

H1 2018 Results

Presentation, conference call and webcast for investors and analysts

26 July 2018



Forward-looking statements

In order, among other things, to utilise the 'safe harbour' provisions of the US Private Securities Litigation Reform Act 1995, we are providing the following cautionary statement: this document contains certain forward-looking statements with respect to the operations, performance and financial condition of the Group, including, among other things, statements about expected revenues, margins, earnings per share or other financial or other measures. Although we believe our expectations are based on reasonable assumptions, any forward-looking statements, by their very nature, involve risks and uncertainties and may be influenced by factors that could cause actual outcomes and results to be materially different from those predicted. The forward-looking statements reflect knowledge and information available at the date of preparation of this document and AstraZeneca undertakes no obligation to update these forward-looking statements. We identify the forward-looking statements by using the words 'anticipates', 'believes', 'expects', 'intends' and similar expressions in such statements. Important factors that could cause actual results to differ materially from those contained in forward-looking statements, certain of which are beyond our control, include, among other things: the loss or expiration of, or limitations to, patents, marketing exclusivity or trademarks, or the risk of failure to obtain and enforce patent protection; effects of patent litigation in respect of IP rights; the impact of any delays in the manufacturing, distribution and sale of any of our products; the impact of any failure by third parties to supply materials or services; the risk of failure of outsourcing; the risks associated with manufacturing biologics; the risk that R&D will not yield new products that achieve commercial success; the risk of delay to new product launches; the risk that new products do not perform as we expect; the risk that strategic alliances and acquisitions, including licensing and collaborations, will be unsuccessful; the risks from pressures resulting from generic competition; the impact of competition, price controls and price reductions; the risks associated with developing our business in emerging markets; the risk of illegal trade in our products; the difficulties of obtaining and maintaining regulatory approvals for products; the risk that regulatory approval processes for biosimilars could have an adverse effect on future commercial prospects; the risk of failure to successfully implement planned cost reduction measures through productivity initiatives and restructuring programmes; the risk of failure of critical processes affecting business continuity; economic, regulatory and political pressures to limit or reduce the cost of our products; failure to achieve strategic priorities or to meet targets or expectations; the risk of substantial adverse litigation/government investigation claims and insufficient insurance coverage; the risk of substantial product liability claims; the risk of failure to adhere to applicable laws, rules and regulations; the risk of failure to adhere to applicable laws, rules and regulations relating to anti-competitive behaviour; the impact of increasing implementation and enforcement of more stringent anti-bribery and anti-corruption legislation; taxation risks; exchange rate fluctuations; the risk of an adverse impact of a sustained economic downturn; political and socio-economic conditions; the risk of environmental liabilities; the risk of occupational health and safety liabilities; the risk associated with pensions liabilities; the impact of failing to attract and retain key personnel and to successfully engage with our employees; the risk of misuse of social media platforms and new technology; and the risk of failure of information technology and cybercrime. Nothing in this presentation / webcast should be construed as a profit forecast.



Presenters



Pascal Soriot
Executive Director and
Chief Executive Officer



Dave Fredrickson
Executive Vice President,
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Mark Mallon
Executive Vice President,
Global Products & Portfolio
Strategy, Global Medical
Affairs, Corporate Affairs



Marc Dunoyer
Executive Director and
Chief Financial Officer



Sean Bohen
Executive Vice President,
Global Medicines Development
and Chief Medical Officer



