



APRIL 24, 2018

2018 EARNINGS

Q1 ○ ○ ○

AGENDA



INTRODUCTION AND KEY RECENT EVENTS

Dave Ricks, Chairman and Chief Executive Officer

Q1 FINANCIAL RESULTS AND FINANCIAL GUIDANCE

Josh Smiley, Senior Vice President, Finance and Chief Financial Officer

PIPELINE AND KEY FUTURE EVENTS

Dave Ricks, Chairman and Chief Executive Officer

QUESTION AND ANSWER SESSION

SAFE HARBOR PROVISION



This presentation contains forward-looking statements that are based on management's current expectations, but actual results may differ materially due to various factors. The company's results may be affected by factors including, but not limited to, the risks and uncertainties in pharmaceutical research and development; competitive developments; regulatory actions; litigation and investigations; business development transactions; economic conditions; and changes in laws and regulations, including health care reform.

For additional information about the factors that affect the company's business, please see the company's latest Forms 10-K and 10-Q filed with the Securities and Exchange Commission.

The company undertakes no duty to update forward-looking statements.

STRATEGIC DELIVERABLES

PROGRESS SINCE THE LAST EARNINGS CALL

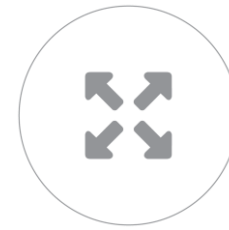


GROW REVENUE



- 9% revenue growth driven by new products
 - 4% Pharma volume growth
 - 30% Diabetes volume growth

EXPAND MARGINS



- Excluding FX on international inventories sold, non-GAAP:
 - gross margin as a % of revenue increased nearly 70bp
 - operating income % of revenue was 30.4%, an increase of 775bp

DEPLOY CAPITAL TO CREATE VALUE



- Collaboration with Sigilon to develop encapsulated cell therapies
- Distributed nearly \$600 million via the dividend
- Repurchased \$1.1 billion of stock

SUSTAIN FLOW OF INNOVATION



- Approval and launch of new Verzenio™ indication in 1L mBC
- Positive Phase 3 NILEX readouts for Taltz® and Cyramza®

KEY EVENTS SINCE THE LAST EARNINGS CALL



COMMERCIAL

- Launched Verzenio (abemaciclib) in the U.S. for the treatment of first line metastatic breast cancer;
- Launched Taltz (ixekizumab) in Germany for the treatment of psoriatic arthritis; and
- Launched Credelio® (lotilaner) in the U.S., to treat and control ticks and fleas in dogs.

REGULATORY

- The U.S. FDA approved Verzenio (abemaciclib) in combination with an aromatase inhibitor for the treatment of breast cancer based on the MONARCH 3 study; and
- The U.S. FDA's Arthritis Advisory Committee recommended approval of the 2mg, but not the 4mg, dose of baricitinib for the treatment of moderately-to-severely active rheumatoid arthritis.

CLINICAL

- Announced top-line results of the COAST-V study for Taltz (ixekizumab) for the treatment of Ankylosing Spondylitis (AS). The trial met the primary and all key secondary endpoints;
- Announced top-line results of the REACH-2 study for Cyramza (ramucirumab) in the second-line treatment of patients with Hepatocellular Carcinoma (liver cancer). The trial met its primary endpoint of overall survival (OS) as well as the secondary endpoint of progression-free survival (PFS);

CLINICAL (continued)

- Announced top-line results of the RANGE study for Cyramza (ramucirumab) in the treatment of patients with advanced or metastatic Urothelial Cancer (bladder cancer). A positive trend was observed in the secondary endpoint of overall survival (OS) which did not reach the level of statistical significance;
- At AACR, presented detailed data from the final MONARCH 3 PFS analysis showing that Verzenio plus an aromatase inhibitor achieved a median PFS of 28.2 months versus 14.8 months in the control arm;
- At AACR, presented updated Verzenio subgroup analyses from MONARCH 2 and 3 which further demonstrated that patients with certain concerning clinical characteristics received relatively greater benefit from the addition of Verzenio to endocrine therapy;
- Along with Merck, presented data at AACR from KEYNOTE-189 studying the combination of pemetrexed and pembrolizumab in non-small cell lung cancer; and
- At AAD, presented data showing that treatment with Taltz (ixekizumab) resulted in improvement in impact of genital psoriasis on sexual activity.

BUSINESS DEVELOPMENT & OTHER

- Announced a collaboration with Sigilon to develop encapsulated cell therapies for the potential treatment of type 1 diabetes;
- Distributed nearly \$600 million to shareholders via the dividend; and
- Repurchased \$1.1 billion of stock.

COMPARISON MEASURES



“REPORTED” RESULTS

Include all financial results as reported in accordance with Generally Accepted Accounting Principles (GAAP)

“NON-GAAP” MEASURES

Start with “REPORTED” RESULTS

Include adjustments for items such as:

- Asset impairment, restructuring and other special charges
- Acquired in-process R&D charges and other income and expenses from business development activities
- Amortization of intangible assets

Q1 2018 INCOME STATEMENT - REPORTED



Millions; except per share data

	<u>Q1 2018</u>	<u>Change</u>
Total Revenue	\$5,700	9%
Gross Margin	72.4%	(1.8pp)
Total Operating Expense*	2,755	(29)%
Operating Income	1,374	NM
Other Income (Expense)	67	(14)%
Effective Tax Rate	15.5%	NM
Net Income (Loss)	\$1,217	NM
EPS	\$1.16	NM

* Includes research and development expense, marketing, selling and administrative expense, acquired in-process research and development charges, and asset impairment, restructuring and other special charges.

NM – not meaningful

RECONCILIATION OF GAAP REPORTED TO NON-GAAP ADJUSTED INFORMATION; CERTAIN LINE ITEMS (UNAUDITED)



Millions; except per share data

Q1 2018

	GAAP Reported	Adjust- ments	Non-GAAP Adjusted	Non-GAAP Adjusted Change
Total Revenue	\$5,700	-	\$5,700	9%
Gross Margin	72.4%	2.7%	75.1%	(2.7pp)
Total Operating Expense	2,755	(80)	2,676	(5)%
Operating Income	1,374	231	1,604	29%
Other Income (Expense)	67	-	67	(14)%
Effective Tax Rate	15.5%	0.4%	15.9%	-5.3pp
Net Income (Loss)	\$1,217	\$189	\$1,406	35%
EPS	\$1.16	\$0.18	\$1.34	37%

Note: Numbers may not add due to rounding; see slide 20 for a complete list of significant adjustments.

RECONCILIATION OF GAAP REPORTED TO NON-GAAP ADJUSTED INFORMATION; CERTAIN LINE ITEMS (UNAUDITED)



Millions; except per share data

	Q1 2018	Q1 2017	Change
EPS (reported)	\$1.16	(\$0.10)	NM
Amortization of intangible assets	0.12	0.11	
Asset impairment, restructuring, and other special charges	0.06	0.16	
Acquired in-process R&D	-	0.81	
Inventory Step-Up	-	0.01	
EPS (non-GAAP)	\$1.34	\$0.98	37%

Note: Numbers may not add due to rounding; see slide 20 for more details on these significant adjustments.

EFFECT OF PRICE/RATE/VOLUME ON REVENUE



Millions

Q1 2018

Pharmaceuticals	Amount	Price	FX Rate	Volume	Total	CER
U.S.	\$2,779.0	8%	-	2%	10%	10%
Europe	893.8	(5%)	15%	7%	17%	2%
Japan	536.8	(3%)	5%	5%	6%	1%
Rest of World	729.1	(0%)	5%	5%	9%	4%
Total Pharma	4,938.7	3%	4%	4%	11%	7%
Animal Health	761.3	5%	3%	(8%)	(1%)	(4%)
Total Revenue	\$5,700.0	3%	4%	2%	9%	5%

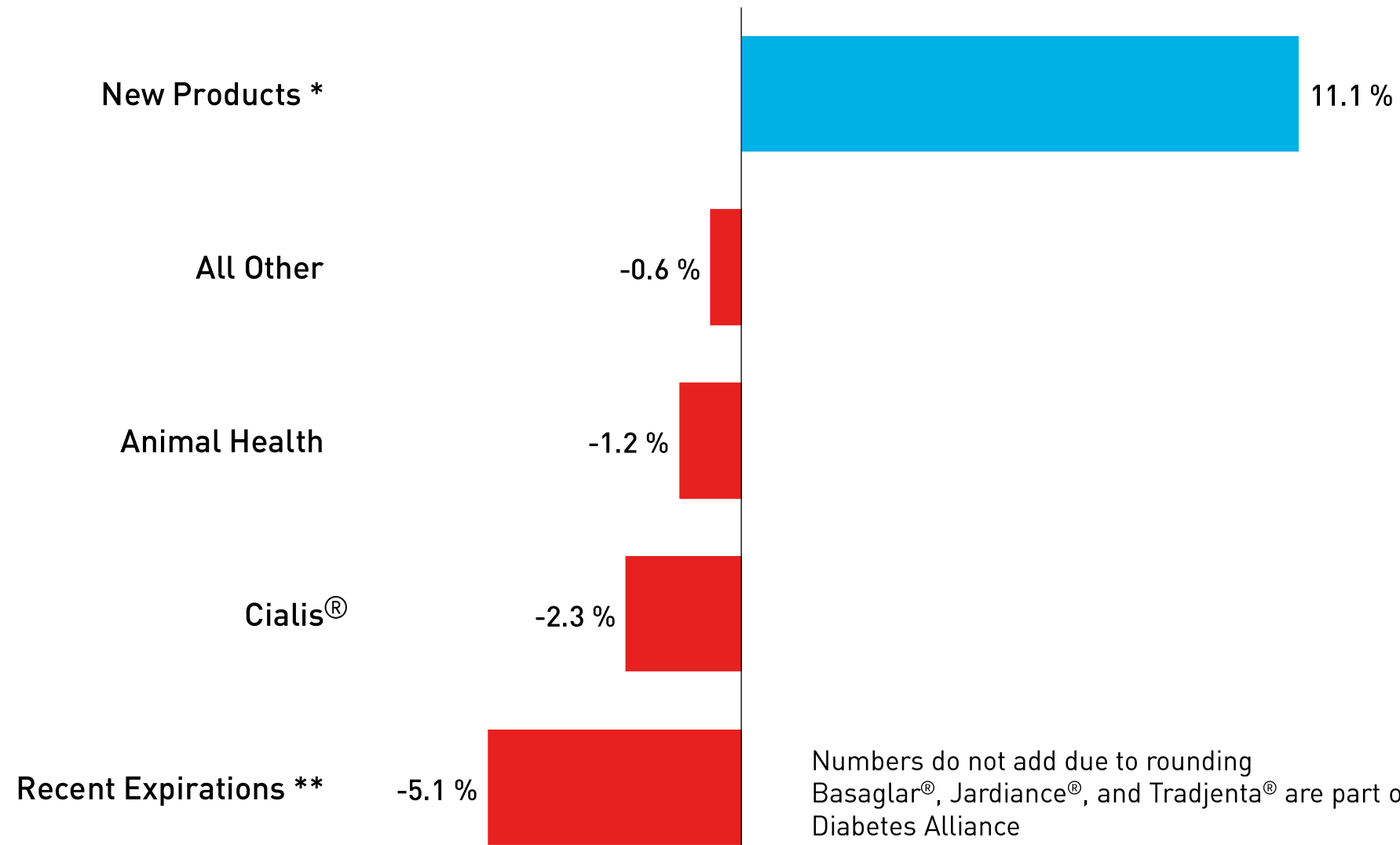
Note: Numbers may not add due to rounding.

CER = price change + volume change

NEW PRODUCTS DRIVING WW VOLUME GROWTH



Contribution to 2% Q1 WW Volume Growth

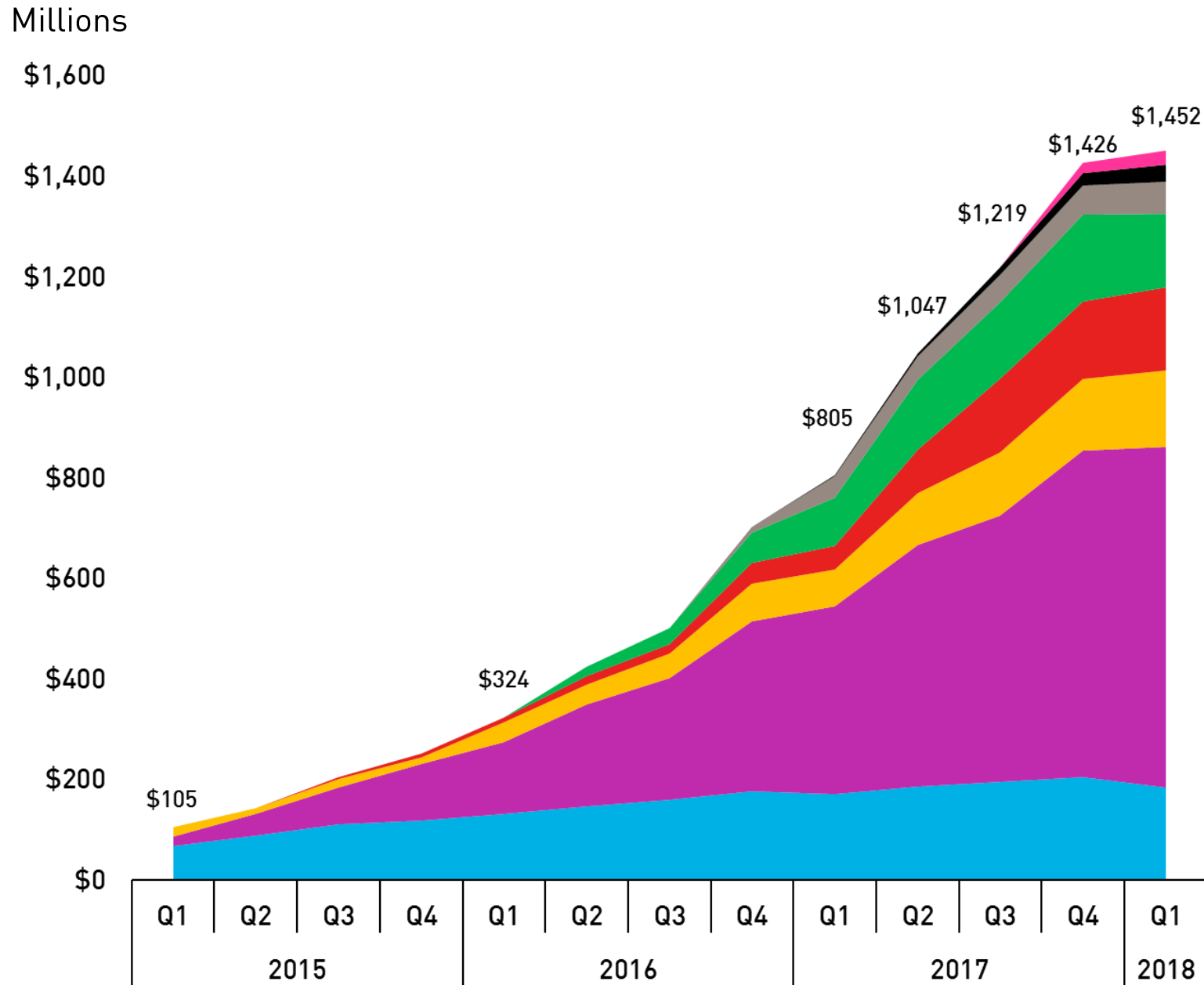


Numbers do not add due to rounding
Basaglar®, Jardiance®, and Tradjenta® are part of the Boehringer Ingelheim and Lilly Diabetes Alliance

