

# 3RD QUARTER 2018 RESULTS

## Q3 2018 SALES

**\$20.3B**

WORLDWIDE INCREASED ▲  
**3.6%**

Excluding acquisitions/  
divestitures on an operational  
basis worldwide sales

INCREASED ▲  
**6.1%\***

DILUTED EARNINGS  
PER SHARE

**\$1.44**

ADJUSTED DILUTED EARNINGS PER SHARE\*

**\$2.05**

INCREASE ▲  
**7.9%**



“We are pleased with our strong third-quarter performance, which reflects continued above-market growth in our Pharmaceutical business, accelerating sales momentum in our Consumer business and consistent progress in our Medical Devices business,” said Alex Gorsky, Chairman and Chief Executive Officer. “I’m confident that with our collaborative and inspired J&J colleagues around the world, unique broad-based business model and strategic investments in innovation, we are well positioned for success today and into the future.”

## WORLDWIDE CONSUMER SALES

**\$3.4B**

Consumer worldwide sales increased: **1.8%**

Primary contributors to growth:



## WORLDWIDE PHARMACEUTICAL SALES

**\$10.3B**

Pharmaceutical worldwide sales increased: **6.7%**

Primary contributors to growth:



## WORLDWIDE MEDICAL DEVICES SALES

**\$6.6B**

Medical Devices worldwide sales decreased: **(0.2%)**

Primary contributors to growth:



ELECTROPHYSIOLOGY



ENDOCUTTERS



BIOSURGERY



ACUVUE®  
CONTACT LENSES



ENERGY



SURGICAL  
VISION



WOUND  
CLOSURE

Note: values may have been rounded

For full financial data and non-GAAP reconciliations, please refer to Johnson & Johnson’s earnings release issued on October 16, 2018, available at <http://www.investor.jnj.com/sales-earnings.cfm>.

\*Non-GAAP financial measure; non-GAAP financial measures should not be considered replacements for, and should be read together with, the most comparable GAAP financial measures.

Caution Concerning Forward-Looking Statements: This document contains “forward-looking statements” as defined in the Private Securities Litigation Reform Act of 1995 regarding future operating and financial performance. You are cautioned not to rely on these forward-looking statements, which are based on current expectations of future events. For important information about the risks and uncertainties that could cause actual results to vary materially from the assumptions, expectations, and projections expressed in any forward-looking statements, review the “Note to Investors Concerning Forward-Looking Statements” included in the Johnson & Johnson earnings release issued on October 16, 2018, as well as the most recently filed Johnson & Johnson Reports on Forms 10-K and 10-Q. Johnson & Johnson does not undertake to update any forward-looking statement as a result of new information or future events or developments.

# 3<sup>rd</sup> Quarter 2018 Earnings Call

October 16, 2018

*Johnson & Johnson*





## **Christopher DeOrefice**

Vice President,  
Investor Relations





# Agenda

Sales Performance and Highlights

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Enterprise Update

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Financial Results Review and Guidance

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Q&A



# Cautionary Note on Forward-looking Statements

This presentation contains “forward-looking statements” as defined in the Private Securities Litigation Reform Act of 1995 regarding, among other things: future operating and financial performance, product development, market position and business strategy. The reader is cautioned not to rely on these forward-looking statements. These statements are based on current expectations of future events. If underlying assumptions prove inaccurate or known or unknown risks or uncertainties materialize, actual results could vary materially from the expectations and projections of Johnson & Johnson. Risks and uncertainties include, but are not limited to: economic factors, such as interest rate and currency exchange rate fluctuations; competition, including technological advances, new products and patents attained by competitors; challenges inherent in new product research and development, including unexpected clinical trial results, additional analysis of existing clinical data, uncertainty of clinical success and obtaining regulatory approvals; uncertainty of commercial success for new and existing products; the impact of business combinations and divestitures; challenges to patents; the impact of patent expirations; the ability of the company to successfully execute strategic plans, including restructuring plans; manufacturing difficulties or delays, internally or within the supply chain; product efficacy or safety concerns resulting in product recalls or regulatory action; significant adverse litigation or government action, including related to product liability claims; changes to applicable laws and regulations, including tax laws, global health care reforms and import/export and trade laws; trends toward health care cost containment; changes in behavior and spending patterns of purchasers of health care products and services; financial instability of international economies and legal systems and sovereign risk; increased scrutiny of the health care industry by government agencies. A further list and descriptions of these risks, uncertainties and other factors can be found in Johnson & Johnson's Annual Report on Form 10-K for the fiscal year ended December 31, 2017, including in the sections captioned “Cautionary Note Regarding Forward-Looking Statements” and “Item 1A. Risk Factors,” in the company’s most recently filed Quarterly Report on Form 10-Q and in the company’s subsequent filings with the Securities and Exchange Commission. Copies of these filings are available online at [www.sec.gov](http://www.sec.gov), [www.jnj.com](http://www.jnj.com) or on request from Johnson & Johnson. Any forward-looking statement made in this presentation speaks only as of the date of this presentation. Johnson & Johnson does not undertake to update any forward-looking statement as a result of new information or future events or developments.

# Cautionary Note on Non-GAAP Financial Measures

This presentation refers to certain non-GAAP financial measures. These non-GAAP financial measures should not be considered replacements for, and should be read together with, the most comparable GAAP financial measures.

A reconciliation of these non-GAAP financial measures to the most directly comparable GAAP financial measures can be found in the accompanying financial schedules of the earnings release and the Investor Relations section of the Company’s website at [www.investor.jnj.com](http://www.investor.jnj.com).



# Strategic Partnerships, Collaborations & Licensing Arrangements

During the course of this morning's presentation, we will discuss a number of products and compounds developed in collaboration with strategic partners or licensed from other companies. Following is an acknowledgement of those relationships:

## Consumer

RHINOCORT licensed from AstraZeneca; DR. CI:LABO brand skincare products in collaboration with Ci:z Holdings Co., Ltd.