Agenda



Overview



Oncology



CVRM, Respiratory, EMs



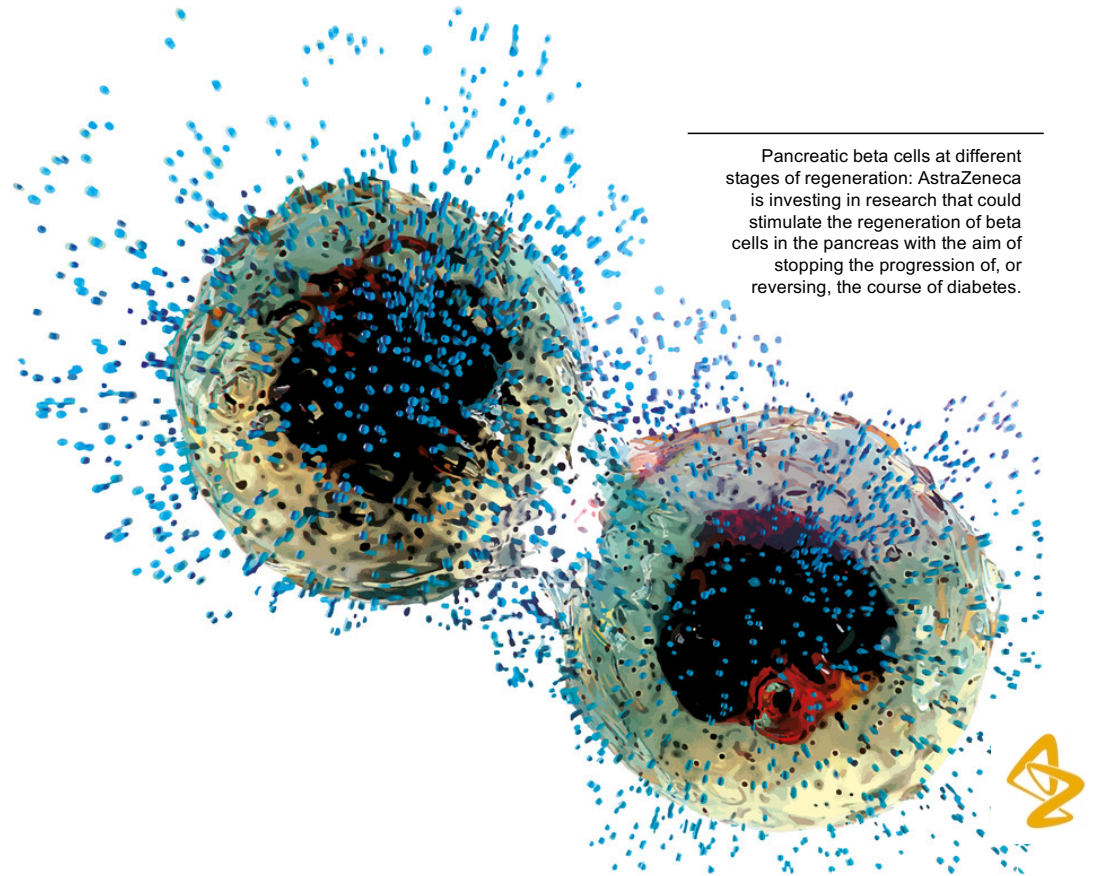
Finance



Pipeline and news flow



Closing and Q&A



Pancreatic beta cells at different stages of regeneration: AstraZeneca is investing in research that could stimulate the regeneration of beta cells in the pancreas with the aim of stopping the progression of, or reversing, the course of diabetes.

Strategic business focus is paying off

The main therapy areas accelerated growth

Product Sales
growth (CER¹)

Q2 2018

H1 2018

Oncology,
New CVRM²,
Respiratory

+19% 

+14% 

Other

-32%

-25%

1. Constant exchange rates.

2. New Cardiovascular, Renal and Metabolism incorporating *Brilinta* and Diabetes.



Launches continue to support 2018 return to growth

Portfolio transformation of AstraZeneca is nearing completion

Business & financials

Product Sales declined by 2% and only by 1% in the quarter

- Strong performance of new medicines¹ (+69%) and China
- Offset by divestments (~2%) and EU/JP *Crestor* generics

Total Revenue declined by 5%

New medicines¹ continued forward: >\$1bn additional sales vs. H1 2017

- Oncology: +37%; continued strong sales of *Lynparza*, *Tagrisso* and *Imfinzi*
- New CVRM: +9%; *Brilinta* (+18%); *Farxiga* (+36%)
- Respiratory: stabilised; *Symbicort* competition; *Pulmicort* supply normalised; *Fasenra* continued strong launch
- Emerging Markets: +10%
 - China: +24%; another very strong quarter (+26%)

Core EPS \$1.17 and FY 2018 guidance reiterated

1. *Lynparza*, *Tagrisso*, *Imfinzi*, *Calquence*, *Brilinta*, *Farxiga*, *Bevespi* and *Fasenra*. Absolute growth at CER and compared to H1 2017. Absolute values at actual exchange rates; changes at CER and for H1 2018, unless otherwise stated. Guidance at CER.



The pipeline continued to deliver

Late-stage pipeline quarterly highlights

Pipeline news

Oncology	• <i>Lynparza</i>	breast cancer ovarian cancer 1L	Approval (JP) Met primary endpoint
	• <i>Tagrisso</i>	lung cancer 1L	Approval (EU)
	• <i>Imfinzi</i>	unresectable, Stage III NSCLC ¹	Approval (JP) Met primary OS endpoint
	• selumetinib	thyroid cancer	Did not meet primary endpoint
Cardiovascular, Renal and Metabolism	• <i>Forxiga</i>	type-1 diabetes	Regulatory submission (JP)
	• combo w/ <i>Onglyza</i> and metformin	type-2 diabetes	Regulatory submission acceptance (EU)
	• <i>Bydureon</i>	type-2 diabetes CVOT ²	Regulatory submission acceptance (US)
	• <i>Bydureon BCise</i>	type-2 diabetes; new device	Positive CHMP opinion (EU)
	• <i>Lokelma</i>	hyperkalaemia	Approval (US)
Respiratory	• <i>Fasenra</i>	COPD ³	Did not meet primary endpoints
Other	• <i>Ianabecestat</i>	Alzheimer's disease	Termination of Phase III programme

1. Non-small cell lung cancer.

2. Cardiovascular outcomes trial.

3. Chronic obstructive pulmonary disease.

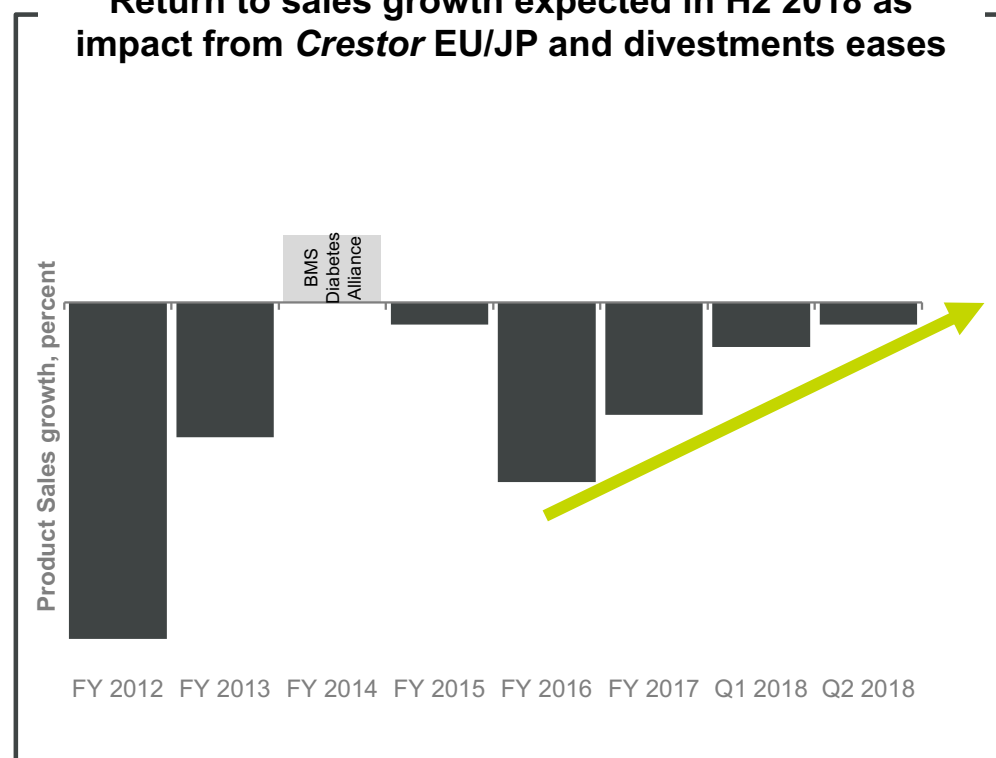
Status since the last results announcement on 18 May 2018.