* includes Basaglar, Cyramza, Jardiance, Lartruvo™, Olumiant®, Portrazza®, Taltz, Trulicity®, and Verzenio

** includes Axiron®, Cymbalta®, Effient®, Evista®, Strattera®, and Zyprexa®

UPDATE ON NEW PRODUCT LAUNCH PROGRESS



- VERZENIO**
 - Launched in U.S. in Q4'17; U.S. NBRx at nearly 15%
 - Launched 1L metastatic breast cancer in Q1'18 in U.S.
- OLUMIANT**
 - Strong early uptake in Germany
 - Launched in over 40 markets
- LARTRUVO**
 - Strong uptake in U.S.; European launches ongoing
- TALTZ**
 - IL-17 NBRx class at approx. 30% SOM in dermatology
 - Launched PsA in Q1'18 in U.S. and Germany
- BASAGLAR**
 - 2nd in U.S. NBRx with over 27% SOM
 - U.S. TRx SOM gain of 500bp within Q1'18
- JARDIANCE**
 - Market leader in U.S. TRx (37% SOM) and NBRx (47% SOM)
- TRULICITY**
 - U.S. TRx grew 70% over Q1'17
 - GLP-1 class TRx growing nearly 27% in U.S., up from 23% in Q4'17
- CYRAMZA**
 - Nearly 67% SOM in 2L metastatic gastric cancer in Japan

Note: Jardiance is sold by Boehringer Ingelheim; Lilly records as revenue its share of Jardiance gross margin
 Jardiance and Basaglar are part of the Boehringer Ingelheim and Lilly Diabetes Alliance

EFFECT OF FOREIGN EXCHANGE ON 2018 RESULTS



Year-on-Year Growth

Q1 2018

Reported	With FX	w/o FX
Total Revenue	9%	5%
Cost of Sales	17%	(1)%
Gross Margin	6%	7%
Operating Expense	(29)%	(31)%
Operating Income	NM	NM
EPS	NM	NM

Non-GAAP

Total Revenue	9%	5%
Cost of Sales	22%	2%
Gross Margin	5%	6%
Operating Expense	(5)%	(8)%
Operating Income	29%	40%
EPS	37%	47%

2018 GUIDANCE



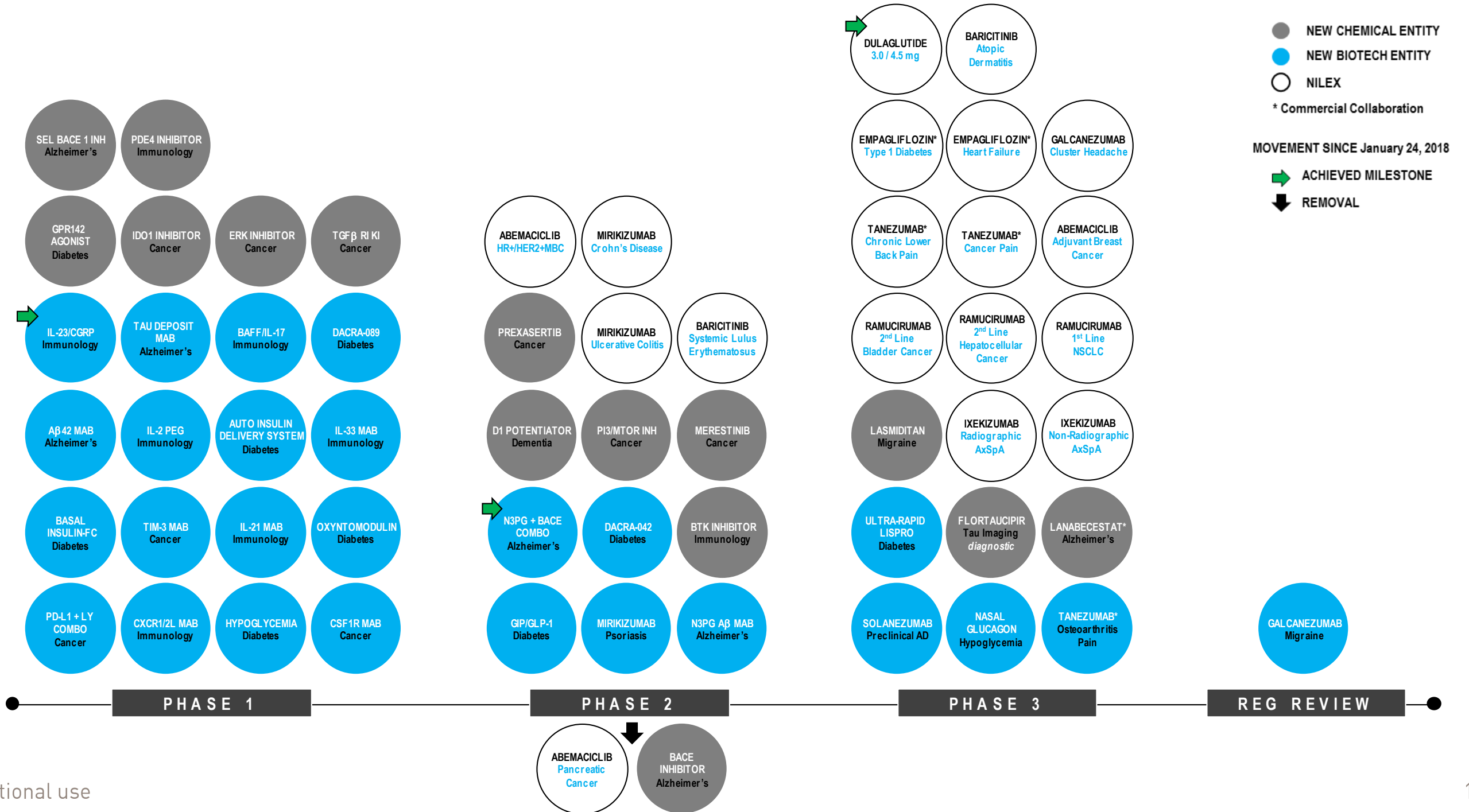
	<u>Prior</u>	<u>Updated</u>	<u>Comments</u>
Total Revenue	\$23.0 - \$23.5 billion	\$23.7 - \$24.2 billion	Favorable segment & payer mix and FX rate movements
Gross Margin % (GAAP)	Approx. 73%	Unchanged	
Gross Margin % (non-GAAP)	Approx. 75%	Unchanged	
Mktg, Selling & Admin.	\$6.1 - \$6.4 billion	\$6.2 - \$6.5 billion	FX rates
Research & Development	\$5.0 - \$5.2 billion	\$5.2 - \$5.4 billion	Funding additional NILEX and FX rates
Other Income/(Expense)	\$75 - \$175 million	\$75 - \$200 million	
Tax Rate (GAAP)	Approx. 18%	Approx. 17%	Favorable jurisdictional mix
Tax Rate (non-GAAP)	Approx. 18%	Approx. 17%	Favorable jurisdictional mix
Earnings per Share (GAAP)	\$4.39 - \$4.49	\$4.52 - \$4.62	See above
Earnings per Share (non-GAAP)	\$4.81 - \$4.91	\$5.10 - \$5.20	See above
Capital Expenditures	Approx. \$1.2 billion	Unchanged	

FX rates for current guidance:

- Euro at 1.23
- Yen at 106.5
- Pound at 1.41

LILLY SELECT NME AND NILEX PIPELINE

APRIL 20, 2018



POTENTIAL KEY EVENTS 2018



PHASE 3 INITIATIONS

Baricitinib for psoriatic arthritis
Mirikizumab for psoriasis
Mirikizumab for ulcerative colitis

✓+ Dulaglutide alternate doses for type 2 diabetes

Empagliflozin for chronic kidney disease¹

PHASE 3 DATA INTERNAL READOUTS

Flortaucipir (18F AV-1451) tau imaging agent
Tanezumab for osteoarthritis pain (dosing study)²
Tradjenta CAROLINA CV outcomes study¹
Trulicity REWIND CV outcomes study
Ultra rapid insulin for type 1 and type 2 diabetes

✓- Ramucirumab RANGE for 2L bladder cancer (final analysis)

Ramucirumab RELAY for 1L EGFR NSCLC cancer (PFS readout)

PHASE 3 DATA EXTERNAL DISCLOSURES

Galcanzumab for cluster headache
Ixekizumab for axial spondyloarthritis
Empagliflozin for type 1 diabetes¹
Tradjenta CARMELINA CV outcomes study¹
Ramucirumab REACH 2 in 2L high AFP hepatocellular cancer

✓+ Alimta+platinum+Keytruda in 1L nonsquamous NSCLC (KN-189)³

REGULATORY SUBMISSIONS

Lasmiditan for acute migraine
Empagliflozin + linagliptin + metformin XR (US)¹
Nasal glucagon for hypoglycemia

REGULATORY ACTIONS

Baricitinib for rheumatoid arthritis (US)
Galcanzumab for migraine prevention
✓+ Ixekizumab for psoriatic arthritis (EU)
Abemaciclib + fulvestrant for 2L breast cancer (MONARCH 2) (EU/J)
Abemaciclib + AIs for 1L breast cancer (MONARCH 3) (US ✓+ /EU/J)
Alimta[®] sNDA to include KEYNOTE-021G data (US)³
Fruquintinib for 3L metastatic colorectal cancer (China)⁴

OTHER

Rulings in ongoing Alimta patent litigation:
US IPR Appeal to CAFC
US alternative salt forms
Japan (Nipro)
Germany

¹ in collaboration with Boehringer Ingelheim

² in collaboration with Pfizer

³ in collaboration with Merck

⁴ in collaboration with Hutchison China MediTech

SUMMARY



- Q1 2018 **revenue growth** of 9%, driven by new products
- Excluding FX, non-GAAP EPS growth of 47% and **operating margin expansion** of 775 basis points
- Progress on our **innovation-based strategy** included: the launch of an additional indication for Verzenio and positive Phase 3 readouts for Taltz and Cyramza
- Deployed \$1.7 billion back to shareholders through the dividend and stock repurchases



GROW REVENUE

- Minimum average annual revenue growth of 5% in constant currency from 2015 through 2020



EXPAND MARGINS

- Excluding FX on int'l inventories sold, minimum operating margin % of revenue of 30% in 2020



SUSTAIN FLOW OF INNOVATION

- Potential to launch 20+ new molecules in 10 years (2014-2023)
- On average, could launch 2+ new indications or line extensions per year

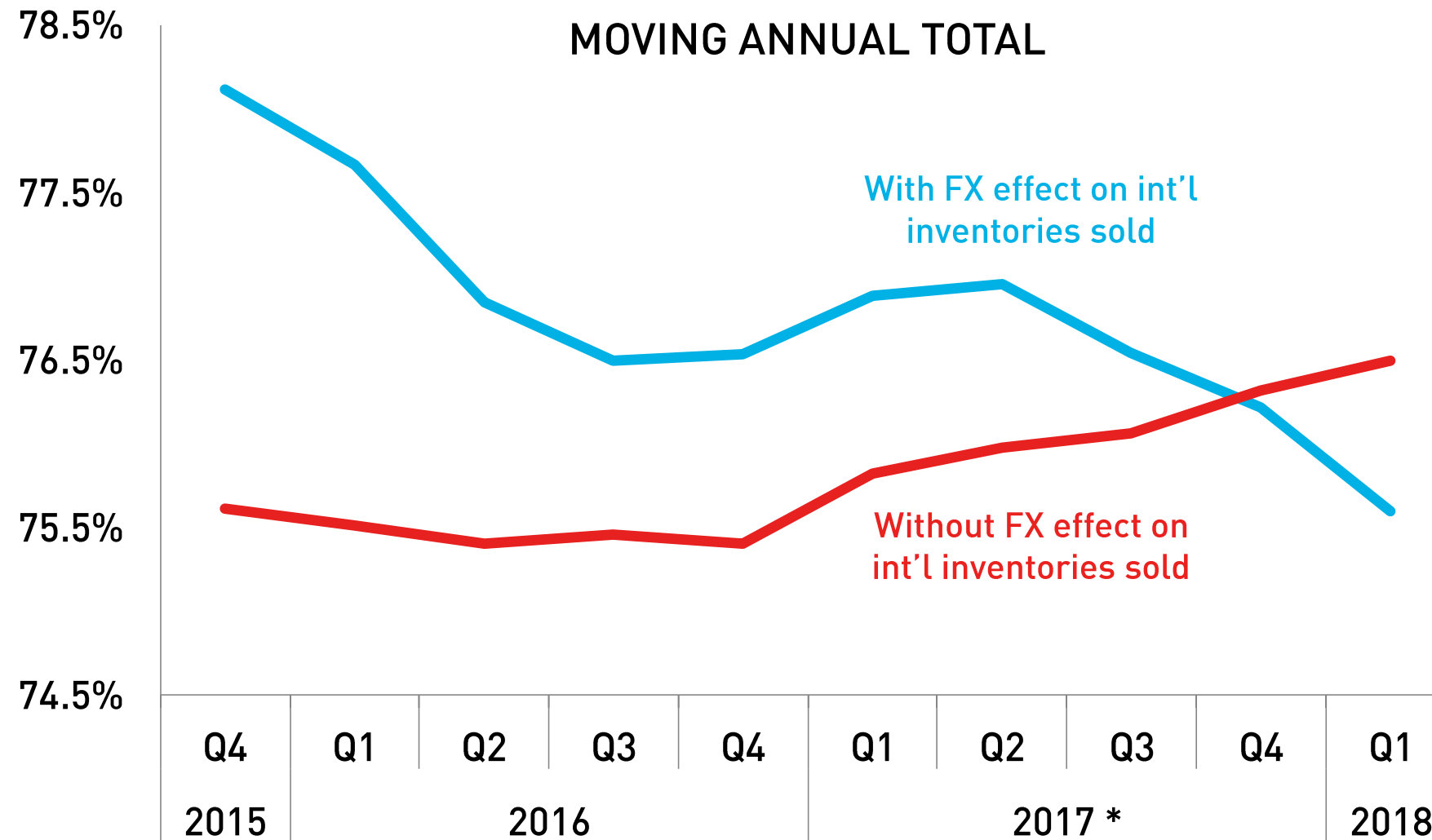


DEPLOY CAPITAL TO CREATE VALUE

- Fund existing marketed and pipeline products
- Bolster growth prospects via business development in focus areas
- Annual dividend increases

Supplementary Slides

NON-GAAP GROSS MARGIN % OF REVENUE



Individual quarter GM % of Revenue:

with FX effect on int'l inv sold	77.3%	76.3%	76.0%	76.4%	77.4%	77.8%	76.3%	74.8%	76.1%	75.1%
w/o FX effect on int'l inv sold	75.7%	74.9%	75.7%	75.5%	75.5%	76.7%	76.3%	75.8%	76.5%	77.4%

Note: The lines in the graph are moving annual totals (i.e. trailing 4 quarters) while the two rows of numbers are from specific quarters.