## Immunology

REMICADE and SIMPONI/ SIMPONI ARIA marketing partners are Schering-Plough (Ireland) Company, a subsidiary of Merck & Co., Inc. and Mitsubishi Tanabe Pharma Corporation, and TREMFYA discovered using MorphoSys AG antibody technology

## Neuroscience

INVEGA SUSTENNA/ XEPLION/ INVEGA TRINZA/ TREVICTA includes technology licensed from Alkermes Pharma Ireland Limited

## Infectious Diseases & Virology

PREZCOBIX/ REZOLSTA fixed-dose combination, SYMTUZA and ODEFSEY developed in collaboration with Gilead Sciences, Inc., and JULUCA developed in collaboration with ViiV Healthcare UK

## Cardiovascular/ Metabolism/Other

INVOKANA/ INVOKAMET/ VOKANAMET/ INVOKAMET XR fixed-dose combination licensed from Mitsubishi Tanabe Pharma Corporation, XARELTO co-developed with Bayer HealthCare AG, and PROCRI/EPREX licensed from Amgen Inc.

## Oncology

IMBRUVICA developed in collaboration and co-marketed in the U.S. with Pharmacyclics, LLC, an AbbVie company, ZYTIGA licensed from BTG International Ltd., VELCADE developed in collaboration with Millennium: The Takeda Oncology Company, DARZALEX licensed from Genmab A/S, and erdafitinib discovered in collaboration with Astex Pharmaceuticals, Inc.

## Pulmonary Hypertension

UPTRAVI license and supply agreement with Nippon Shinyaku (co-promotion in Japan), OPSUMIT co-promotion agreement with Nippon Shinyaku in Japan





# 3<sup>rd</sup> Quarter 2018 Sales

\$ U.S. Billions

Total Company	3Q 2018	3Q 2017	% Change	
			Reported	Operational <sup>1</sup>
U.S	\$10.7	\$10.3	3.6%	3.6%
Europe	4.4	4.3	2.5	5.1
Western Hemisphere (ex U.S.)	1.6	1.6	(1.2)	11.2
Asia-Pacific, Africa	3.7	3.5	6.7	8.6
International	9.7	9.4	3.5	7.5
Worldwide (WW)	\$20.3	\$19.7	3.6%	5.5%

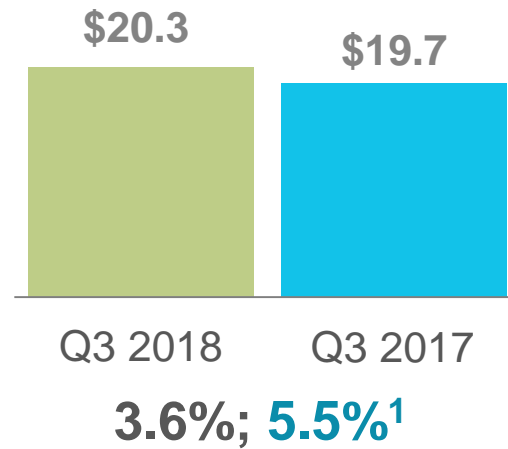


<sup>1</sup> Excludes impact of translational currency  
Note: values may have been rounded

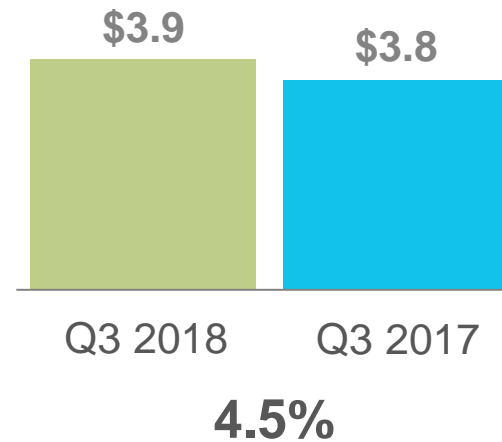
# 3<sup>rd</sup> Quarter 2018 Financial Highlights

\$ U.S. Billions, except EPS  
Reported %; **Operational %**

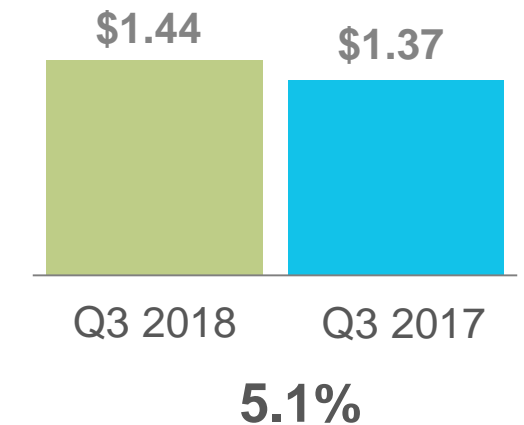
## Sales



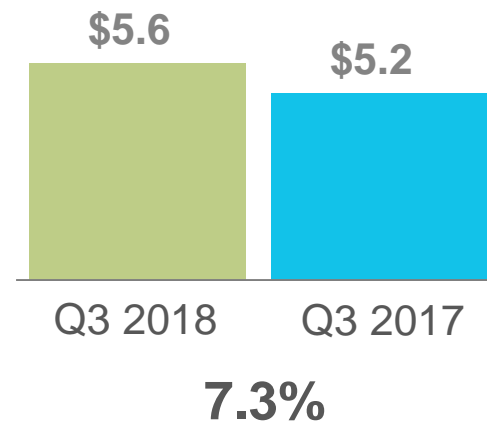
## GAAP Earnings



## GAAP EPS



## Adjusted Earnings<sup>2</sup>



## Adjusted EPS<sup>2</sup>



<sup>1</sup> Excludes impact of translational currency

<sup>2</sup> Non-GAAP measure; excludes intangible amortization expense and special items; see reconciliation