2018: return to sales growth on track

The sales momentum continued to improve

Return to sales growth expected in H2 2018 as impact from *Crestor* EU/JP and divestments eases



Medicines impacting Product Sales in 2018

Lynparza

ongoing launch of tablet in ovarian and breast cancer

Tagrisso

ongoing launch in 1st-line lung cancer

Imfinzi

ongoing launch in unresect., sIII lung cancer

Brilinta

continued global growth

Farxiga

continued global growth and the DECLARE trial

Crestor

loss of exclusivity (EU, JP)

Fasenra

ongoing launch in severe, uncontrolled asthma

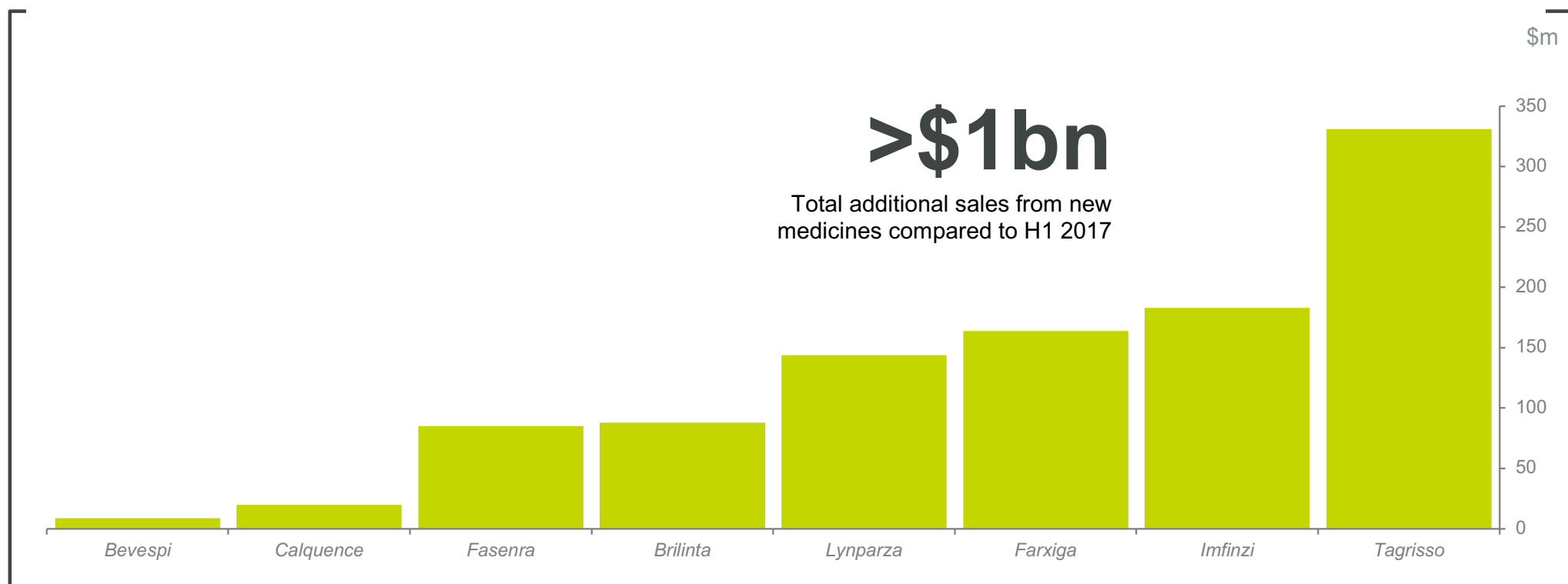
2018: low single-digit growth in Product Sales

Change (Product Sales growth) and FY 2018 guidance at CER.



Product Sales: new medicines continued forward

>\$1bn in additional sales and growth of 69% in H1 2018



Absolute values at CER.



Product Sales: Oncology and China performed strongly

Global performance impacted by *Crestor* EU/JP and divestments

	Q2 2018 \$m	% change	% Product Sales	H1 2018 \$m	% change	% Product Sales
Product Sales	5,030	(1)	100	10,015	(2)	100
 Oncology	1,434	40	29	2,664	37	27
 New CVRM	974	9	19	1,874	9	19
 Respiratory	1,226	7	24	2,407	-	24
Other	1,396	(32)	28	3,070	(25)	31
 Emerging Markets	1,659	12	33	3,424	10	34
-of which China	868	26	17	1,893	24	19

Product Sales values at actual exchange rates; changes at CER and for H1 2018, unless otherwise stated.