* 2017 has been reclassified to reflect changes to pension and post-retirement benefit cost accounting effective Jan 1, 2018

Q1 2018 INCOME STATEMENT NOTES



Q1 2018 NON-GAAP INFORMATION HAS BEEN ADJUSTED TO ELIMINATE:

- amortization of intangible assets primarily associated with costs of marketed products acquired or licensed from third parties totaling \$152.4 million (pretax), or \$0.12 per share (after-tax); and
- asset impairment, restructuring and other special charges of \$78.3 million (pretax), or \$0.06 per share (after-tax), primarily associated with the decision to end Posilac[®] (rbST) production at the Augusta, Georgia manufacturing site, as well as expenses associated with the review of strategic alternatives for the Elanco Animal Health business.

Q1 2017 NON-GAAP INFORMATION HAS BEEN ADJUSTED TO ELIMINATE:

- amortization of intangible assets primarily associated with costs of marketed products acquired or licensed from third parties totaling \$176.1 million (pretax), or \$0.11 per share (after-tax);
- charges primarily related to severance costs incurred as a result of actions taken to reduce the company's cost structure, as well as integration costs related to the acquisition of Novartis Animal Health, totaling \$213.9 million, or \$0.16 per share (after-tax);
- an acquired in-process research and development charge related to the acquisition of CoLucid Pharmaceuticals totaling \$857.6 million (pretax), or \$0.81 per share (after-tax); and
- inventory step-up costs associated with the acquisition of Boehringer Ingelheim Vetmedica's U.S. feline, canine, and rabies vaccine portfolio totaling \$10.4 million (pretax), or \$0.01 per share (after-tax).

COMPARATIVE EPS SUMMARY 2017/2018



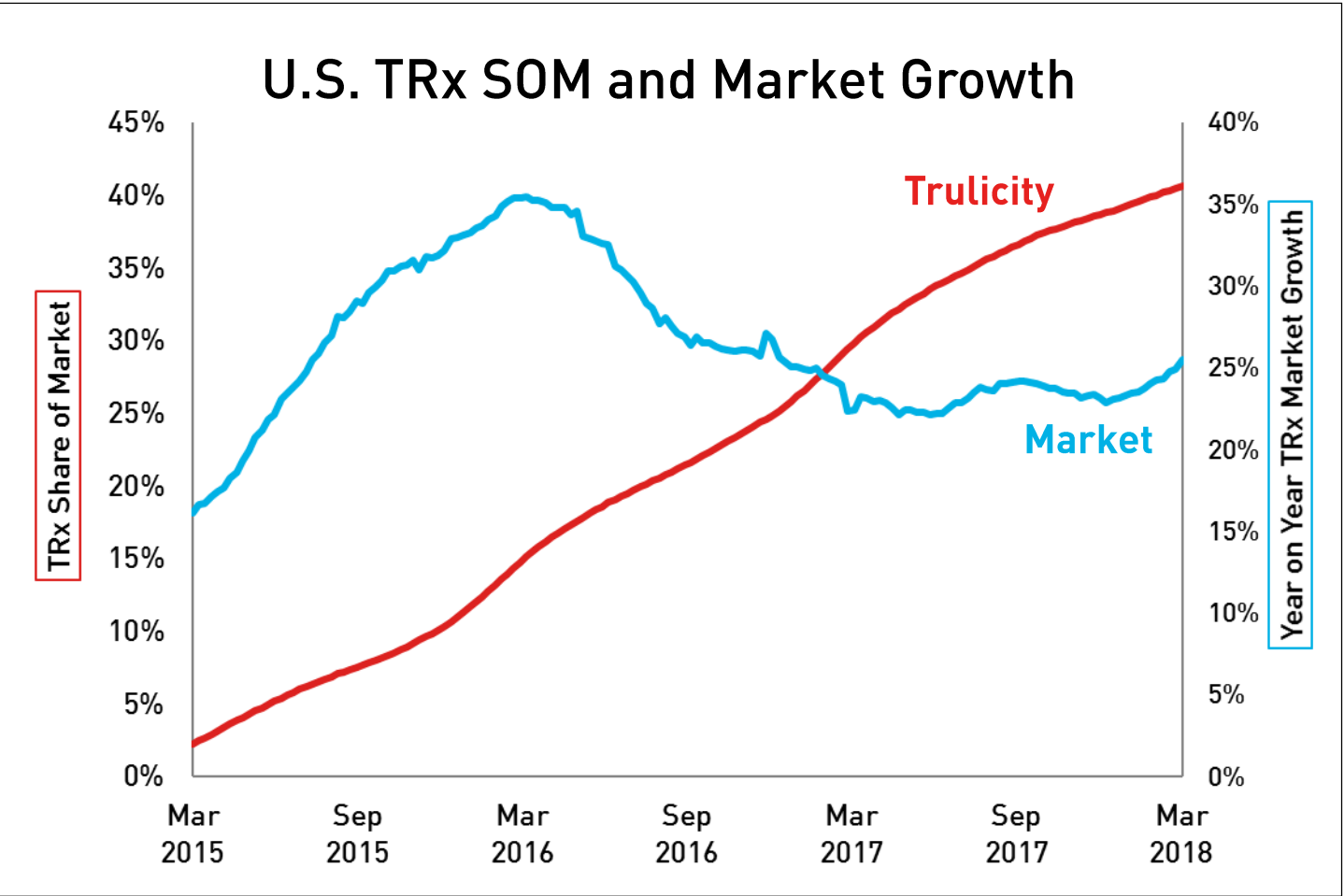
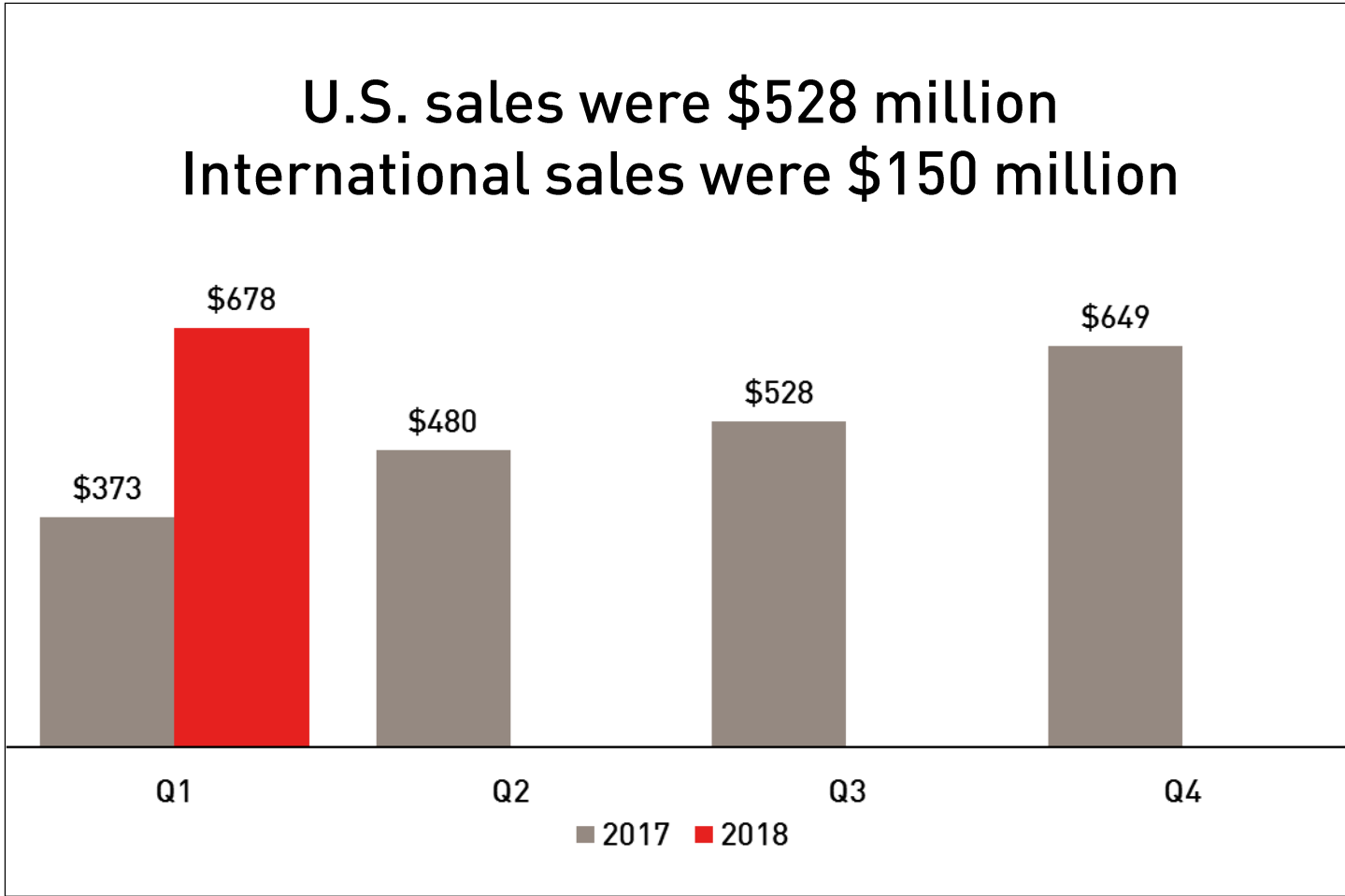
	1Q17	2Q17	3Q17	4Q17	2017	1Q18	2Q18	3Q18	4Q18	2018
Reported	(0.10)	0.95	0.53	(1.58)	(0.19)	1.16				
Non-GAAP	0.98	1.11	1.05	1.14	4.28	1.34				

Note: Numbers may not add due to rounding.
 For a complete reconciliation to reported earnings, see slide 20 and our earnings press release dated April 24, 2018.

Q1 2018 TRULICITY SALES INCREASED 82%



Millions



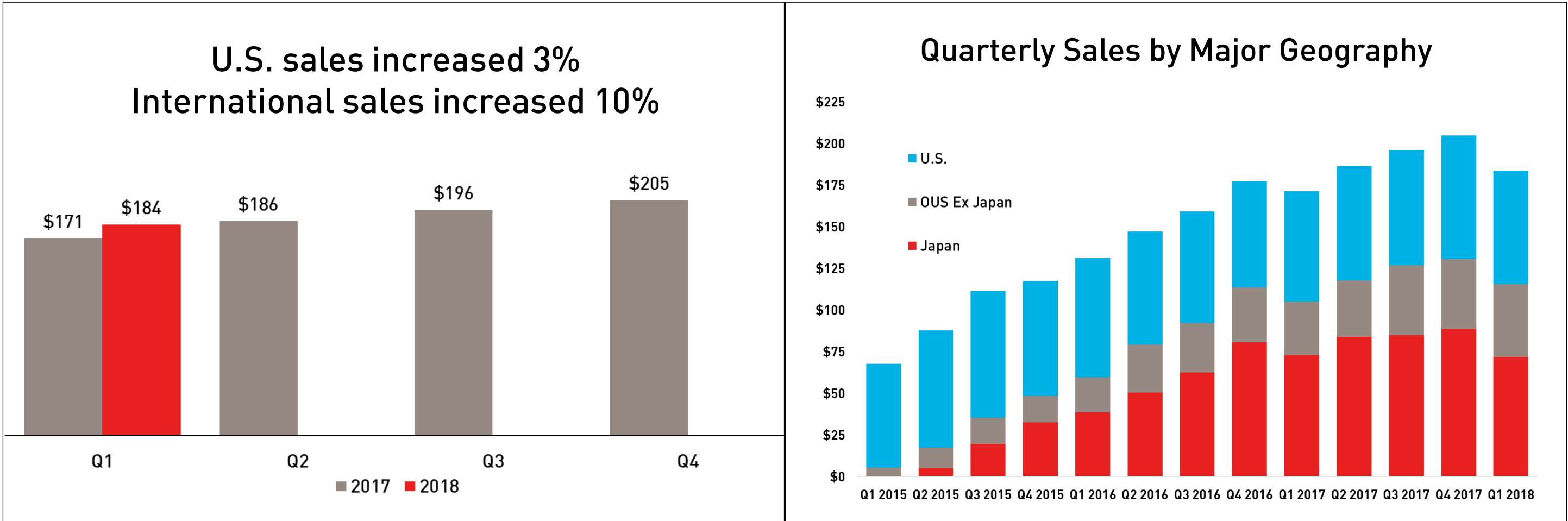
Note: Numbers may not add due to rounding.