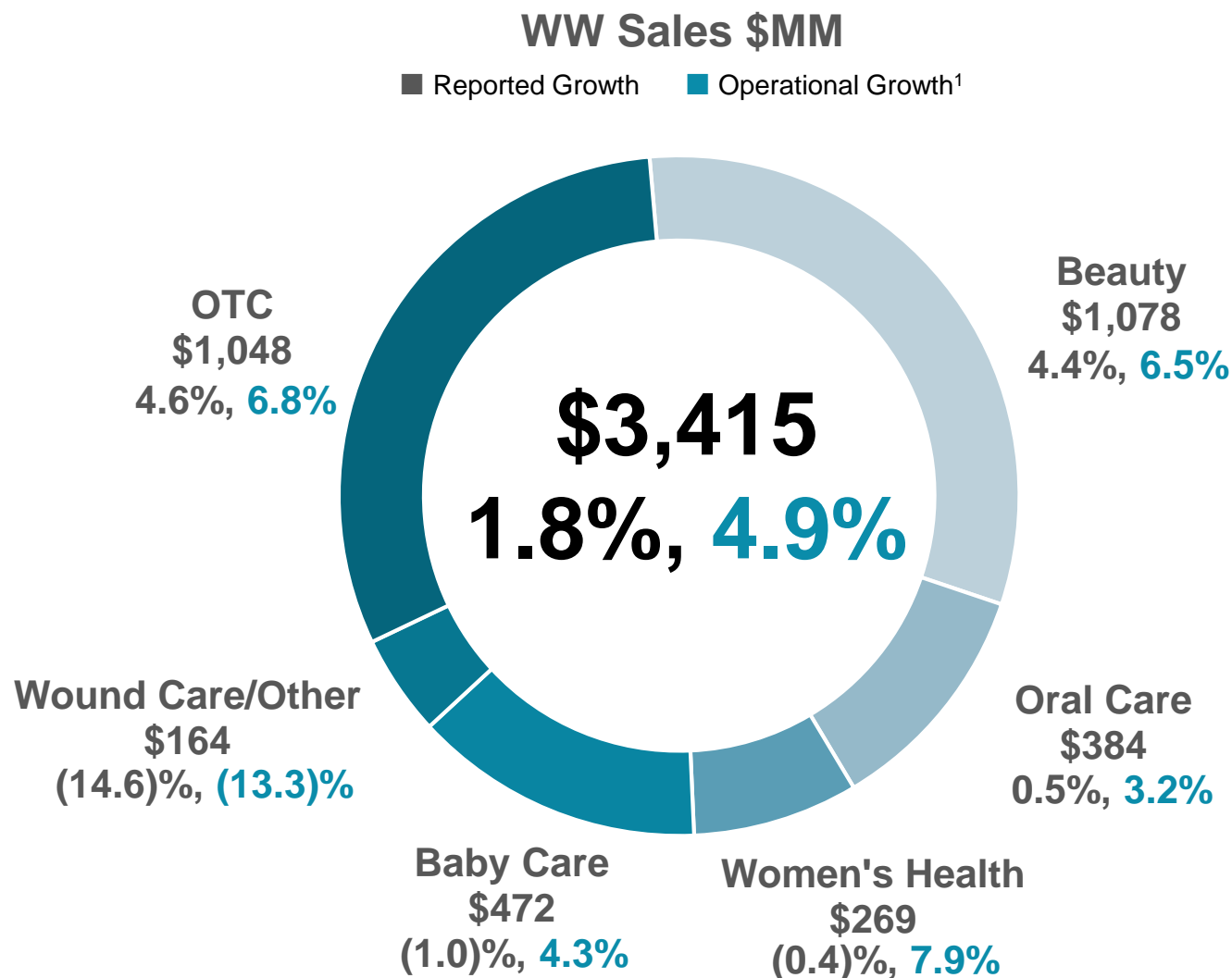


# Consumer Highlights – 3<sup>rd</sup> Quarter 2018

Improved operational performance across all franchises and all major regions driving above-market growth

Reported: WW 1.8%, U.S. 6.6%, Int'l (1.3)%

Operational<sup>1</sup>: WW 4.9%, U.S. 6.6%, Int'l 3.7%



## Key Drivers of Operational Performance<sup>1</sup>

<b>Baby Care</b>	<ul style="list-style-type: none"> <li>JOHNSON's U.S. relaunch pipeline replenishment; excluding pipeline fill WW results are flat</li> </ul>
<b>Beauty</b>	<ul style="list-style-type: none"> <li>Growth driven by ASPAC premium expansion (DR. CI:LABO) and DABAO strong consumption, NEUTROGENA market and sales growth through channel expansion and HYDROBOOST performance coupled with OGX, MAUI MOISTURE share gains and NEOSTRATA growth primarily due to market/product expansion</li> </ul>
<b>Oral Care</b>	<ul style="list-style-type: none"> <li>LISTERINE new product launches such as Ready Tabs in the US, reformulated Total Care, LISTERINE Kids and Nightly Reset in ASPAC</li> </ul>
<b>OTC</b>	<ul style="list-style-type: none"> <li>Share and consumption growth primarily in ZYRTEC, TYLENOL, IMODIUM and Children's MOTRIN as well as NICORETTE Quickmist performance and the ZARBEE's acquisition</li> </ul>
<b>Women's Health</b>	<ul style="list-style-type: none"> <li>Growth primarily driven by liners in LATAM</li> </ul>
<b>Wound Care/Other</b>	<ul style="list-style-type: none"> <li>COMPEED OUS divestiture</li> </ul>

Sales excl. acquisition and divestiture<sup>2</sup>: WW 6.1%, U.S. 6.4%, OUS 5.9%



