since its presentation set forth in Item 7A, “Quantitative and Qualitative Disclosures About Market Risk,” in its Annual Report on Form 10-K for the fiscal year ended January 1, 2023.

Item 4 — CONTROLS AND PROCEDURES

Disclosure controls and procedures. At the end of the period covered by this report, the Company evaluated the effectiveness of the design and operation of its disclosure controls and procedures. The Company’s disclosure controls and procedures are designed to ensure that information required to be disclosed by the Company in the reports that it files or submits under the Securities Exchange Act is recorded, processed, summarized and reported, within the time periods specified in the SEC’s rules and forms. Disclosure controls and procedures include, without limitation, controls and procedures designed to ensure that information required to be disclosed by the Company in the reports that it files or submits under the Securities Exchange Act is accumulated and communicated to the Company’s management, including its principal executive and principal financial officers, or persons performing similar functions, as appropriate, to allow timely decisions regarding required disclosure. Joaquin Duato, Chief Executive Officer; Chairman, Executive Committee and Joseph J. Wolk, Executive Vice President, Chief Financial Officer, reviewed and participated in this evaluation. Based on this evaluation, Messrs. Duato and Wolk concluded that, as of the end of the period covered by this report, the Company’s disclosure controls and procedures were effective.

Internal control. During the period covered by this report, there were no changes in the Company’s internal control over financial reporting that have materially affected, or are reasonably likely to materially affect, the Company’s internal control over financial reporting. The Company continues to monitor and assess the effectiveness of the design and operation of its disclosure controls and procedures.

Part II — OTHER INFORMATION

Item 1 — LEGAL PROCEEDINGS

The information called for by this item is incorporated herein by reference to Note 11 included in Part I, Item 1, Financial Statements (unaudited) — Notes to Consolidated Financial Statements.

Item 2 — UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS

(c) Purchases of Equity Securities by the Issuer and Affiliated Purchasers.

On September 14, 2022, the Company announced that its Board of Directors approved a share repurchase program, authorizing the Company to purchase up to \$5.0 billion of the Company’s Common Stock. The repurchase program was completed during the fiscal first quarter of 2023.

The following table provides information with respect to Common Stock purchases by the Company during the fiscal second quarter of 2023. Common stock purchases on the open market are made as part of a systematic plan to meet the needs of the Company’s compensation programs. The repurchases below also include the stock-for-stock option exercises that settled in the fiscal second quarter.

Fiscal Month Period	Total Number of Shares Purchased ⁽¹⁾	Avg. Price Per Share	Total Number of Shares Purchased as Part of Publicly Announced Plans or Programs ⁽²⁾	Maximum Number of Shares that May Yet Be Purchased Under the Plans or Programs
April 3, 2023 through April 30, 2023	1,011,500	162.35	—	—
May 1, 2023 through May 28, 2023	383,500	161.73	—	—
May 29, 2023 through July 2, 2023	962,825	160.35	—	—
Total	2,357,825	161.44	—	—

- (1) During the fiscal second quarter of 2023, the Company repurchased an aggregate of 2,357,825 shares of Johnson & Johnson Common Stock in open-market transactions, all of which were purchased pursuant as part of a systematic plan to meet the needs of the Company's compensation programs.
- (2) As of April 2, 2023, an aggregate of 30,546,218 shares were purchased for a total of \$5.0 billion since the inception of the repurchase program announced on September 14, 2022.

Item 6 — EXHIBITS

[Exhibit 31.1](#) Certification of Chief Executive Officer under Rule 13a-14(a) of the Securities Exchange Act pursuant to Section 302 of the Sarbanes-Oxley Act of 2002 — Filed with this document.

[Exhibit 31.2](#) Certification of Chief Financial Officer under Rule 13a-14(a) of the Securities Exchange Act pursuant to Section 302 of the Sarbanes-Oxley Act of 2002 — Filed with this document.

[Exhibit 32.1](#) Certification of Chief Executive Officer pursuant to Section 906 of the Sarbanes-Oxley Act of 2002 — Furnished with this document.

[Exhibit 32.2](#) Certification of Chief Financial Officer pursuant to Section 906 of the Sarbanes-Oxley Act of 2002 — Furnished with this document.