Source: IQVIA NPA TRx 3MMA, weekly data March 30, 2018

Q1 2018 CYRAMZA SALES INCREASED 7%



Millions

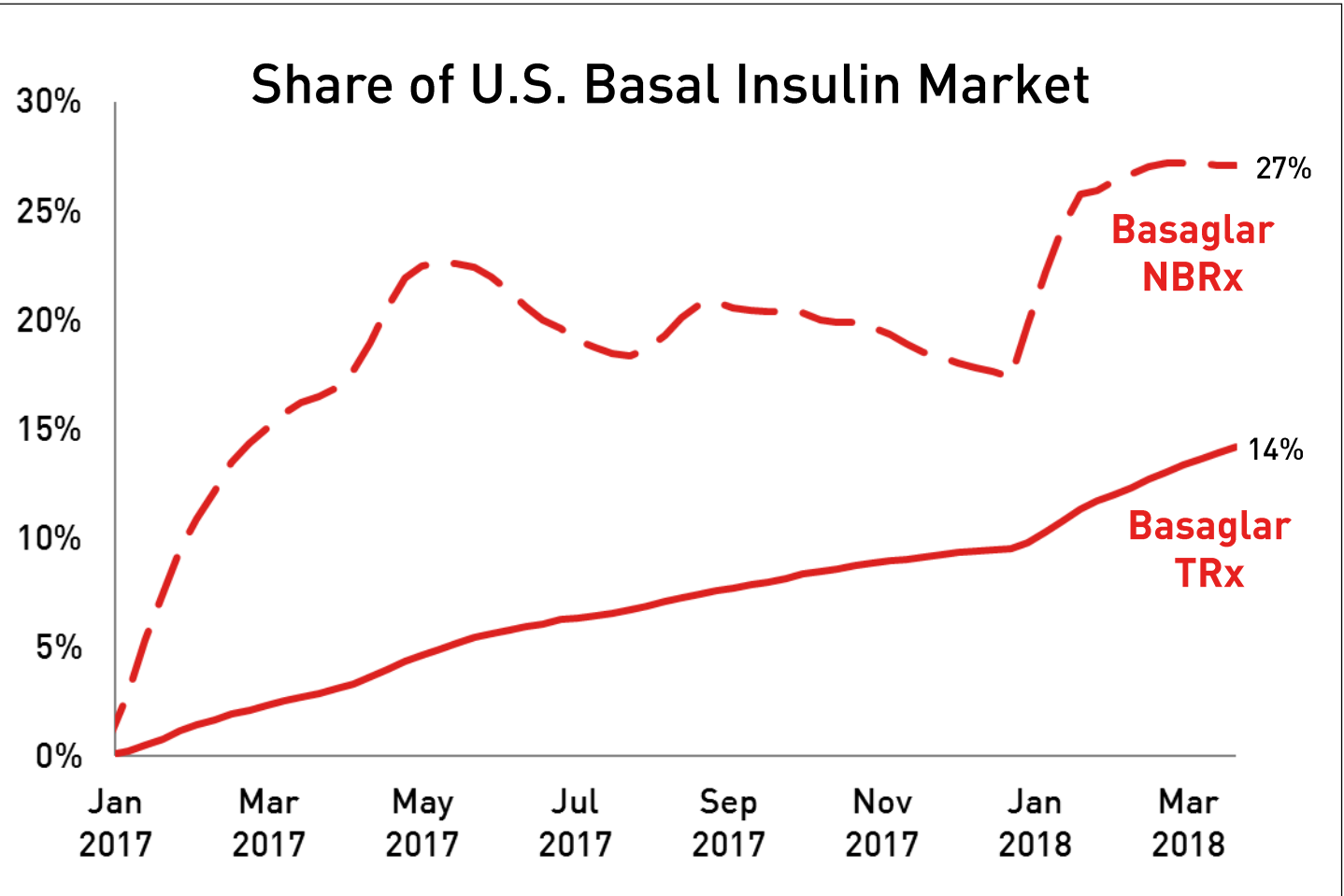
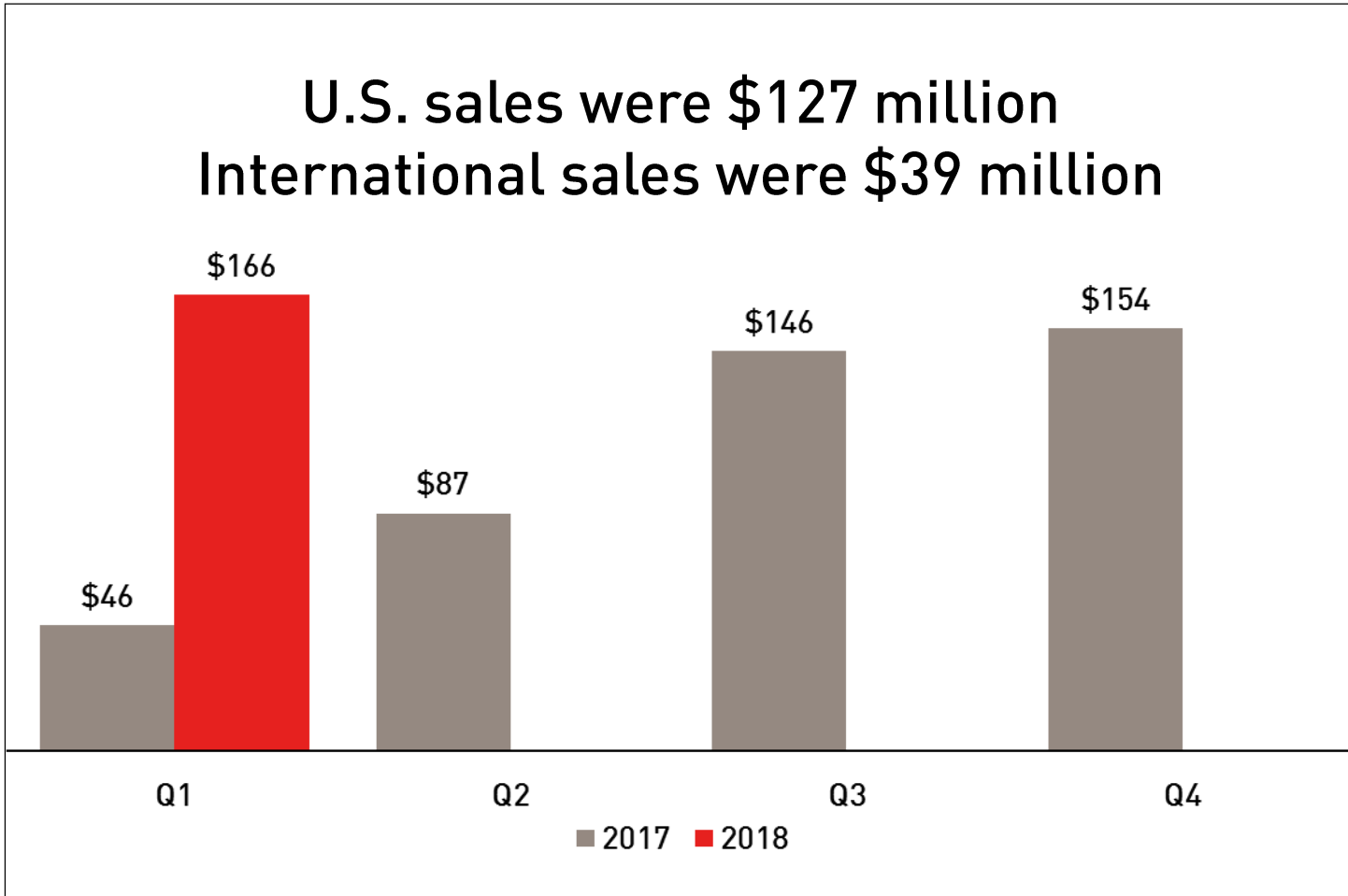


Note: Numbers may not add due to rounding.

Q1 2018 BASAGLAR SALES WERE \$166 MILLION



Millions



Note: Numbers may not add due to rounding.

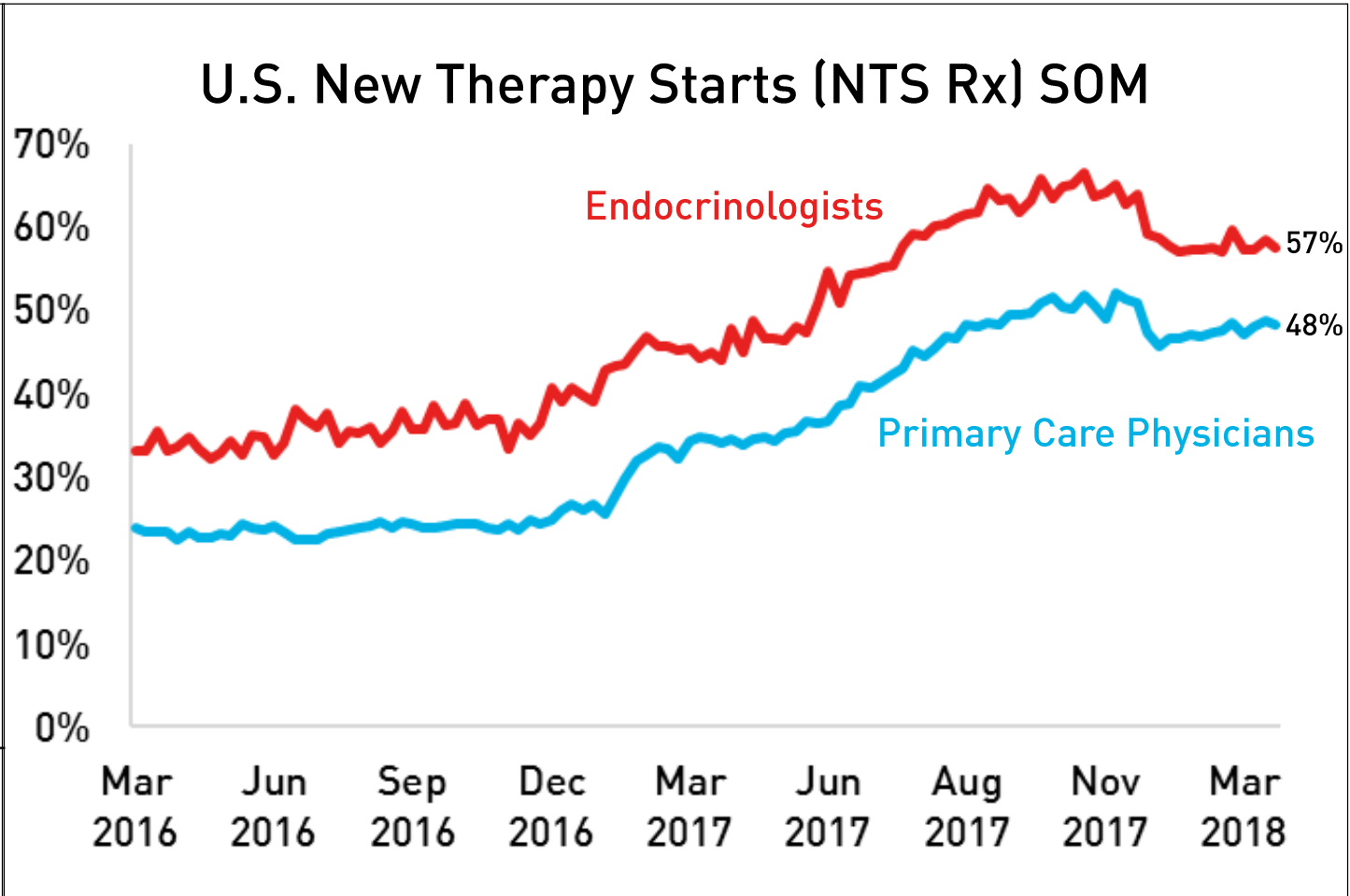
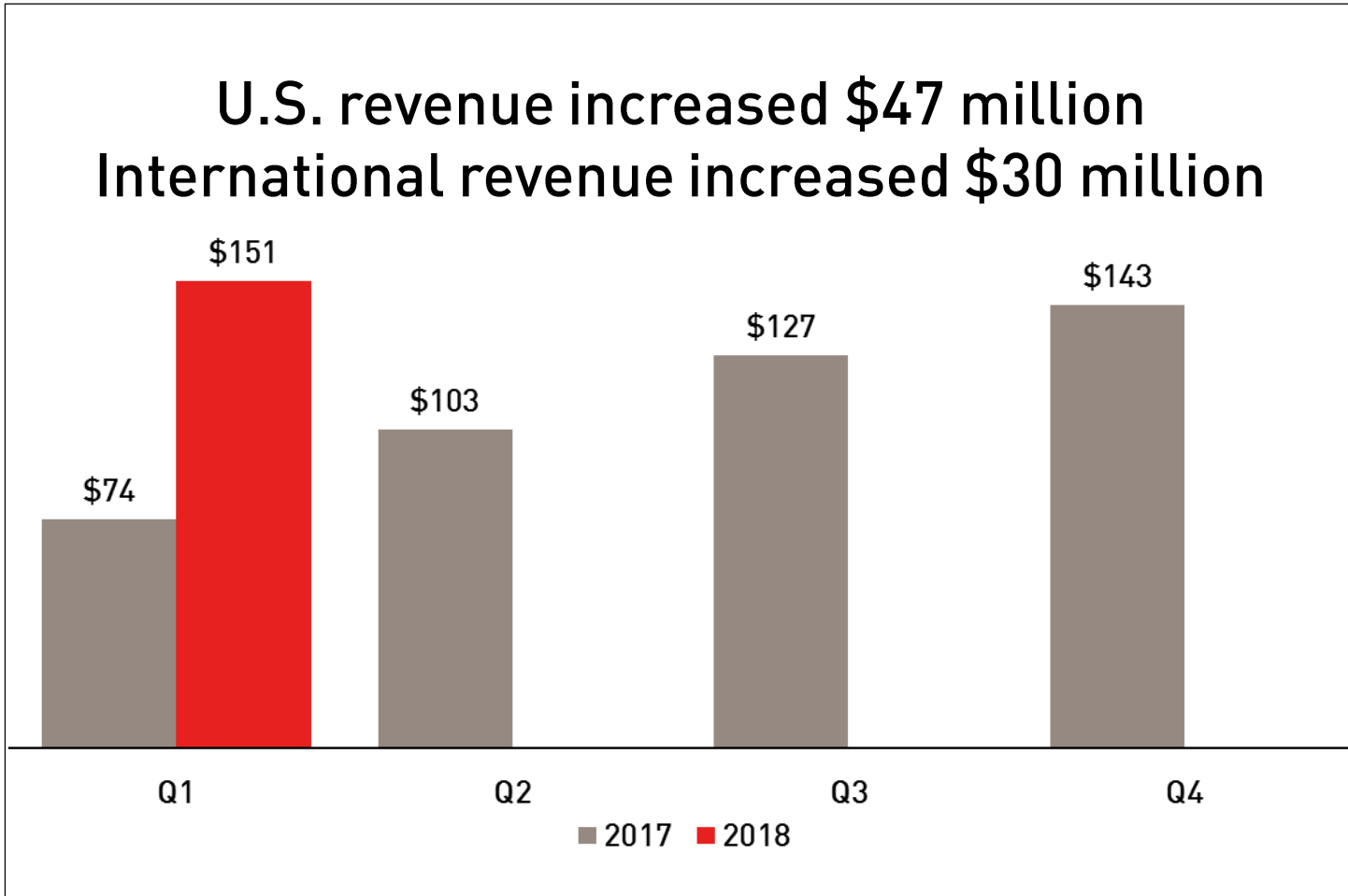
Source: IQVIA NPA TRx 3MMA, weekly data March 30, 2018

Note: Basaglar is part of the Boehringer Ingelheim and Lilly Diabetes Alliance

Q1 2018 JARDIANCE REVENUE WAS \$151 MILLION



Millions



Note: Numbers may not add due to rounding.