Agenda



Overview



Oncology



CVRM, Respiratory, EMs



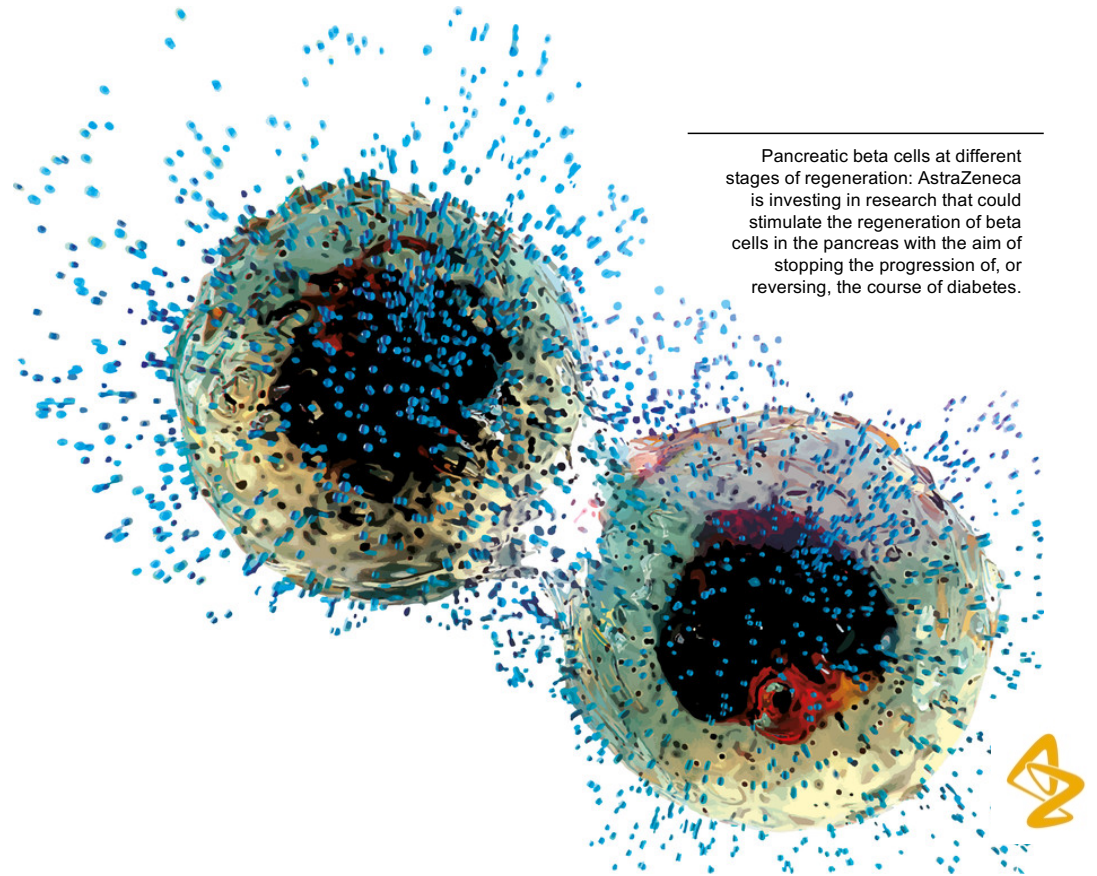
Finance



Pipeline and news flow



Closing and Q&A

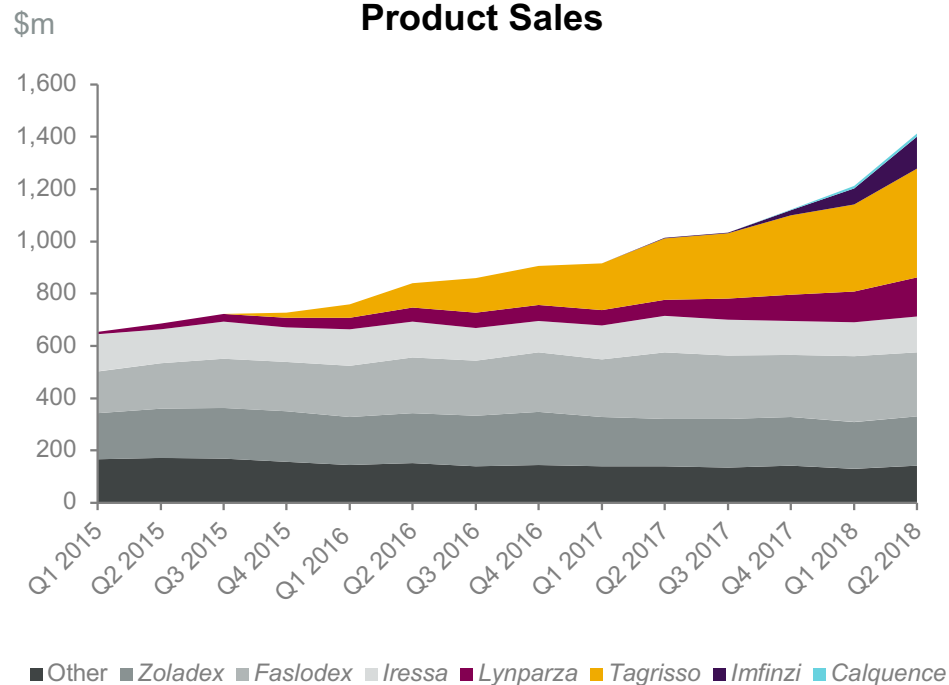


Pancreatic beta cells at different stages of regeneration: AstraZeneca is investing in research that could stimulate the regeneration of beta cells in the pancreas with the aim of stopping the progression of, or reversing, the course of diabetes.