<sup>1</sup> Excludes impact of translational currency

<sup>2</sup> Non-GAAP measure; see reconciliation



Neutrogena



Aveeno  
ACTIVE NATURALS



Carefree

LISTERINE

BAND-AID  
BRAND ADHESIVE BANDAGES



TYLENOL

Motrin<sup>IB</sup>



# Pharmaceutical Highlights – 3<sup>rd</sup> Quarter 2018

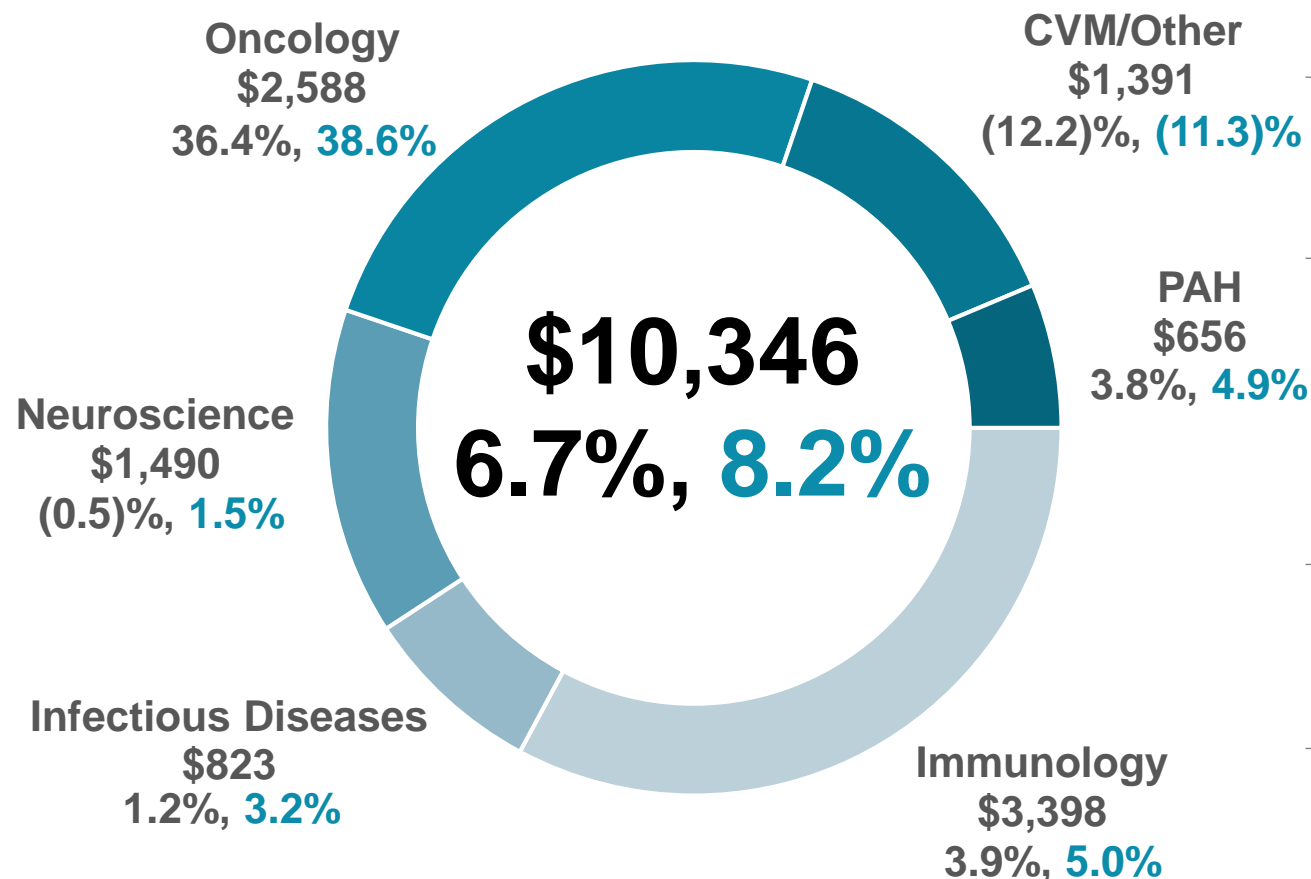
Above market performance driven by double-digit growth in nine key products

Reported: WW 6.7%, U.S. 4.8%, Int'l 9.5%

Operational<sup>1</sup>: WW 8.2%, U.S. 4.8%, Int'l 13.2%

## WW Sales \$MM

■ Reported Growth ■ Operational Growth<sup>1</sup>



## Key Drivers of Operational Performance<sup>1</sup>

<b>Immunology</b>	<ul style="list-style-type: none"> <li>Growth driven by strong uptake of STELARA in Crohn's Disease, expanded indications of SIMPONI/SIMPONI ARIA, strong launch uptake of TREMFYA and U.S. immunology market growth</li> <li>Lower sales of REMICADE due to increased discounts/rebates and biosimilar competition</li> </ul>
<b>Infectious Diseases</b>	<ul style="list-style-type: none"> <li>Strong sales of PREZCOBIX/REZOLSTA, continued success of ODEFSEY and launch of SYMTUZA and JULUCA, offset by lower sales of PREZISTA</li> </ul>
<b>Neuroscience</b>	<ul style="list-style-type: none"> <li>Paliperidone long-acting injectables growth due to strength of INVEGA TRINZA/TREVICTA and INVEGA SUSTENNA/XEPLION, partially offset by cannibalization of RISPERDAL CONSTA and generic competition for CONCERTA</li> </ul>
<b>Oncology</b>	<ul style="list-style-type: none"> <li>DARZALEX continued strong uptake and share gain in the U.S. and EU, especially 1-prior line setting; commercially available in 31 countries in EMEA. Strong growth in Japan</li> <li>Strong sales of IMBRUVICA due to increased patient uptake globally; higher market share and growth across multiple indications; U.S. new and total patient share leader for 1st and 2nd line CLL and 2nd line MCL and total patient share leader in WM</li> <li>Strong sales and share growth of ZYTIGA driven by LATITUDE data and market growth</li> <li>Addition of newly launched ERLEADA sales in non-metastatic CRPC</li> </ul>
<b>Cardiovascular/ Metabolism/ Other (CVM/Other)</b>	<ul style="list-style-type: none"> <li>XARELTO lower sales driven by higher discounts and rebates, partially offset by increase in market share</li> <li>INVOKANA/INVOKAMET lower sales due to increased contracting discounts and higher managed care rebates and share loss due to competitive pressures</li> </ul>
<b>Pulmonary Hypertension (PAH)</b>	<ul style="list-style-type: none"> <li>Growth driven by strong sales of OPSUMIT and UPTRAVI due to continued market growth and share gains</li> <li>Lower sales of TRACLEER due to market share loss to OPSUMIT and generics</li> </ul>

Sales excl. acquisition and divestiture<sup>2</sup>: WW 8.2%, U.S. 4.8%, OUS 13.2%