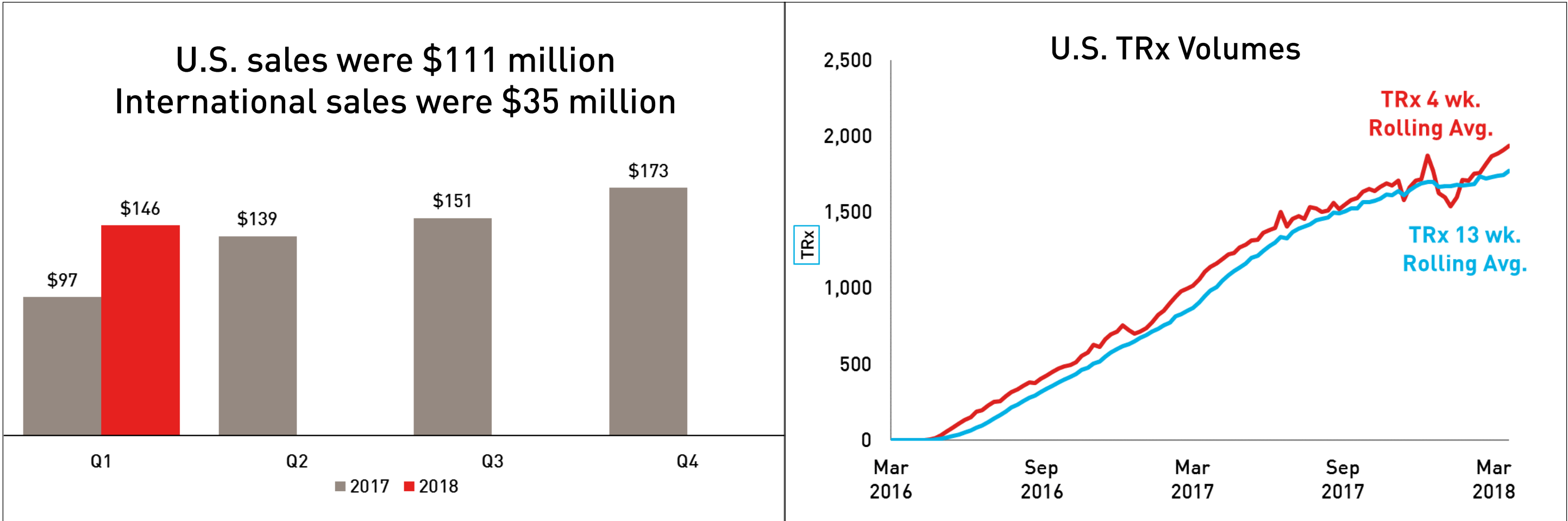
Source: IQVIA NPA TRx 3MMA, weekly data March 30, 2018

Note: Jardiance is part of the Boehringer Ingelheim and Lilly Diabetes Alliance

Q1 2018 TALTZ SALES INCREASED 52%



Millions



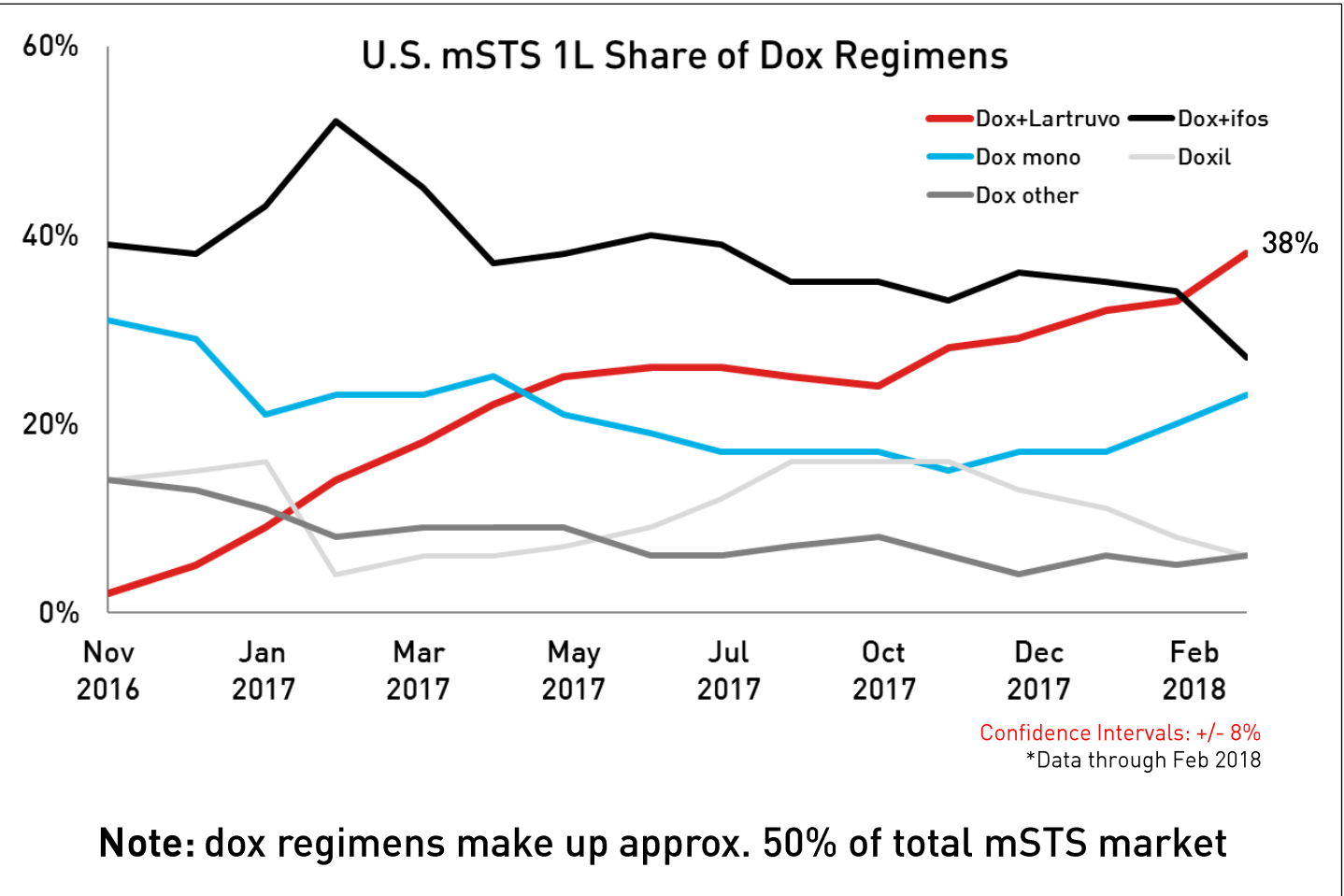
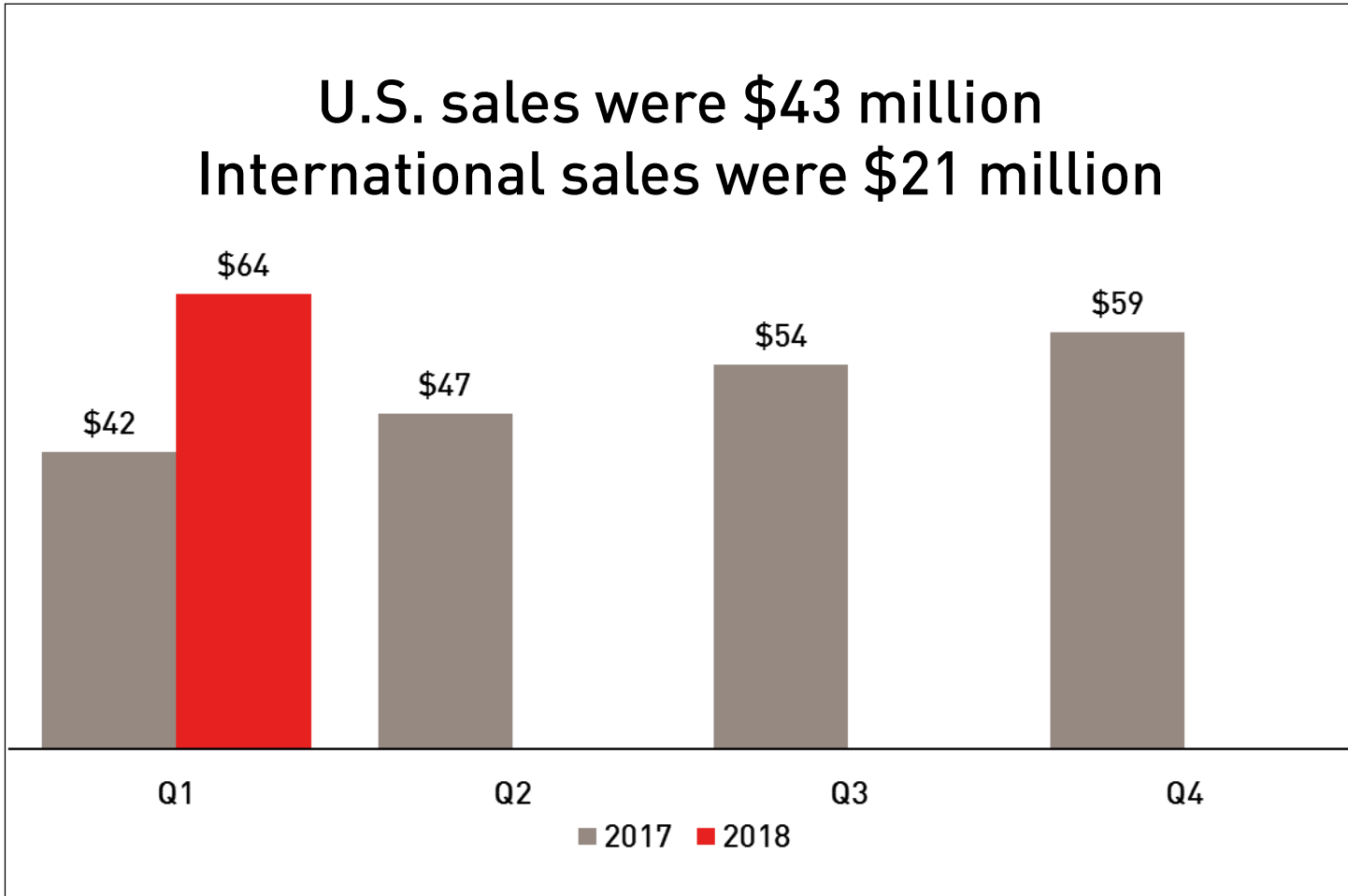
Note: Numbers may not add due to rounding.

Source: IQVIA NPA TRx 3MMA, weekly data March 30, 2018

Q1 2018 LARTRUVO SALES WERE \$64 MILLION



Millions

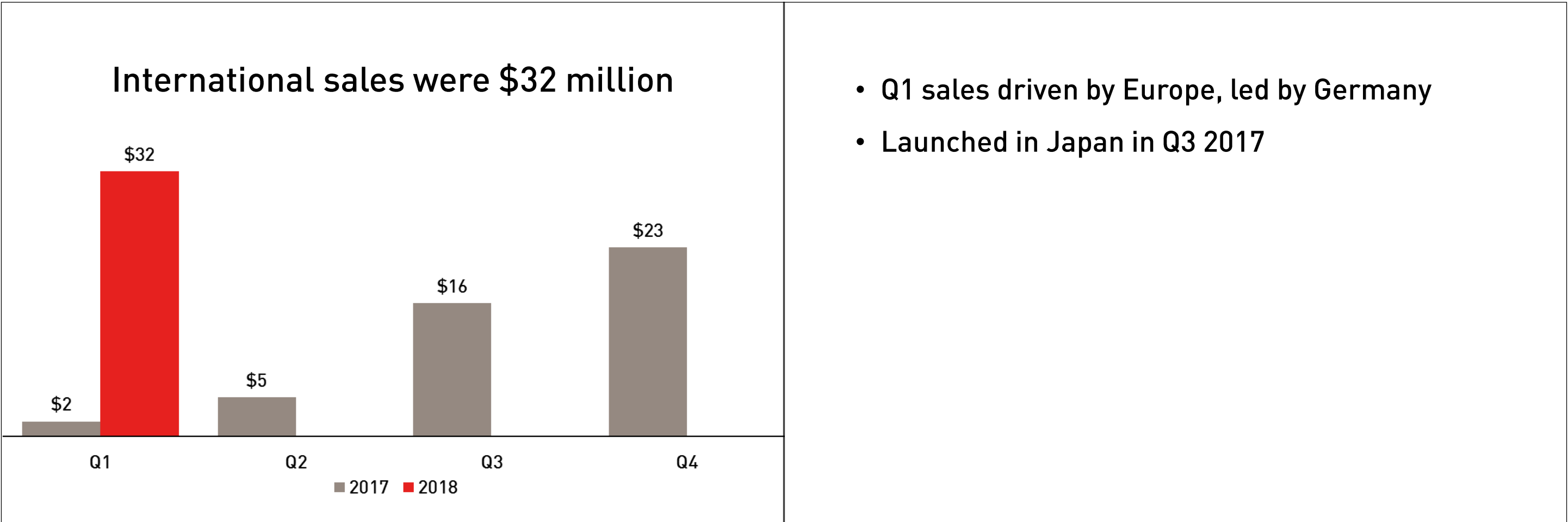


Note: Numbers may not add due to rounding.