Oncology

New medicines continued to drive strong performance

Total Oncology
Product Sales



New medicines *Lynparza*, *Tagrisso*,
Imfinzi and *Calquence* added \$0.7bn

- Oncology +37%; now 27% of total Product Sales
- New medicines contributed \$0.7bn in additional sales vs. H1 2017
 - **Lynparza**: accelerated growth globally; promising launch in Japan
 - **Tagrisso**: sustained very high growth; increasing use in 2nd line; encouraging start in the 1st-line setting
 - **Imfinzi**: quarterly sales ~doubled in lung cancer
 - **Calquence**: launch progressed solidly with increased use in BTKi-naïve patients

Absolute values and change at CER and for H1 2018, unless otherwise stated.



Four quarters of strong growth: +147% in Q2

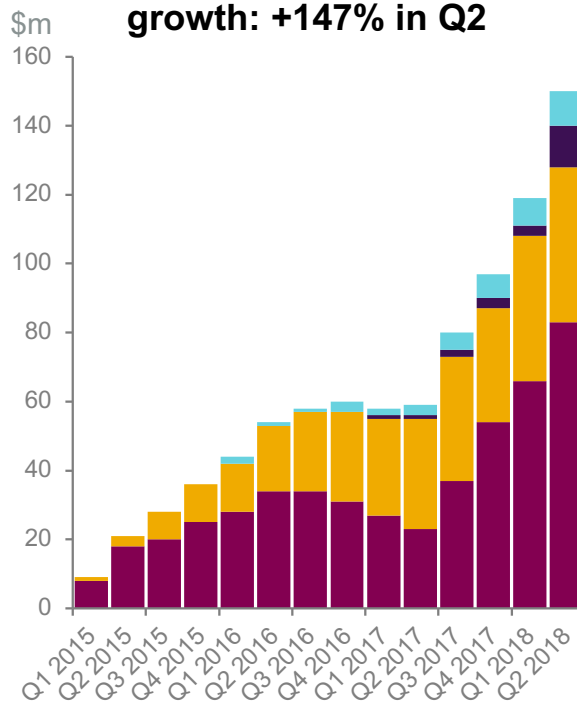


Chart legend: **US** **Europe** **Established RoW** **Emerging Markets**.
Absolute values at actual exchange rates; changes at CER and for H1 2018, unless otherwise stated.

Leading PARP inhibitor approved in >50 countries

- **US +198%**
Tablet formulation, broad label in ovarian cancer and launch in breast cancer accelerated growth
- **Europe +36%**
Increased testing rates, duration and early adoption of tablet and broad label in ovarian cancer
- **Established RoW**
Successful launch in Japan (\$10m); breast cancer approved

Upcoming key milestones

- 1st-line ovarian cancer (*BRCAm*) data presentation in H2 2018; regulatory submission soon
- China first regulatory decision expected in H2 2018 in ovarian cancer
- EU breast cancer regulatory decision expected in H1 2019

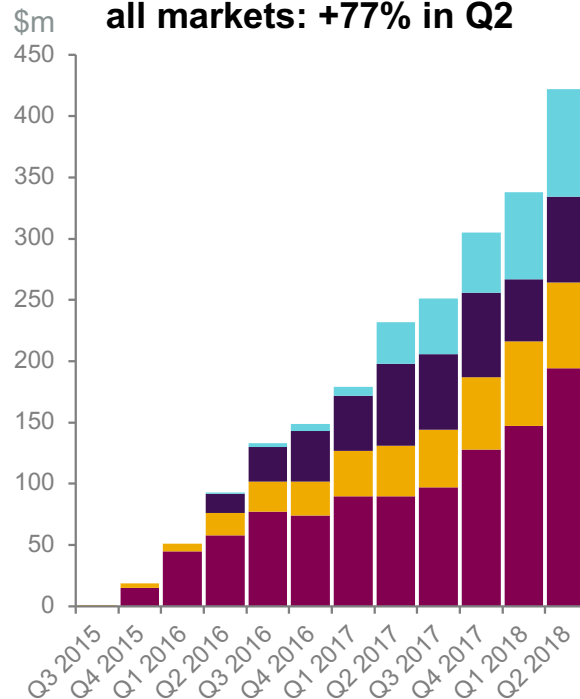


Lung cancer: *Tagrisso*



Strong 2nd-line business; step change from 1st-line launches

Strong performance in all markets: +77% in Q2



Approved in >75 countries worldwide

- **US +89%**
Continued momentum in 2nd line with a boost from 1st-line launch
- **Europe +63%**
Continued 2nd line momentum; early 1st-line launches
- **Japan +11%**
Sequential quarterly growth back following intense 2nd-line focus
- **Emerging Markets**
Continued strong uptake in China

1st-line launches will widen patient benefits

- Unprecedented 1st-line progression-free survival data
- Approved in Brazil, US, EU, Russia, Australia, Canada, Egypt
- Reimbursement underway in the EU; launched in France, Germany
- JP regulatory decision expected in H2 2018 with subsequent launch
- China regulatory decision expected from next year