<sup>1</sup> Excludes impact of translational currency

<sup>2</sup> Non-GAAP measure; see reconciliation



# Medical Devices Highlights – 3<sup>rd</sup> Quarter 2018

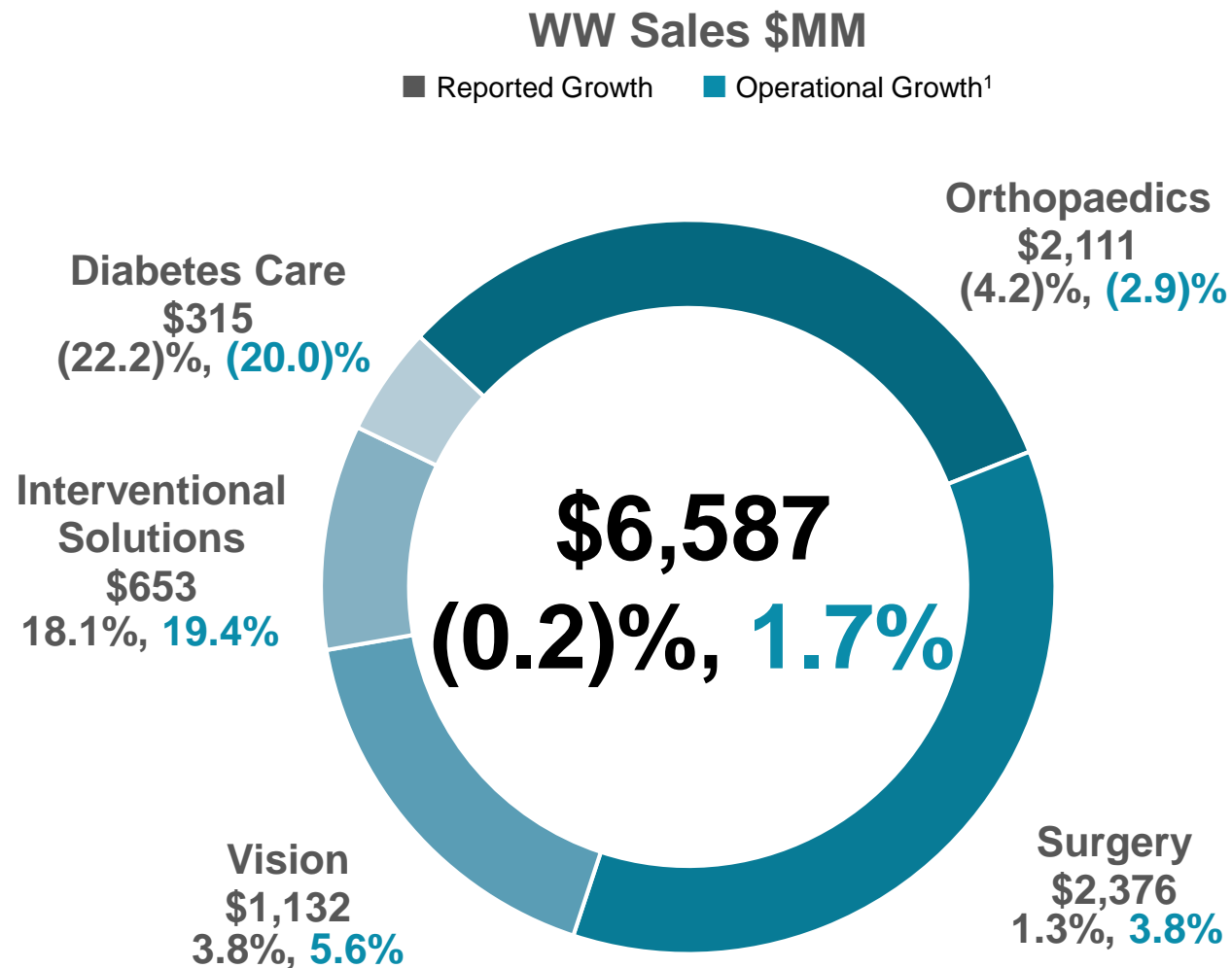
Consistent sales momentum fueled by *Interventional Solutions, Advanced Surgery and Vision*

Reported: WW (0.2)%, U.S. 0.3%, Int'l (0.6)%

Operational<sup>1</sup>: WW 1.7%, U.S. 0.3%, Int'l 3.0%

## Key Drivers of Operational Performance<sup>1</sup>

Interventional Solutions	<ul style="list-style-type: none"> <li>Electrophysiology growth of +23% primarily driven by Atrial Fibrillation procedure growth coupled with strong THERMOCOOL SMARTTOUCH SF Contact Force Sensing Catheter and diagnostic catheter sales</li> </ul>
Diabetes Care	<ul style="list-style-type: none"> <li>Pump discontinuation, BGM price declines in the U.S., as well as category and share softness in EMEA</li> </ul>
Orthopaedics	<ul style="list-style-type: none"> <li><b>Hips:</b> Driven by leadership position in the anterior approach and strong market demand for the ACTIS stem</li> <li><b>Knees:</b> Growth driven by ATTUNE Revision uptake and aided by one-time prior year impact of price legislation in India</li> <li><b>Trauma:</b> Flat due to lower market growth along with continued pricing pressure, primarily in the U.S.</li> <li><b>Spine &amp; Other:</b> Codman divestiture and share decline in Spine partially offset by new product launches                             <ul style="list-style-type: none"> <li><b>Spine:</b> WW: ~(3%), U.S. ~(2%), OUS: ~(4%)</li> </ul> </li> </ul>
Surgery	<ul style="list-style-type: none"> <li><b>Advanced:</b> <ul style="list-style-type: none"> <li><b>Endocutters:</b> +10% driven by double-digit procedure growth in China and continued success of new products across all OUS markets</li> <li><b>Biosurgery:</b> +9% driven by Topical Absorbable Hemostat and Biologics</li> <li><b>Energy:</b> +4.5% driven by OUS growth, primarily double-digit growth in ASPAC</li> </ul> </li> <li><b>General:</b> Led by Wound Closure growth driven by strength in ASPAC aided by the WHO &amp; CDC guidelines recommending our Plus Suture product</li> <li><b>Specialty:</b> Primarily driven by ASP growth</li> </ul>
Vision	<ul style="list-style-type: none"> <li><b>Contact Lenses/Other:</b> Growth due to the strength of the astigmatism and daily disposable lenses in the OASYS family partially offset by prior year comparisons</li> <li><b>Surgical:</b> Strength in international cataracts, primarily in Asia Pac / Japan</li> </ul>



Sales excl. acquisition and divestiture<sup>2</sup>: WW 2.9%, U.S. 1.2%, OUS 4.4%