Exhibit 101:

EX-101.INS	Instance Document - the instance document does not appear in the Interactive Data File because its XBRL tags are embedded within the Inline XBRL document
EX-101.SCH	Inline XBRL Taxonomy Extension Schema
EX-101.CAL	Inline XBRL Taxonomy Extension Calculation Linkbase
EX-101.LAB	Inline XBRL Taxonomy Extension Label Linkbase
EX-101.PRE	Inline XBRL Taxonomy Extension Presentation Linkbase
EX-101.DEF	Inline XBRL Taxonomy Extension Definition Document

Exhibit 104: Cover Page Interactive Data File—the cover page interactive data file does not appear in the Interactive Data File because its XBRL tags are embedded within the Inline XBRL document.

SIGNATURES

Pursuant to the requirements 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.

Date: July 31, 2023

JOHNSON & JOHNSON
(Registrant)

By /s/ J. J. WOLK

J. J. WOLK

Executive Vice President, Chief Financial Officer (Principal Financial Officer)

Date: July 31, 2023

By /s/ R. J. DECKER Jr.

R. J. DECKER Jr.

Controller (Principal Accounting Officer)

CERTIFICATION OF CHIEF EXECUTIVE OFFICER
PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT

I, Joaquin Duato, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q for the quarterly period ended July 2, 2023 (the "report") of Johnson & Johnson (the "Company");

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 Company as of, and for, the periods presented in this report;

4. The Company's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rule 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rule 13a-15(f) and 15d-15(f)) for the Company and have:

a) 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 Company, 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 Company'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 Company's internal control over financial reporting that occurred during the Company's most recent fiscal quarter (the Company's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the Company's internal control over financial reporting; and

5. The Company's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the Company's auditors and the audit committee of the Company'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 Company'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 Company's internal control over financial reporting.

/s/ Joaquin Duato

Joaquin Duato
Chief Executive Officer

Date: July 31, 2023

CERTIFICATION OF CHIEF FINANCIAL OFFICER
PURSUANT TO SECTION 302 OF THE SARBANES-OXLEY ACT

I, Joseph J. Wolk, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q for the quarterly period ended July 2, 2023 (the "report") of Johnson & Johnson (the "Company");

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 Company as of, and for, the periods presented in this report;

4. The Company's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rule 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rule 13a-15(f) and 15d-15(f)) for the Company and have:

a) 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 Company, 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 Company'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 Company's internal control over financial reporting that occurred during the Company's most recent fiscal quarter (the Company's fourth fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the Company's internal control over financial reporting; and

5. The Company's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the Company's auditors and the audit committee of the Company'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 Company'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 Company's internal control over financial reporting.

/s/ Joseph J. Wolk

Joseph J. Wolk
Chief Financial Officer

Date: July 31, 2023

CERTIFICATION OF CHIEF EXECUTIVE OFFICER

PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT

The undersigned, Joaquin Duato, the Chief Executive Officer of Johnson & Johnson, a New Jersey corporation (the “Company”), pursuant to 18 U.S.C. 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, hereby certifies that, to the best of my knowledge:

- (1) the Company's Quarterly Report on Form 10-Q for the quarterly period ended July 2, 2023 (the “Report”) fully complies with the requirements of Section 13(a) 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.

/s/ Joaquin Duato

Joaquin Duato

Chief Executive Officer

Dated: July 31, 2023

This certification is being furnished to the SEC with this Report pursuant to Section 906 of the Sarbanes-Oxley Act of 2002 and shall not, except to the extent required by such Act, be deemed filed by the Company for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to liability of that section.

CERTIFICATION OF CHIEF FINANCIAL OFFICER

PURSUANT TO SECTION 906 OF THE SARBANES-OXLEY ACT

The undersigned, Joseph J. Wolk, the Chief Financial Officer of Johnson & Johnson, a New Jersey corporation (the “Company”), pursuant to 18 U.S.C. 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, hereby certifies that, to the best of my knowledge:

- (1) the Company's Quarterly Report on Form 10-Q for the quarterly period ended July 2, 2023 (the “Report”) fully complies with the requirements of Section 13(a) 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.

/s/ Joseph J. Wolk

Joseph J. Wolk
Chief Financial Officer

Dated: July 31, 2023

This certification is being furnished to the SEC with this Report pursuant to Section 906 of the Sarbanes-Oxley Act of 2002 and shall not, except to the extent required by such Act, be deemed filed by the Company for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to liability of that section.