Chart legend: **US** **Europe** **Established RoW** **Emerging Markets**.
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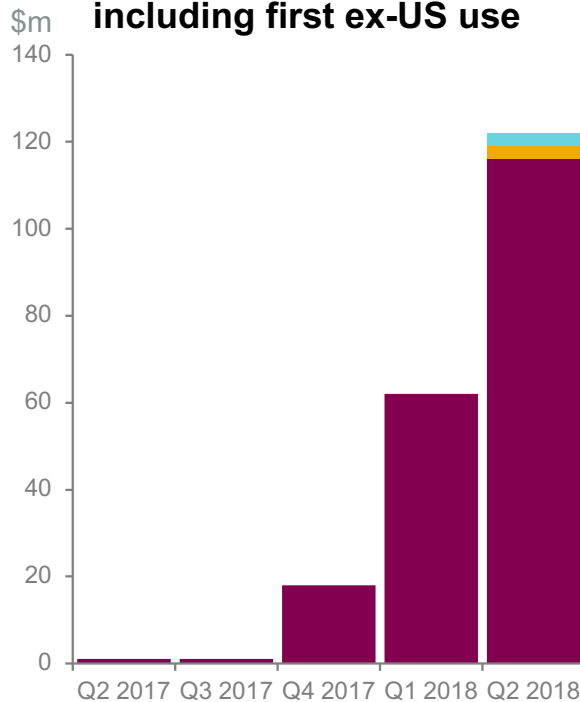


Lung cancer: *Imfinzi*

Continued fast uptake in unresectable, Stage III NSCLC (PACIFIC)



Q2 Product Sales: \$122m, including first ex-US use



PACIFIC launch gaining global momentum

- **Product Sales ~doubled to \$122m in Q2; total \$184m in H1**
Lung cancer the majority of sales; very limited use in bladder cancer
- **Additional approvals obtained**
Japan, Canada, Switzerland, India, Brazil
- **First non-US sales in Q2 2018**
- **~40 more countries expected to approve PACIFIC regimen in H2**

Increasingly more US patients are treated with *Imfinzi*

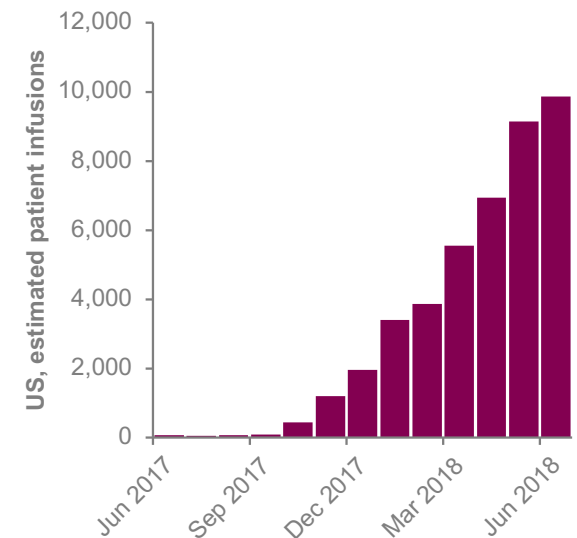
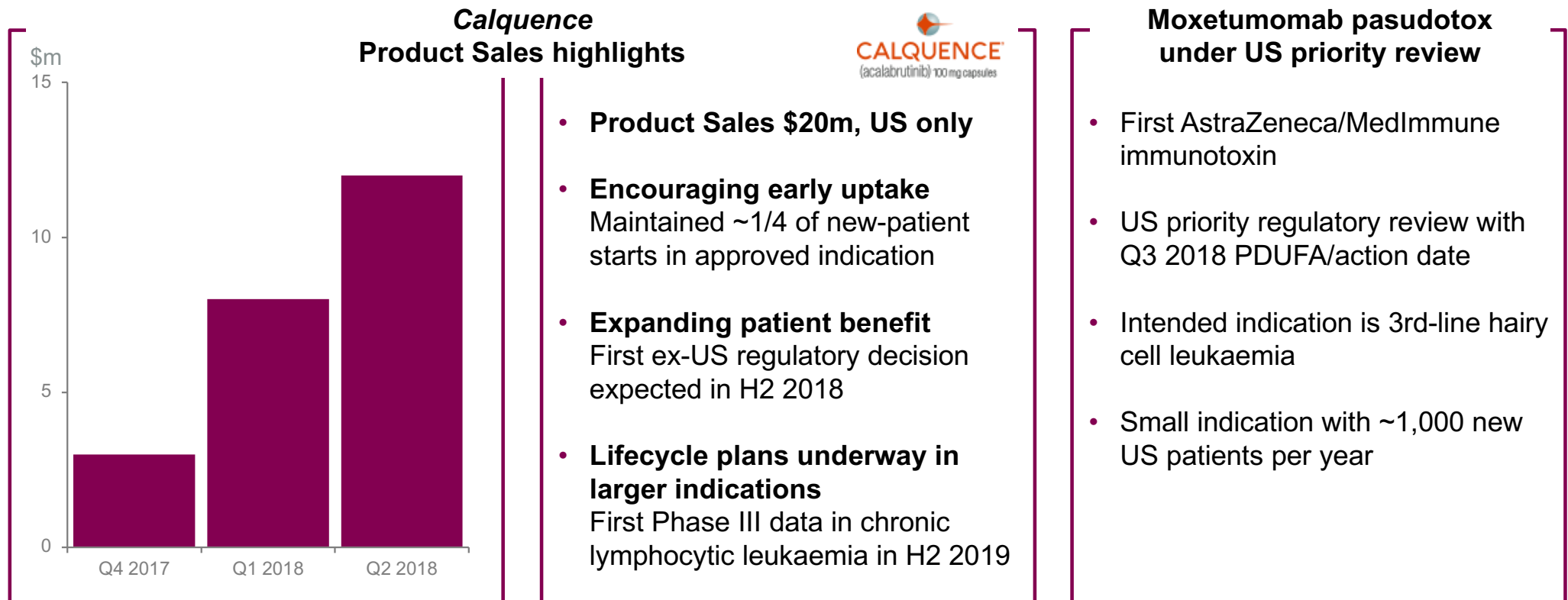


Chart legend: **US** **Europe** **Established RoW** **Emerging Markets**.
Absolute values at actual exchange rates.