Q1 2018 OLUMIANT SALES WERE \$32 MILLION



Millions

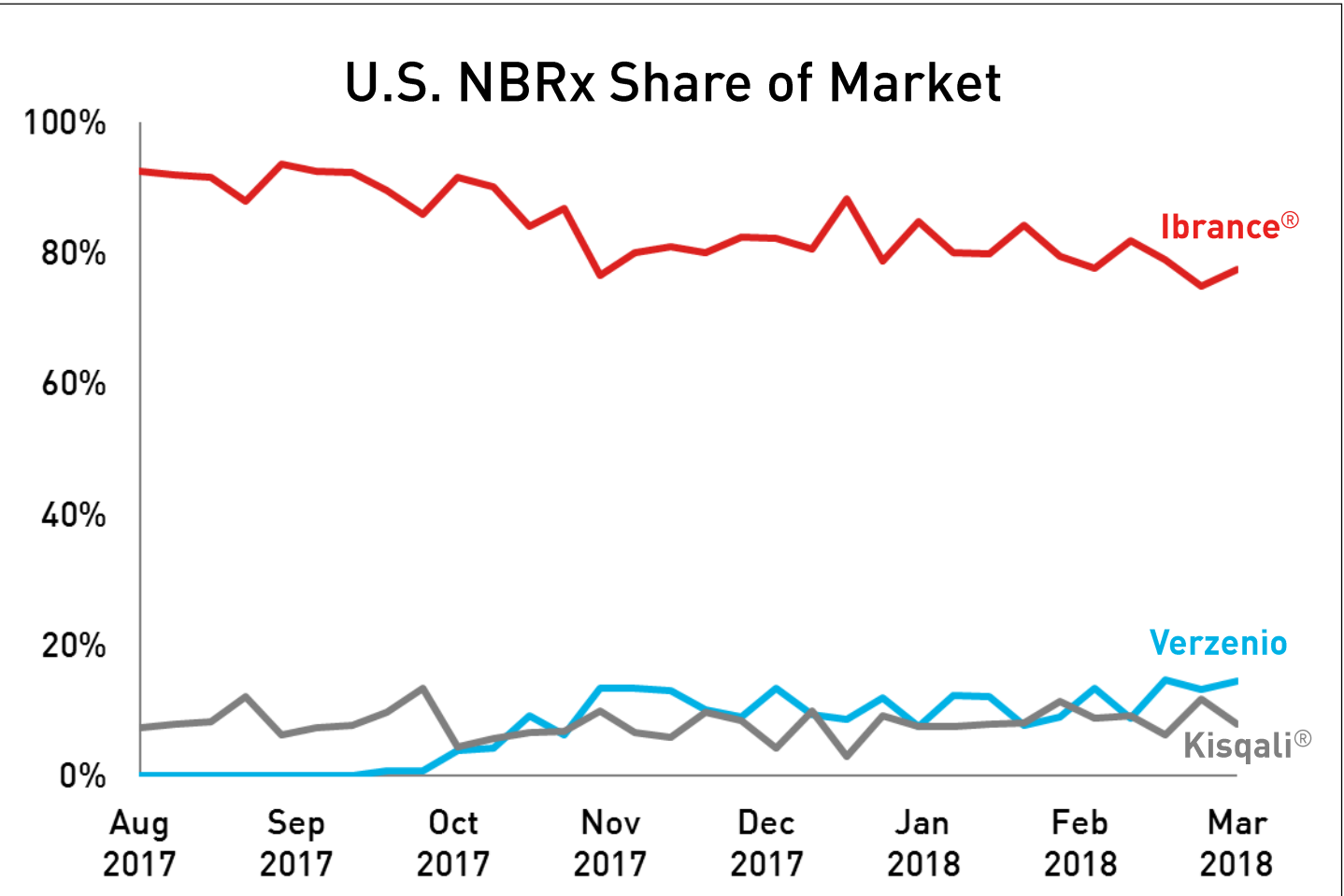
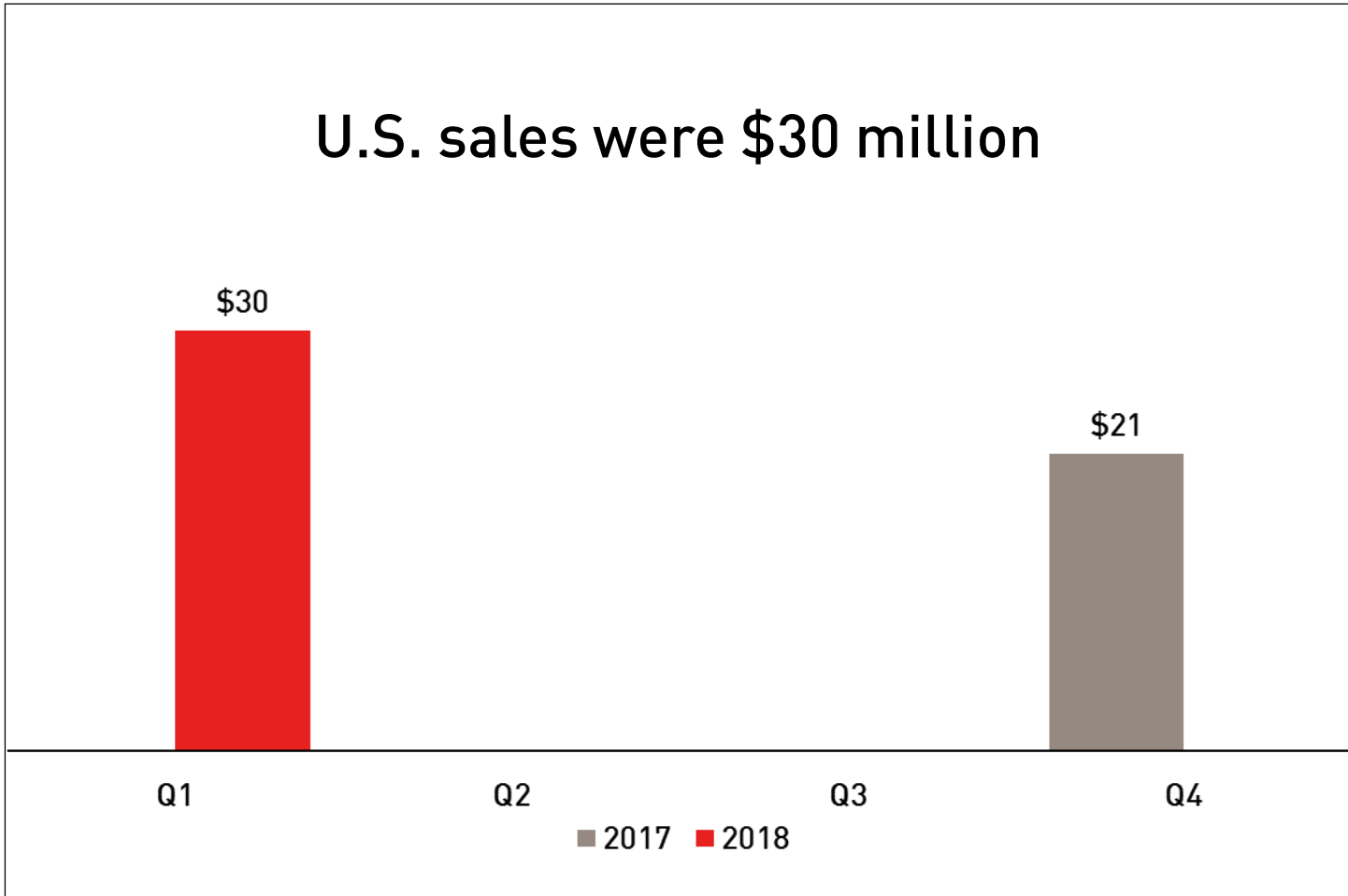


Note: Numbers may not add due to rounding.

Q1 2018 VERZENIO SALES WERE \$30 MILLION



Millions



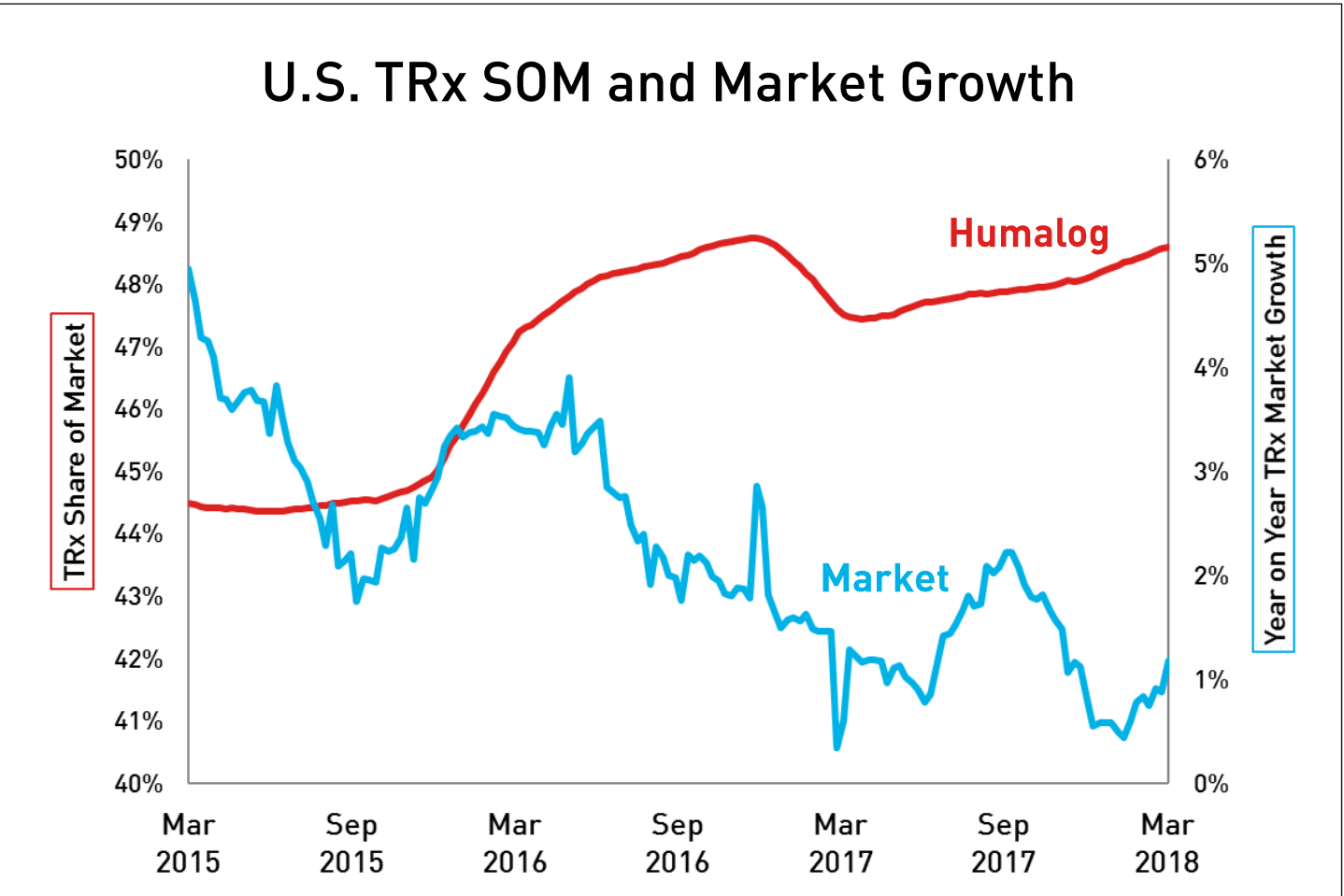
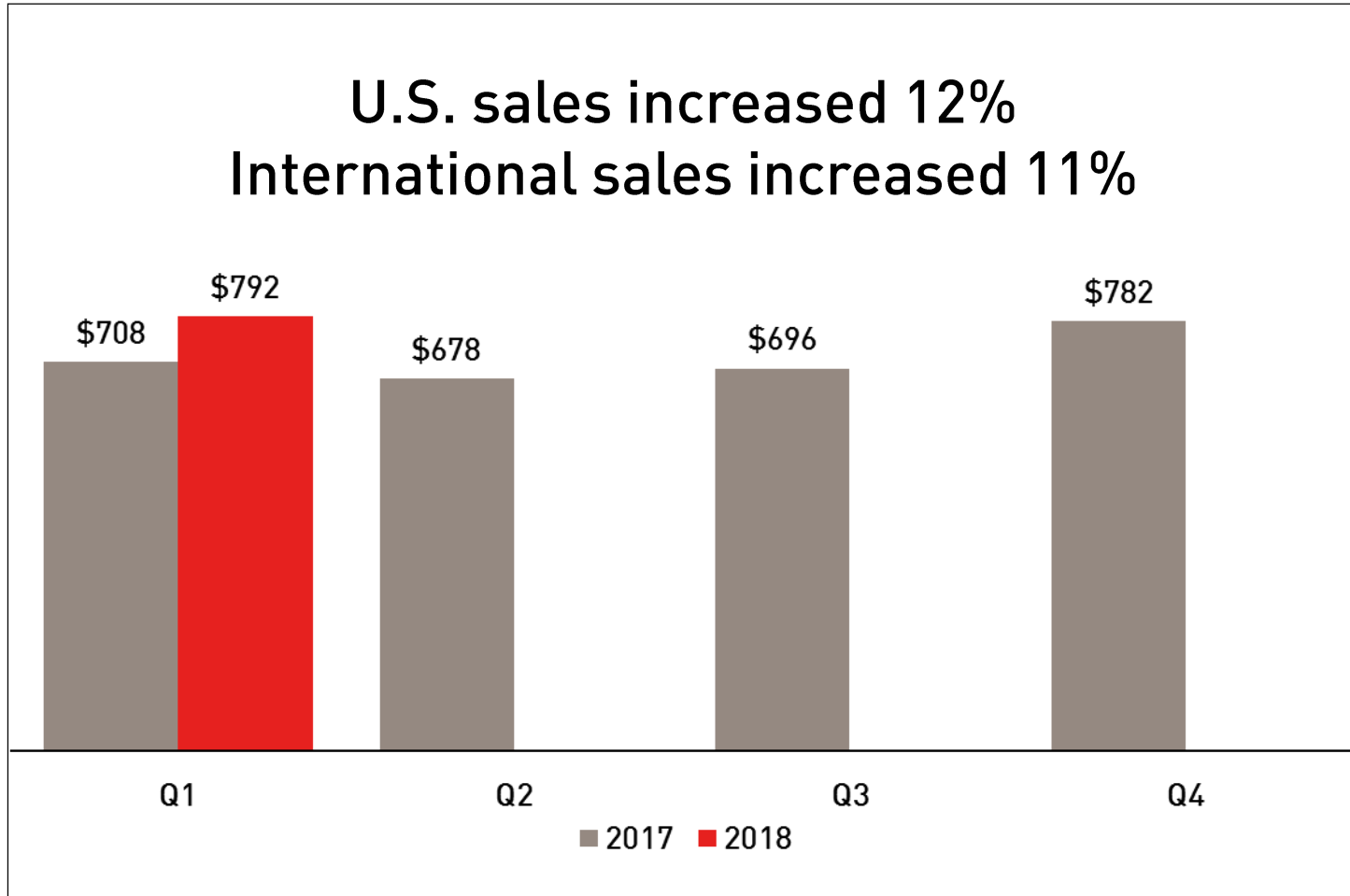
Note: Numbers may not add due to rounding.

Source: IQVIA NPA TRx 3MMA, weekly data March 30, 2018

Q1 2018 HUMALOG® SALES INCREASED 12%



Millions



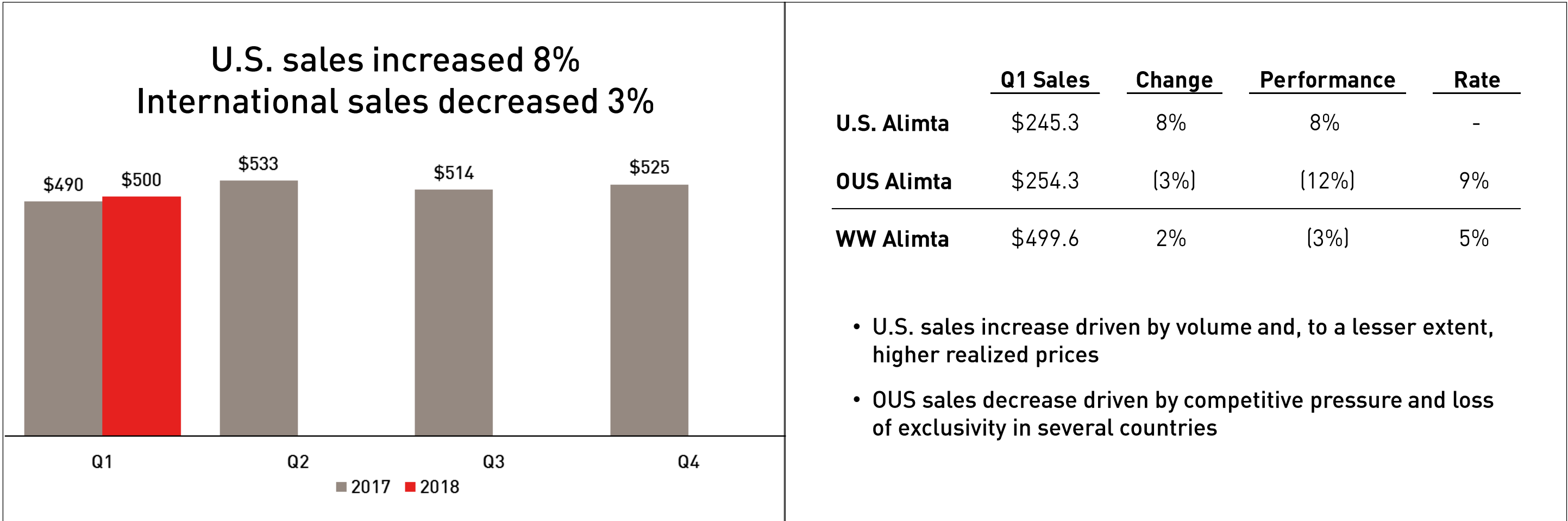
Note: Numbers may not add due to rounding.