<sup>1</sup> Excludes impact of translational currency

<sup>2</sup> Non-GAAP measure; see reconciliation



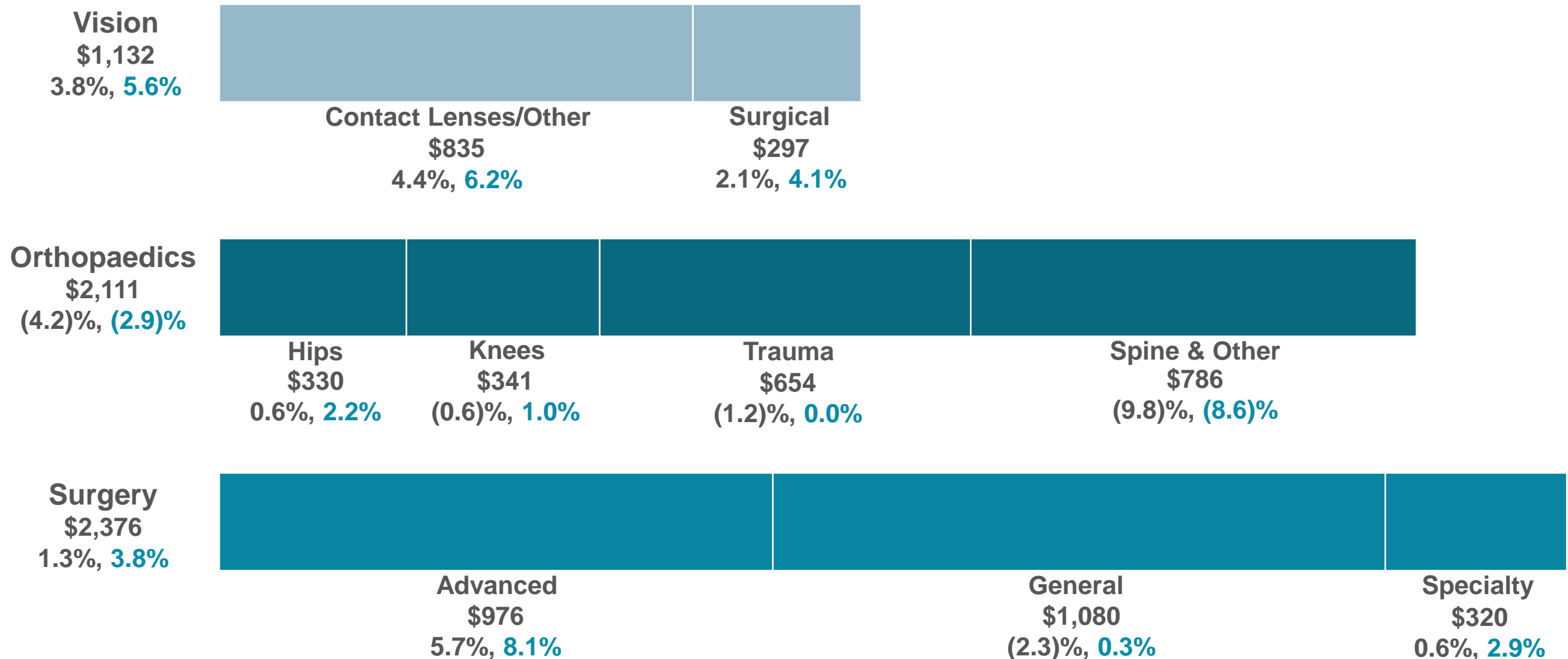


# Medical Devices Platforms

Consistent sales momentum fueled by Interventional Solutions, Advanced Surgery and Vision

## WW Sales \$MM

■ Reported Growth ■ Operational Growth<sup>1</sup>



<sup>1</sup> Excludes impact of translational currency



# Important Developments in 3<sup>rd</sup> Quarter 2018

- The U.S. Food and Drug Administration (FDA) approved and the European Commission (EC) granted marketing authorization for SYMTUZA (D/C/F/TAF), a complete darunavir-based single-tablet regimen for the treatment of HIV-1 infection
- Completed the acquisition of Zarbee's, Inc., a leader in naturally-based healthcare products
- A supplemental Biologics License Application was submitted to the FDA and a Type II Variation to the European Medicines Agency (EMA) seeking approval of a split dosing regimen for DARZALEX (daratumumab)
- Received European CE mark approval for BRAVO Flow Diverter for use in the treatment of patients suffering from intracranial aneurysms
- A supplemental New Drug Application was submitted to the FDA seeking to broaden the use of IMBRUVICA (ibrutinib) in chronic lymphocytic leukemia or small lymphocytic lymphoma to include combination use with a non-chemotherapy agent, obinutuzumab, in the frontline setting
- The FDA approved an additional indication for IMBRUVICA (ibrutinib) in combination with rituximab as a non-chemotherapy combination regimen for patients with Waldenström's Macroglobulinemia, a rare blood cancer
- A Type II Variation was submitted to the EMA seeking to expand the indication of OPSUMIT (macitentan) to include the treatment of adults with inoperable chronic thromboembolic pulmonary hypertension (CTEPH), WHO Group 4, to improve exercise capacity and pulmonary vascular resistance
- The EC granted marketing authorization for DARZALEX (daratumumab) in combination with VELCADE (bortezomib), a proteasome inhibitor, melphalan, an alkylating agent, and prednisone for the treatment of newly diagnosed multiple myeloma patients who are ineligible for autologous stem cell transplant
- A New Drug Application was submitted to the FDA for esketamine nasal spray, a rapidly acting antidepressant for treatment-resistant depression in adults
- A New Drug Application was submitted to the FDA for erdafitinib, a once-daily, oral pan-fibroblast growth factor receptor (FGFR) inhibitor for the treatment of locally advanced or metastatic urothelial cancer
- Completed the acquisition of Emerging Implant Technologies GmbH, a privately held manufacturer of 3D-printed titanium interbody implants for spinal fusion surgery
- A binding offer was accepted from Fortive Corporation to acquire Advanced Sterilization Products business for an aggregate value of approximately \$2.8 billion, subject to customary adjustments
- Launched SENTIO MMG, a first-of-its-kind digital mechanomyography platform designed to assess nerve status and identify and avoid peripheral nerves during spine surgery
- Announced the decision to discontinue the collaboration and license agreement with Geron Corporation for imetelstat
- Reassessed fair value of Alios Biopharma Inc., in-process research and development asset. This resulted in a partial impairment of approximately \$0.6 billion after-tax
- Completed the divestiture of the LifeScan business to Platinum Equity for approximately \$2.1 billion, subject to customary adjustments<sup>1</sup>
- An exclusive worldwide license agreement was entered into with Arrowhead Pharmaceuticals, Inc. to develop and commercialize a new treatment for chronic Hepatitis B viral infection<sup>1</sup>
- The FDA approved an additional indication for XARELTO (rivaroxaban) to reduce the risk of major cardiovascular (CV) events, such as CV death, myocardial infarction and stroke, in people with chronic coronary or peripheral artery disease<sup>1</sup>
- A Marketing Authorization Application was submitted to the EMA for esketamine nasal spray, a rapidly acting antidepressant for treatment-resistant depression in adults<sup>1</sup>