Haematology: *Calquence* and moxetumomab

Emerging franchise; initially in smaller indications



Absolute values at actual exchange rates.



New CVRM

Brilinta and Farxiga delivered strong results

Brilinta Product Sales and growth

Brilinta +18%: Continued double-digit growth across all major regions

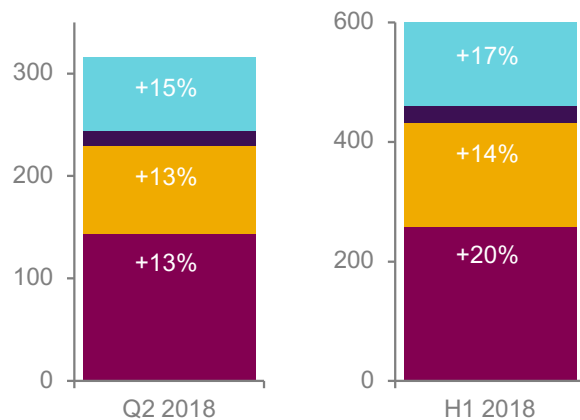


Chart legend: **US** **Europe** **Established RoW** **Emerging Markets**.
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Diabetes Product Sales and growth

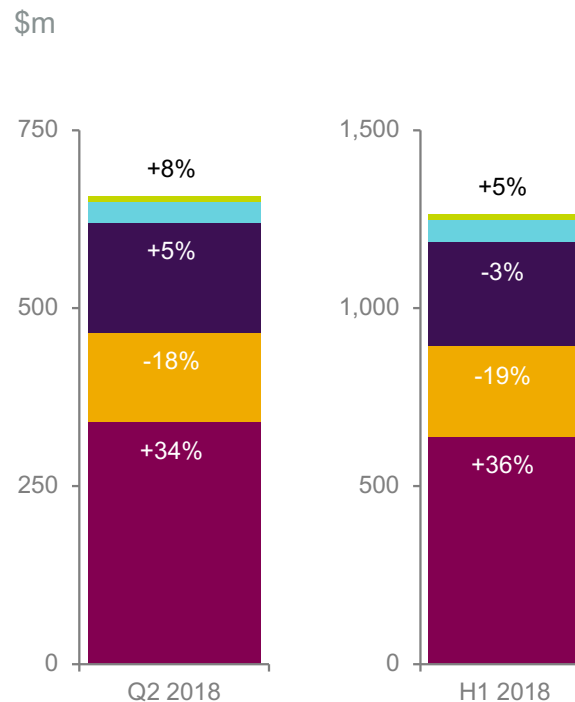


Chart legend: **Farxiga** **Onglyza** **Bydureon** **Byetta** **Other**.

Source: IQVIA. Farxiga: includes fixed-dose combinations.

Farxiga +36%

- US (+29%); increased market share from contract gains; overall market growth slowing
- Ex-US (58% of total; increasing) Strong volume-driven growth continued, e.g. Europe (+28%), Emerging Markets (+59%)

Bydureon -3%, but +5% in Q2

- Strong launch of new *BCise* device
- Volumes starting to offset price



Respiratory

Improving performance; *Fasenra* and *Pulmicort* offsetting *Symbicort*

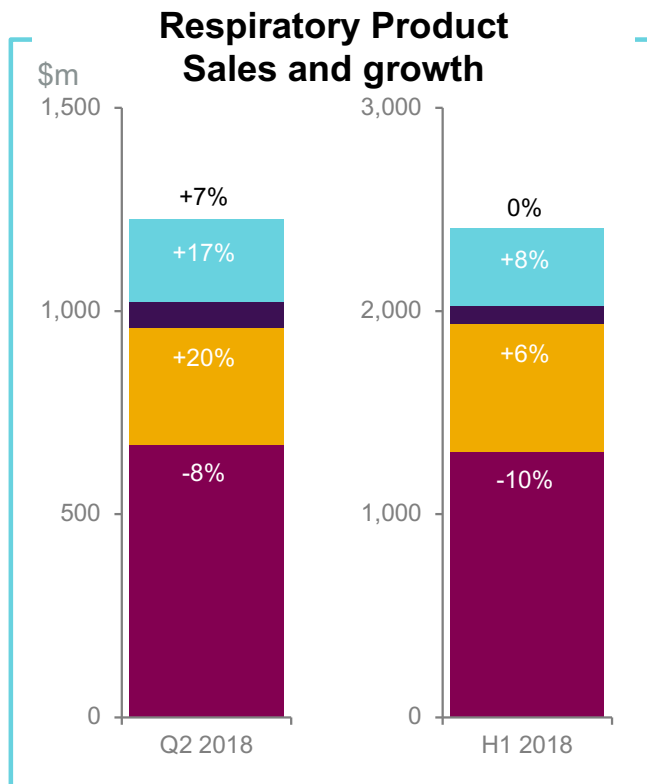


Chart legend: **Symbicort** **Pulmicort** **Fasenra** **Others**
 Absolute values at actual exchange rates; changes at CER and for H1 2018, unless otherwise stated.