Source: IQVIA NPA TRx 3MMA, weekly data March 30, 2018

Q1 2018 ALIMTA SALES INCREASED 2%



Millions



	<u>Q1 Sales</u>	<u>Change</u>	<u>Performance</u>	<u>Rate</u>
U.S. Alimta	\$245.3	8%	8%	-
OUS Alimta	\$254.3	(3%)	(12%)	9%
WW Alimta	\$499.6	2%	(3%)	5%

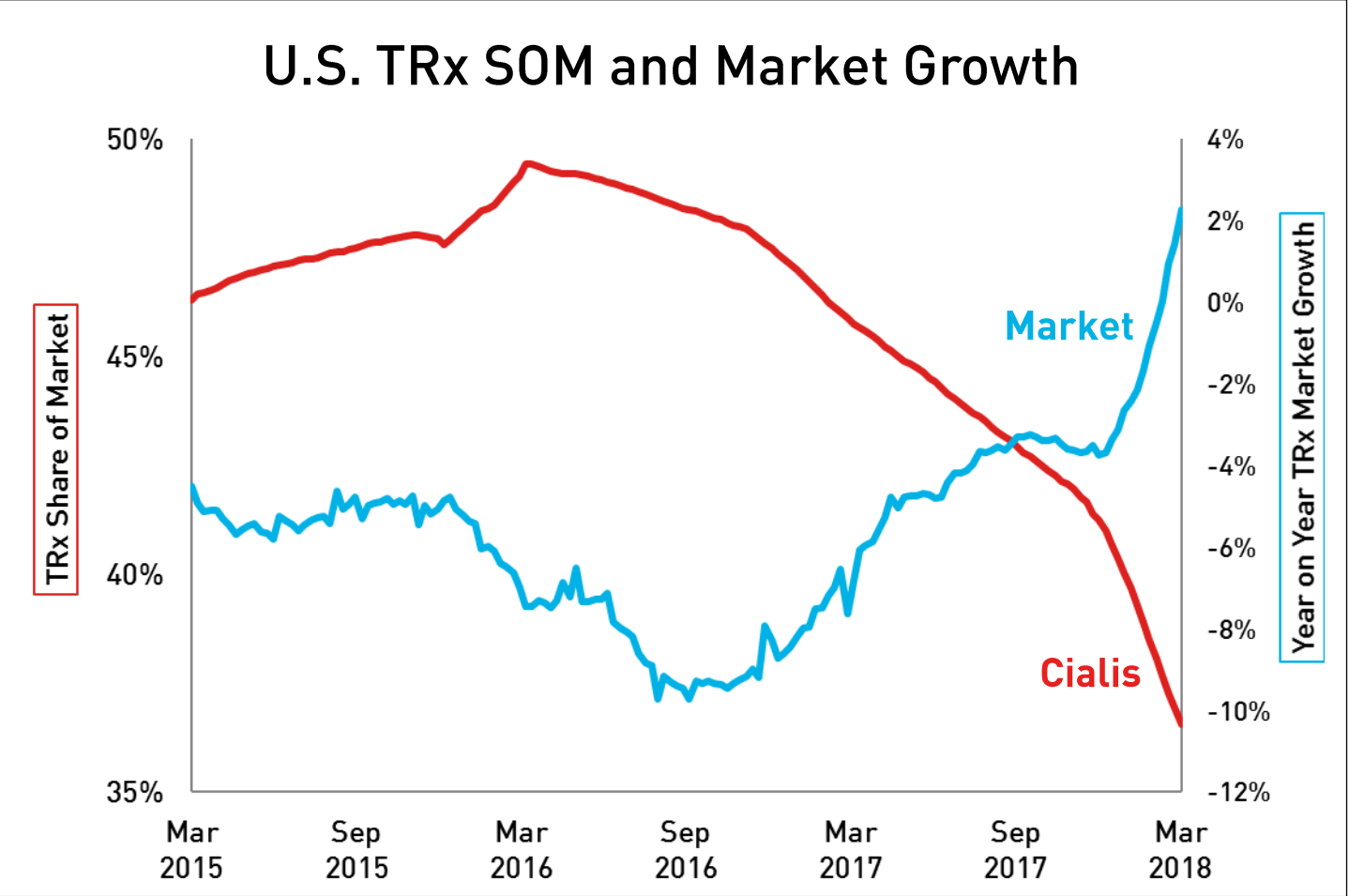
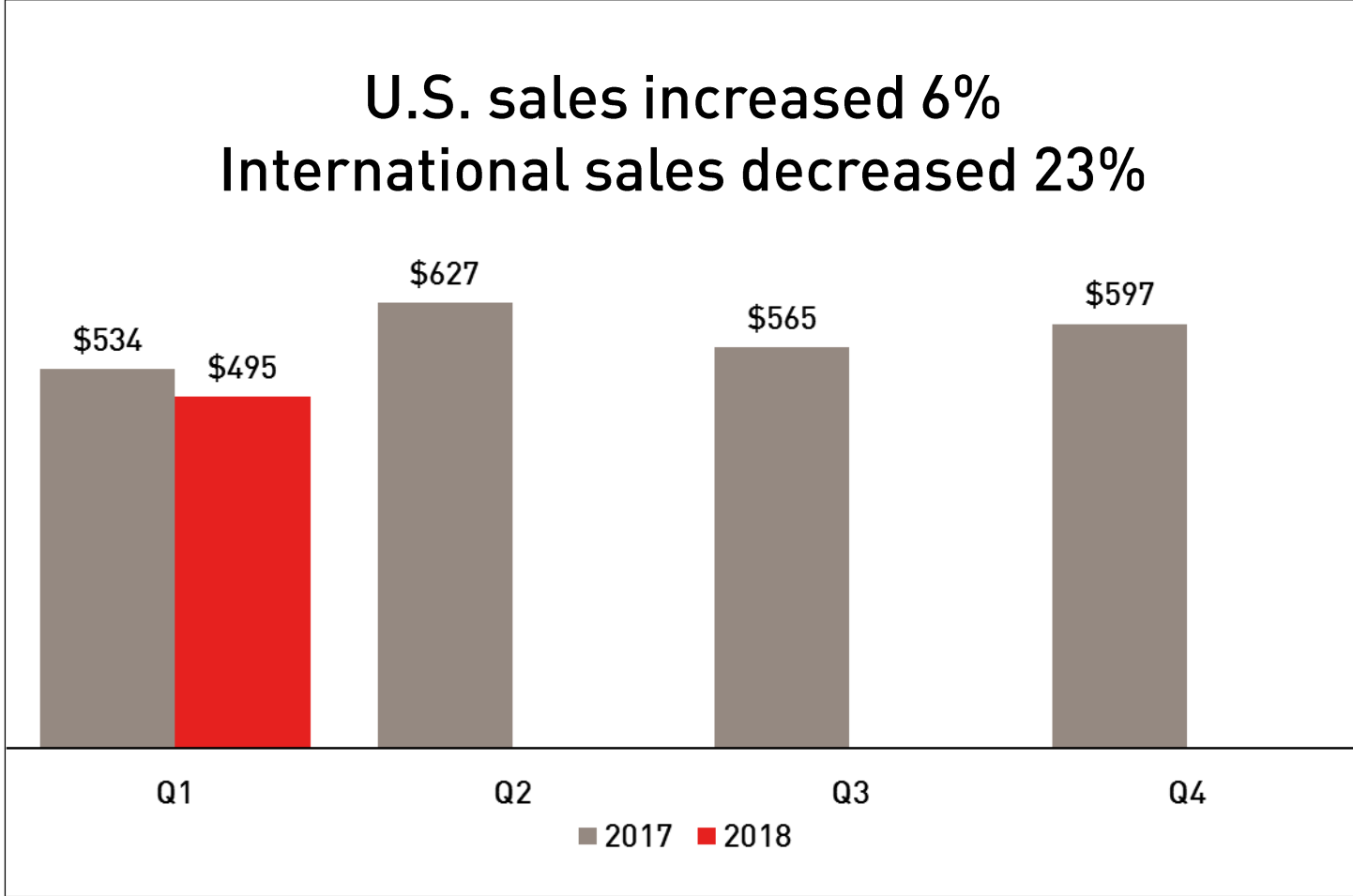
- U.S. sales increase driven by volume and, to a lesser extent, higher realized prices
- OUS sales decrease driven by competitive pressure and loss of exclusivity in several countries

Note: Numbers may not add due to rounding.

Q1 2018 CIALIS SALES DECREASED 7%



Millions



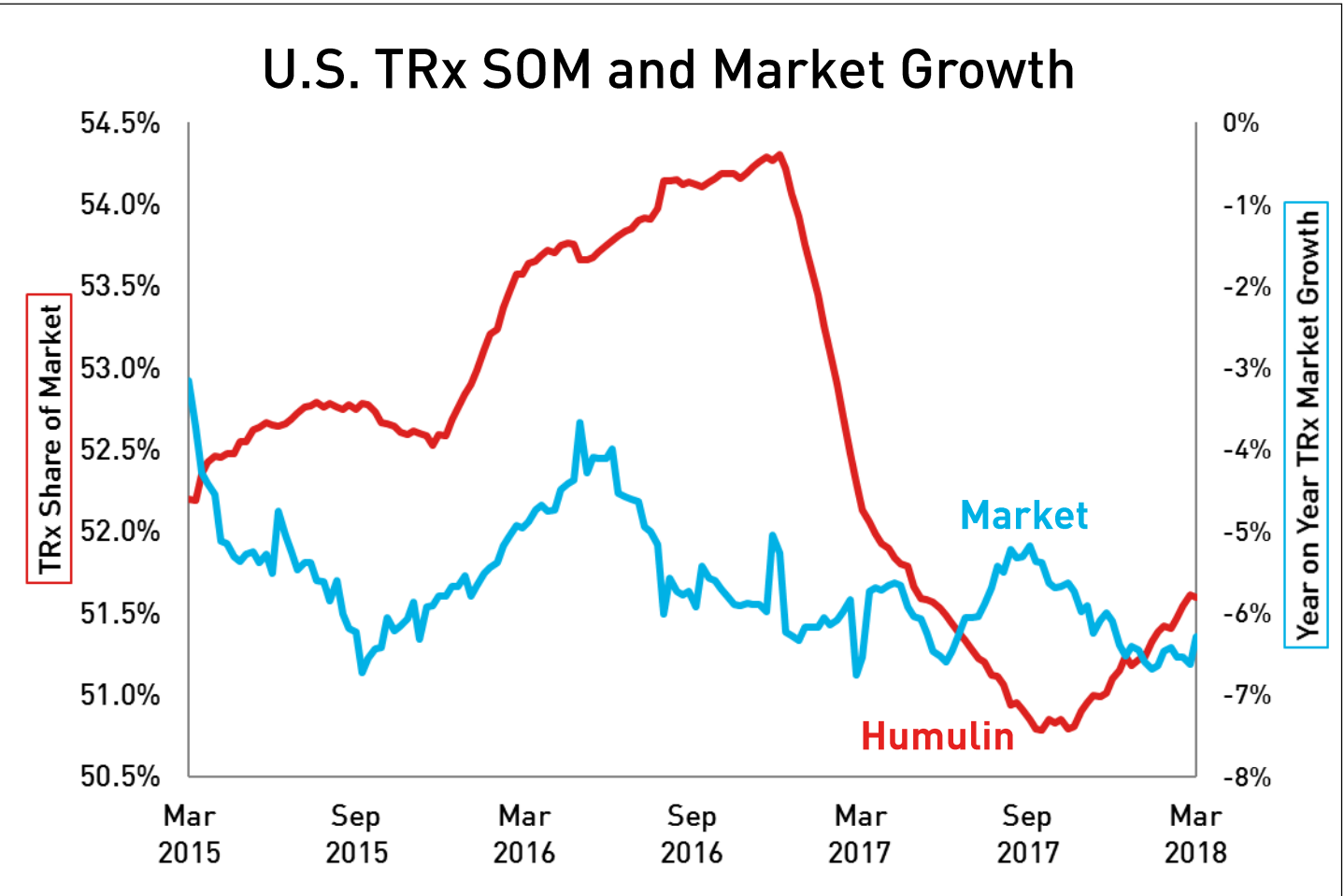
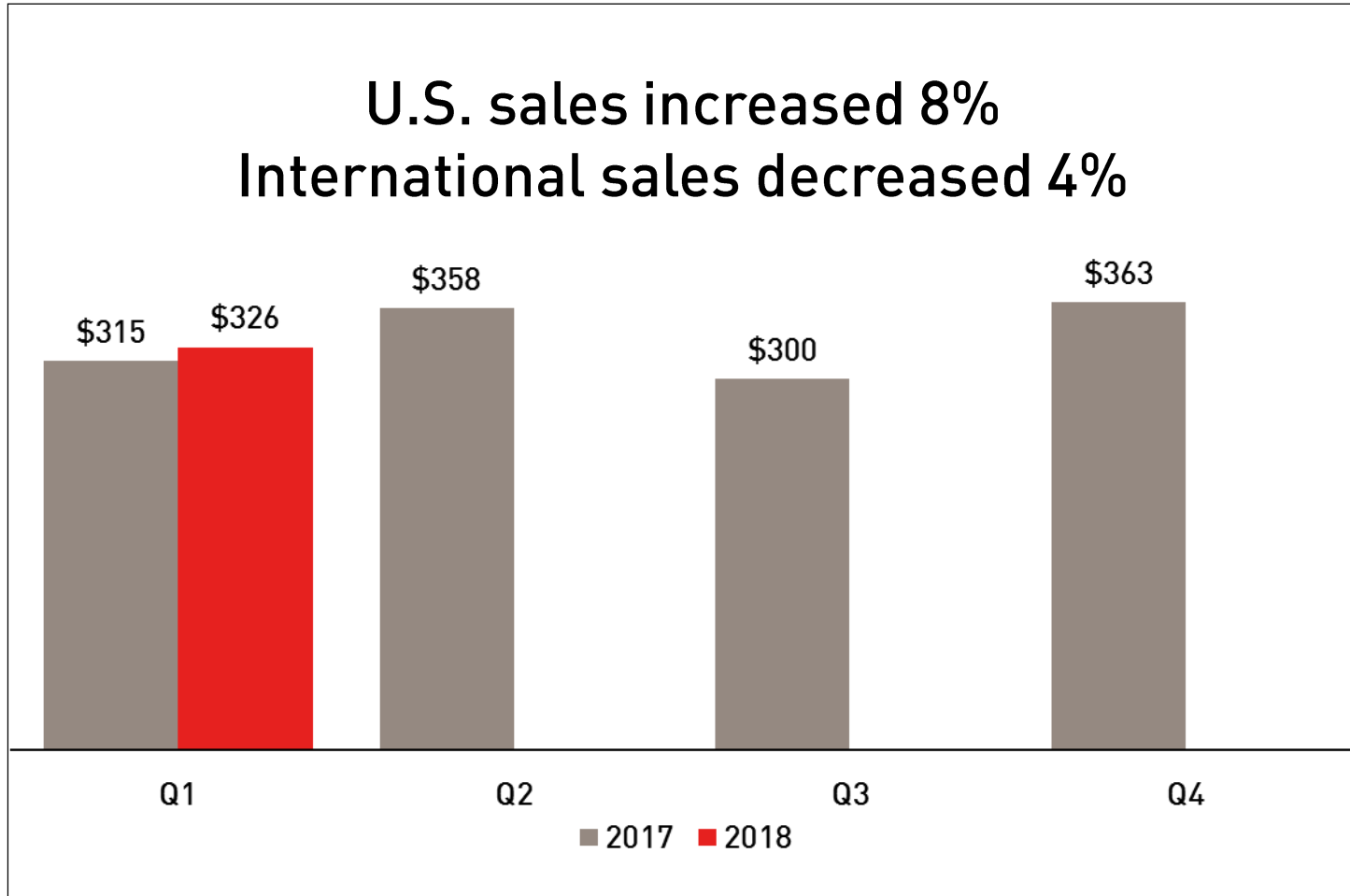
Note: Numbers may not add due to rounding.