<sup>1</sup> Subsequent to the quarter



## **Joseph J. Wolk**

Executive Vice President,  
Chief Financial Officer





# Condensed Consolidated Statement Of Earnings

3<sup>rd</sup> Quarter 2018

(Unaudited; Dollar and Shares in Millions Except Per Share Figures)

	2018		2017*		Percent Increase (Decrease)
	Amount	Percent to Sales	Amount	Percent to Sales	
<b>Sales to customers</b>	\$ 20,348	100.0	\$ 19,650	100.0	3.6
<b>Cost of products sold</b>	6,589	32.4	6,925	35.2	(4.9)
<b>Gross Profit</b>	13,759	67.6	12,725	64.8	8.1
<b>Selling, marketing and administrative expenses</b>	5,543	27.3	5,423	27.6	2.2
<b>Research and development expense</b>	2,508	12.3	2,585	13.2	(3.0)
<b>In-process research and development</b>	1,126	5.6	-	-	
<b>Interest (income) expense, net</b>	68	0.3	155	0.8	
<b>Other (income) expense, net</b>	3	0.0	(297)	(1.5)	
<b>Restructuring</b>	88	0.4	69	0.3	
<b>Earnings before provision for taxes on income</b>	4,423	21.7	4,790	24.4	(7.7)
<b>Provision for taxes on income</b>	489	2.4	1,026	5.2	(52.3)
<b>Net earnings</b>	\$ 3,934	19.3	\$ 3,764	19.2	4.5
<b>Net earnings per share (Diluted)</b>	\$ 1.44		\$ 1.37		5.1
<b>Average shares outstanding (Diluted)</b>	2,727.6		2,737.7		
<b>Effective tax rate</b>	11.1 %		21.4 %		

<b>Adjusted earnings before provision for taxes and net earnings<sup>(1)</sup></b>					
<b>Earnings before provision for taxes on income</b>	\$ 6,780	33.3	\$ 6,573	33.5	3.1
<b>Net earnings</b>	\$ 5,590	27.5	\$ 5,208	26.5	7.3
<b>Net earnings per share (Diluted)</b>	\$ 2.05		\$ 1.90		7.9
<b>Effective tax rate</b>	17.6 %		20.8 %		

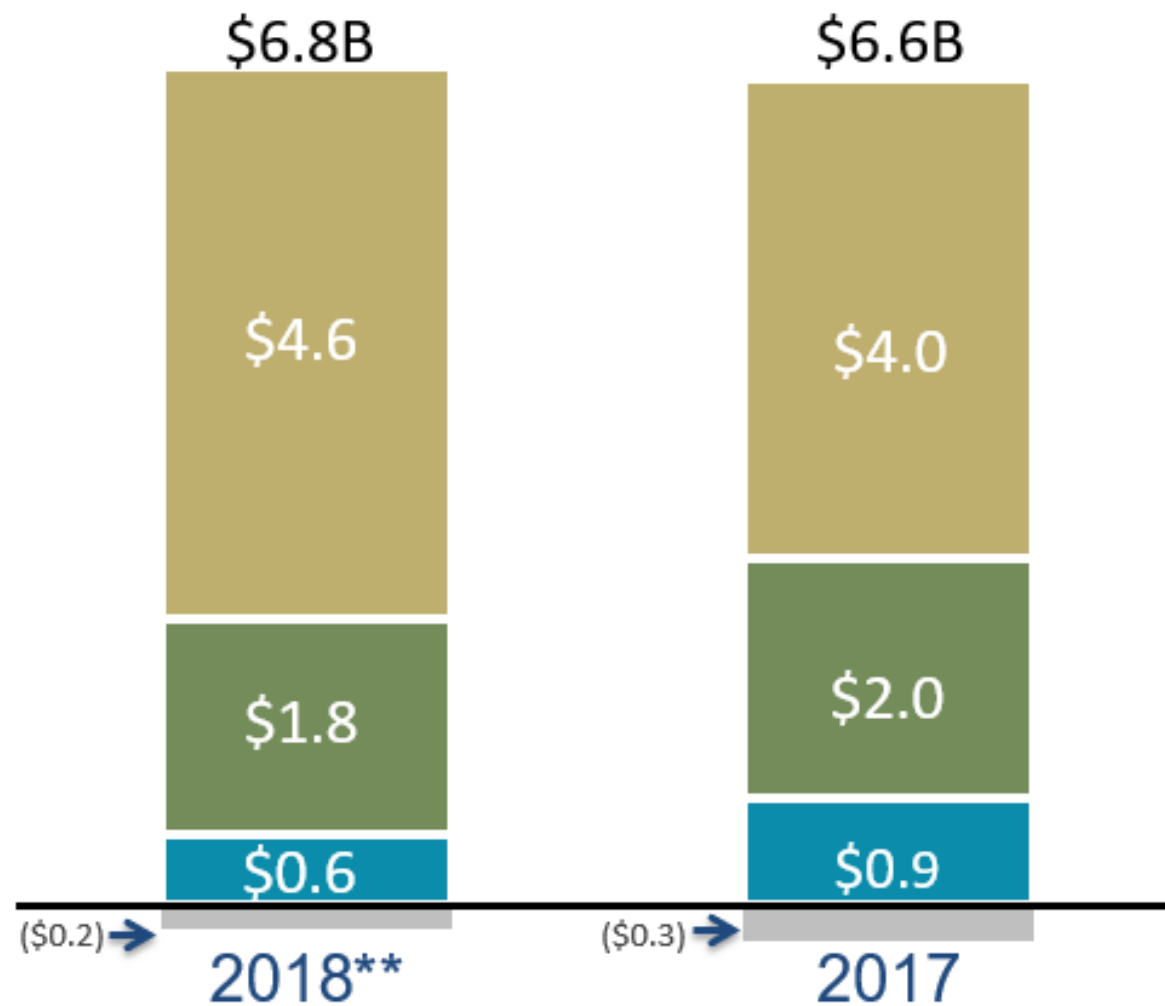


(1) See Reconciliation of Non-GAAP Financial Measures.