Source: IQVIA NPA TRx 3MMA, weekly data March 30, 2018

Q1 2018 HUMULIN[®] SALES INCREASED 4%



Millions



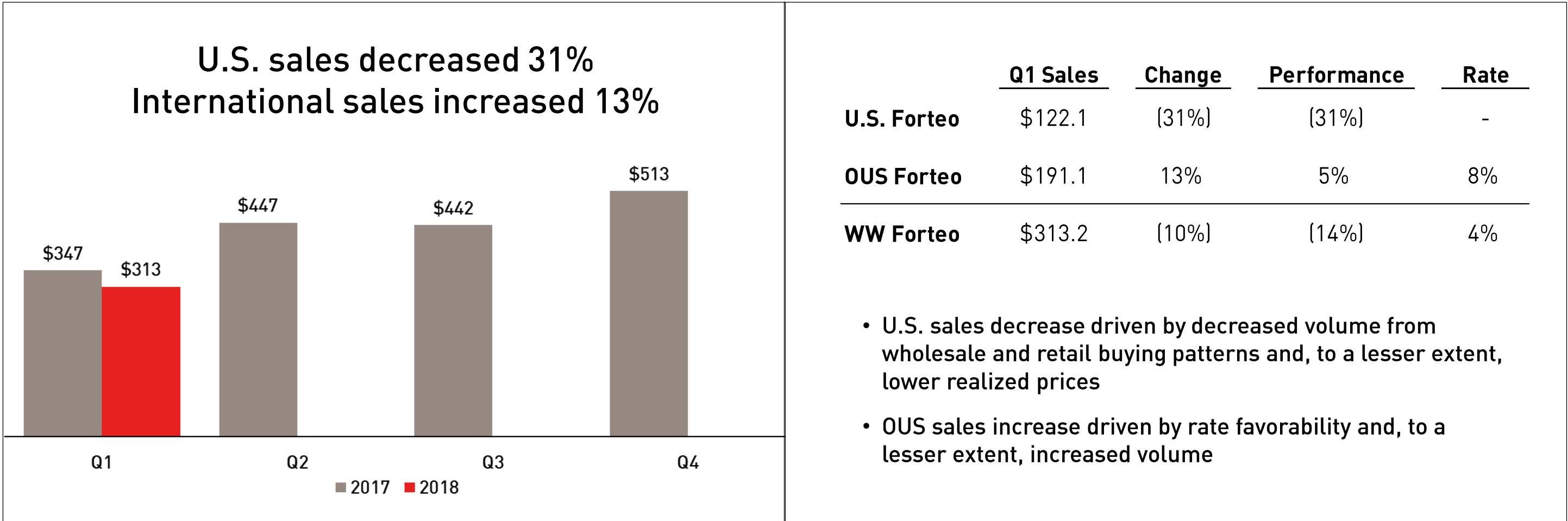
Note: Numbers may not add due to rounding.

Source: IQVIA NPA TRx 3MMA, weekly data March 30, 2018

Q1 2018 FORTEO® SALES DECREASED 10%



Millions

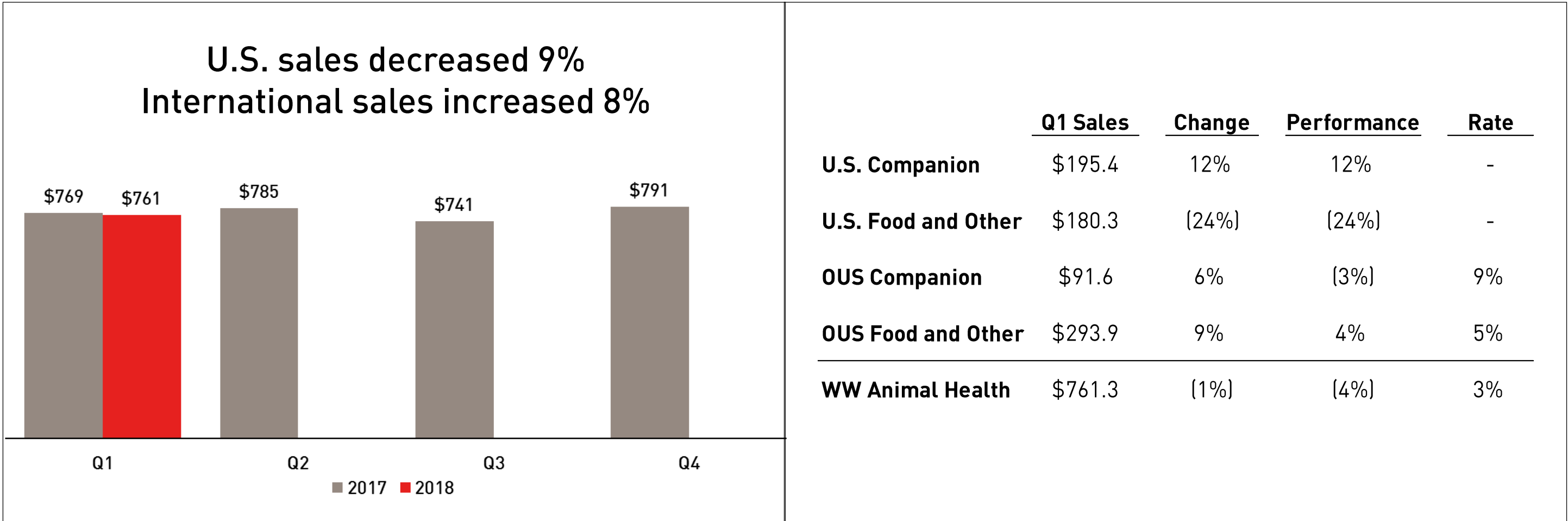


Note: Numbers may not add due to rounding.

Q1 2018 ANIMAL HEALTH SALES DECREASED 1%



Millions

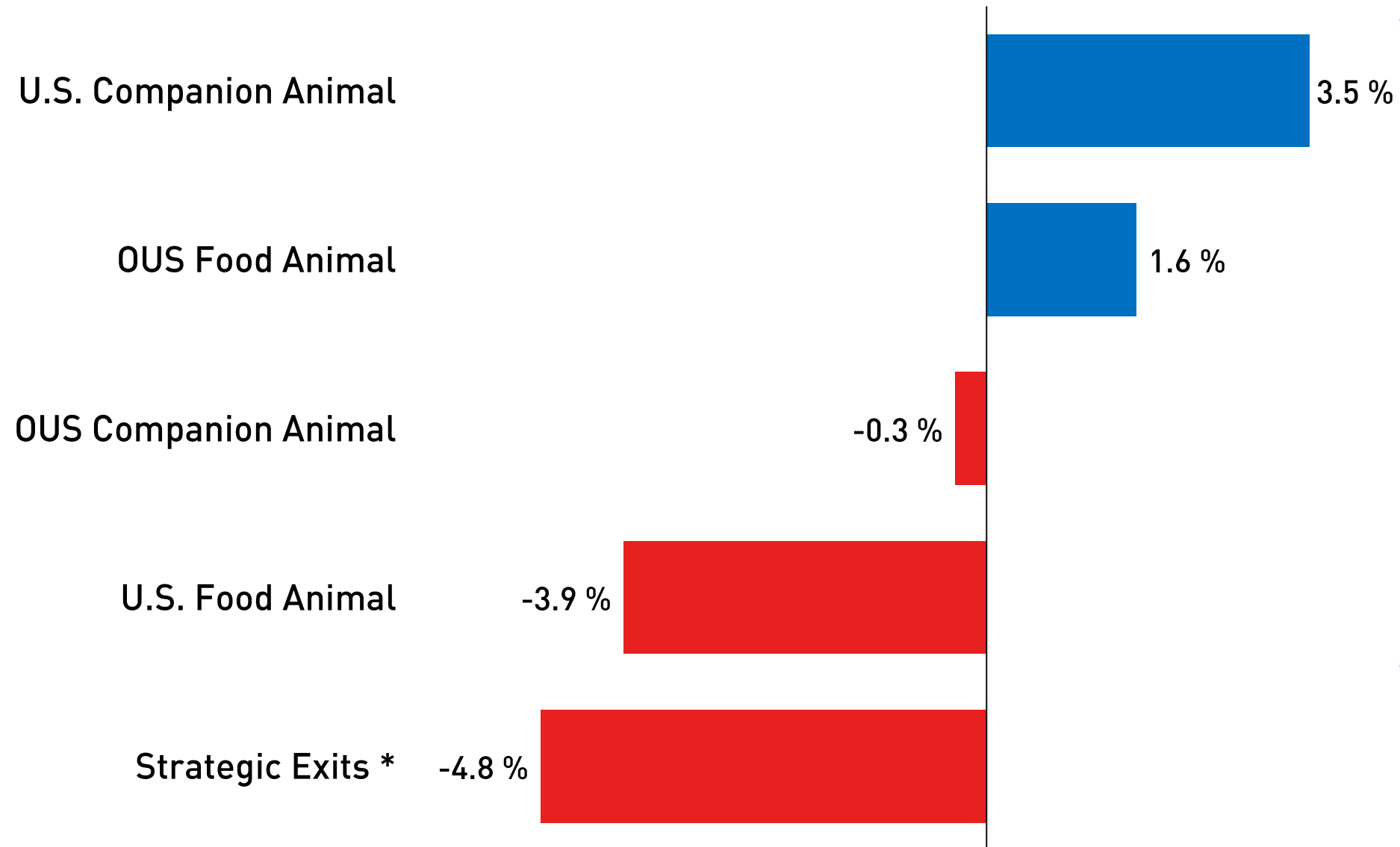


Note: Numbers may not add due to rounding.

DRIVERS OF ELANCO WW REVENUE CHANGE



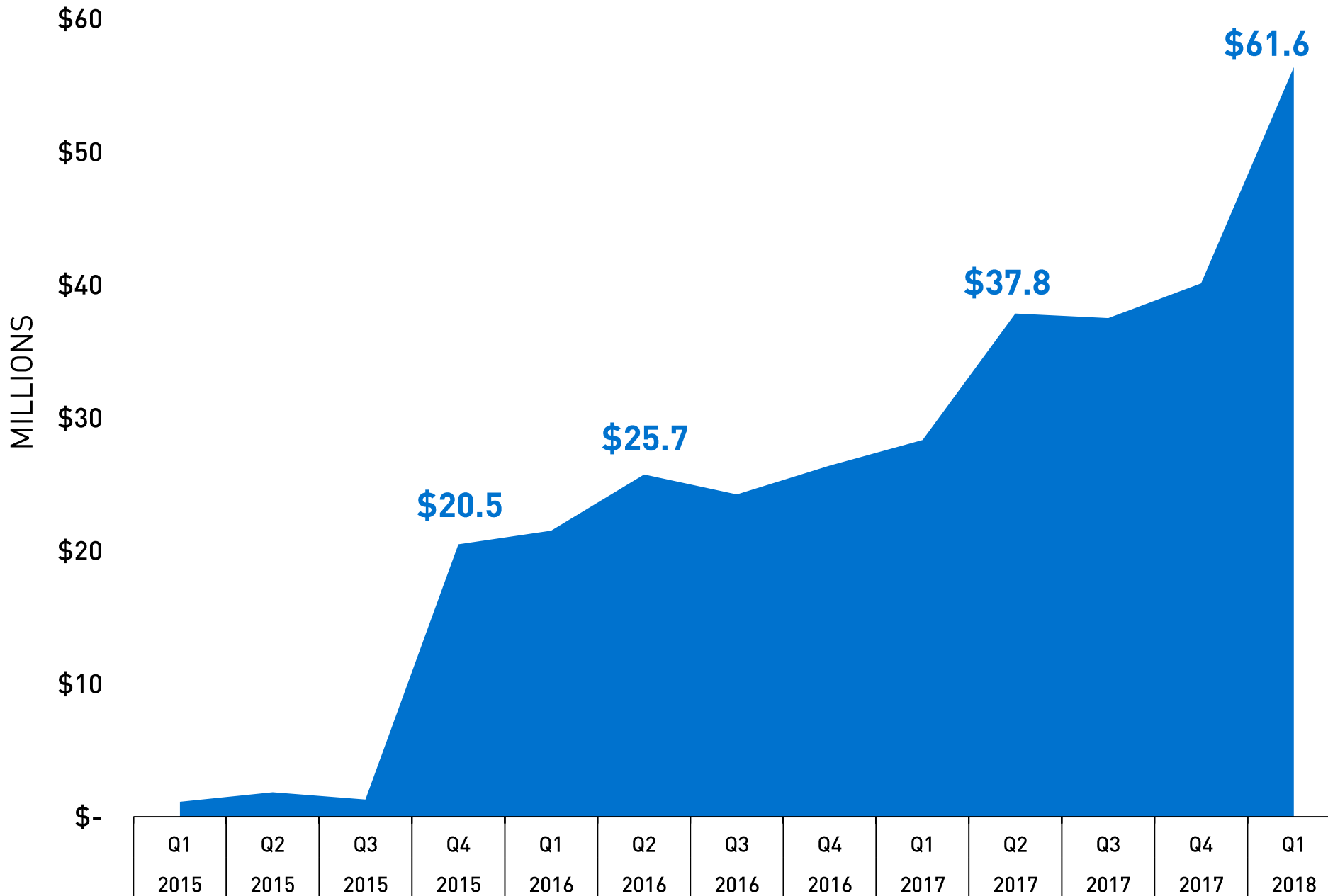
Contribution to 4% Elanco Performance Decline



**excluding Strategic Exits,
Elanco grew 1% in
performance terms**

* Strategic Exits includes Posilac, Ft. Dodge CMO, and Adequan®/Ethicon

ELANCO NEW PRODUCT LAUNCHES



New products drove **\$144M** of sales in 2017

NEW PRODUCTS INCLUDE:

COMPANION ANIMAL

- Interceptor[®] Plus
- Osumnia[®]
- Galliprant[®]
- Credelio

FOOD ANIMAL

- Imrestor[®]
- Invixa[™]
- Kavault[®]/ Integrity[®]
- Clynav[™]



Lillemor