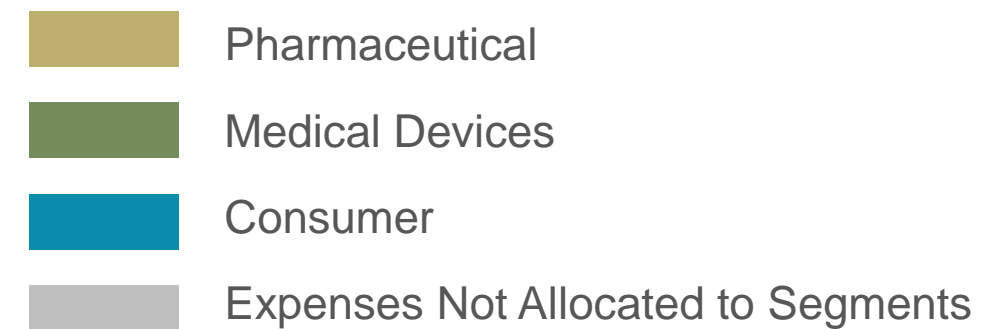
\*2017 Statement of Earnings line items have been restated to reflect impact of ASU 2017-07

# Adjusted Income Before Tax by Segment\*

3<sup>rd</sup> Quarter 2018



	% to Sales	
	Q3 2018	Q3 2017
Pharmaceutical	44.8%	41.0%
Medical Devices	27.1%	30.1%
Consumer	17.1%	28.0%
<b>Total</b>	<b>33.3%</b>	<b>33.5%</b>



\* Non-GAAP measure; excludes amortization expense and special items; see reconciliation at [www.investor.jnj.com](http://www.investor.jnj.com)

\*\* Estimated as of 10/16/18

Johnson & Johnson



# 2018 Guidance - Sales

October 2018	Estimated Operational <sup>1</sup>	Estimated Currency	Estimated Reported <sup>2</sup>
Sales Change vs. PY	\$80.6B - \$81.0B 5.5% - 6.0%	\$0.4B 0.5%	\$81.0B - \$81.4B 6.0% - 6.5%
<i>Net Impact: Acq./Div.</i>	(~1.0%)		
Sales ex. Acq./Div. Change vs. PY	4.5% - 5.0%		

July 2018	Estimated Operational <sup>1</sup>	Estimated Currency	Estimated Reported <sup>2</sup>
Sales Change vs. PY	\$79.9B - \$80.7B 4.5% - 5.5%	\$0.6B 0.8%	\$80.5B - \$81.3B 5.3% - 6.3%
<i>Net Impact: Acq./Div.</i>	(~1.0%)		
Sales ex. Acq./Div. Change vs. PY	3.5% - 4.5%		



<sup>1</sup> Excludes the impact of translational currency

<sup>2</sup> Euro Average Rate: October 2018 = \$1.18; July 2018 = \$1.19

# 2018 Guidance

	October 2018	July 2018
Adjusted Pre-Tax Operating Margin <sup>1,2</sup>	Improve by at least 150 basis points	Improve by approximately 150 basis points
Net Interest Expense	\$450 - \$500 million	\$500 - \$600 million
Net Other Income <sup>1</sup>	\$1.5 - \$1.7 billion	\$1.5 - \$1.7 billion
Effective Tax Rate <sup>1</sup>	17.5% - 18.0%	17.0% - 18.0%



<sup>1</sup> Non-GAAP measure; excludes intangible amortization expense and special items

<sup>2</sup> Sales less: COGS, SM&A and R&D expenses

# 2018 Guidance - EPS

October 2018	Estimated Operational <sup>2</sup>	Estimated Currency	Estimated Reported <sup>3</sup>
Adjusted EPS <sup>1</sup> Change vs. PY	\$7.98 - \$8.03 9.3% - 10.0%	\$0.15 2.0%	\$8.13 - \$8.18 11.4% - 12.1%

July 2018	Estimated Operational <sup>2</sup>	Estimated Currency	Estimated Reported <sup>3</sup>
Adjusted EPS <sup>1</sup> Change vs. PY	\$7.92 - \$8.02 8.5% - 9.9%	\$0.15 2.0%	\$8.07 - \$8.17 10.5% - 11.9%



<sup>1</sup> Non-GAAP measure; excludes intangible amortization expense and special items

<sup>2</sup> Excludes the impact of translational currency

<sup>3</sup> Euro Average Rate: October 2018 = \$1.18; July 2018 = \$1.19

Note: values may have been rounded



# 2018 Guidance – Sales and EPS Summary

October 2018	Estimated Operational <sup>1</sup>	Estimated Currency	Estimated Reported <sup>2</sup>
Sales Change vs. PY	\$80.6B - \$81.0B 5.5% - 6.0%	\$0.4B 0.5%	\$81.0B - \$81.4B 6.0% - 6.5%
Sales ex. Acq./Div. Change vs. PY <sup>3</sup>	4.5% - 5.0%		
Adjusted EPS <sup>4</sup> Change vs. PY	\$7.98 - \$8.03 9.3% - 10.0%	\$0.15 2.0%	\$8.13 - \$8.18 11.4% - 12.1%



<sup>1</sup> Excludes the impact of translational currency

<sup>2</sup> Euro Average Rate: October 2018 = \$1.18

<sup>3</sup> Excludes Acq./Div impact of (~1.0%)

<sup>4</sup> Non-GAAP measure; excludes intangible amortization expense and special items

Note: values may have been rounded

# Q&A



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