US competitive; new medicines, Emerging Markets encouraging

US -10%

- *Symbicort* (-21%); relatively stable volumes in continued price-competitive environment

Europe -2%

- Relatively stable *Symbicort* volume

Japan +7%

Emerging Markets +13%

- *Pulmicort* supply normalised in China

Fasenra launch performing strongly

US \$67m

- Very encouraging launch
- Leading novel biologic (within IL-5 class)

Europe \$8m

- Germany majority of sales
- Launched in other EU markets

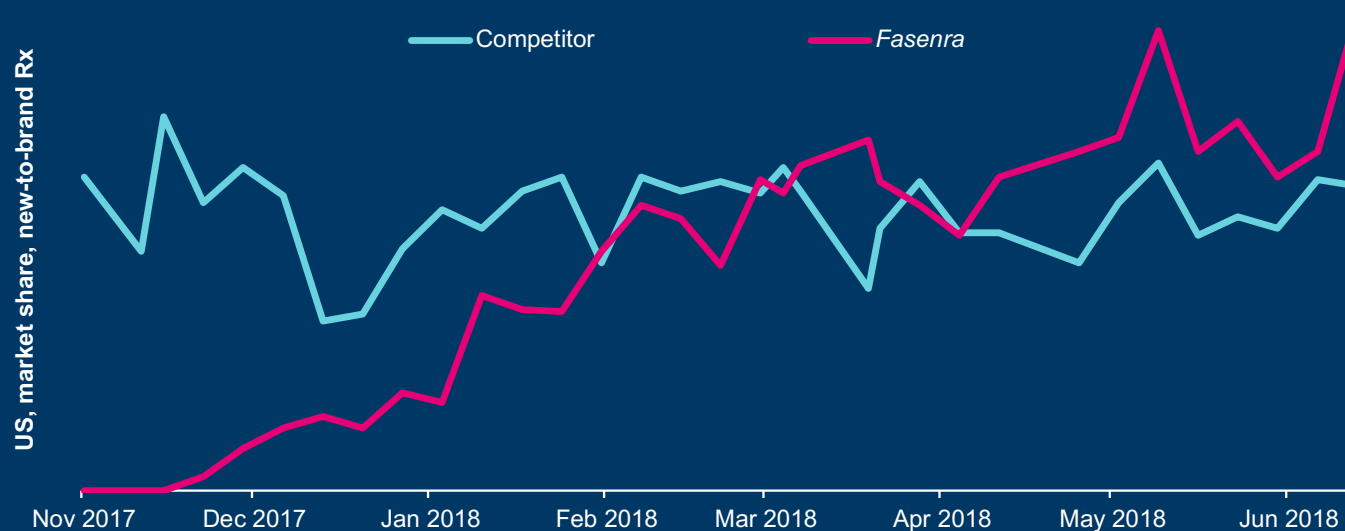
Japan \$11m

- Very strong early uptake



Respiratory: *Fasenra*

The leading novel respiratory biologic



\$65m

Q2 2018 Product Sales
(from \$21m in Q1 2018)

Launched

Fasenra now launched in the
US, Japan and the EU
(Germany, Denmark, Sweden)
with access also in Hong Kong,
Qatar and Saudi Arabia

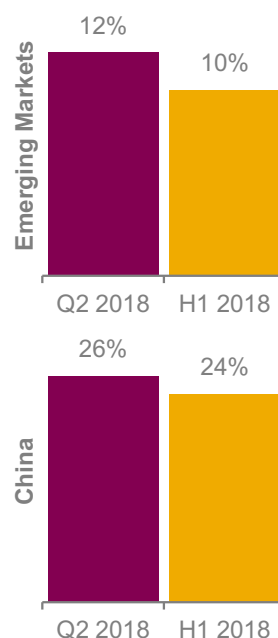
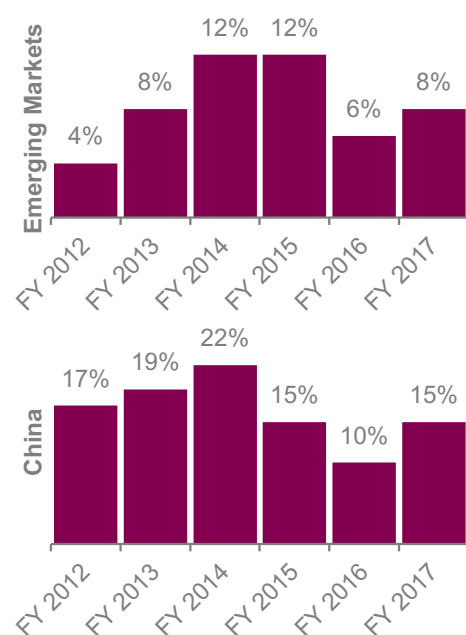
Source: IQVIA



Emerging Markets

China continued strongly

China continued very strongly (+24%)
Ex-China growth (-3%) impacted by divestments



All three main therapy areas performed well

- **Mid to high single-digit growth continued**
 Growth ex-China reduced by divestments (7-8% impact) and general economic conditions in Russia
- **Oncology +37%:** *Tagrisso* (\$159m) now second-biggest Oncology medicine. Hormone-receptor medicines continued growth, with *Faslodex* leading
- **New CVRM +32%:** *Brilinta* (+17%); *Forxiga* (+59%)
- **Respiratory +13%:** *Pulmicort* (+15%, \$482m) normalised supply in China. *Symbicort* (+10%, \$241m)

Absolute values at actual exchange rates; changes at CER and for H1 2018, unless otherwise stated.



Agenda



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Oncology



CVRM, Respiratory, EMs



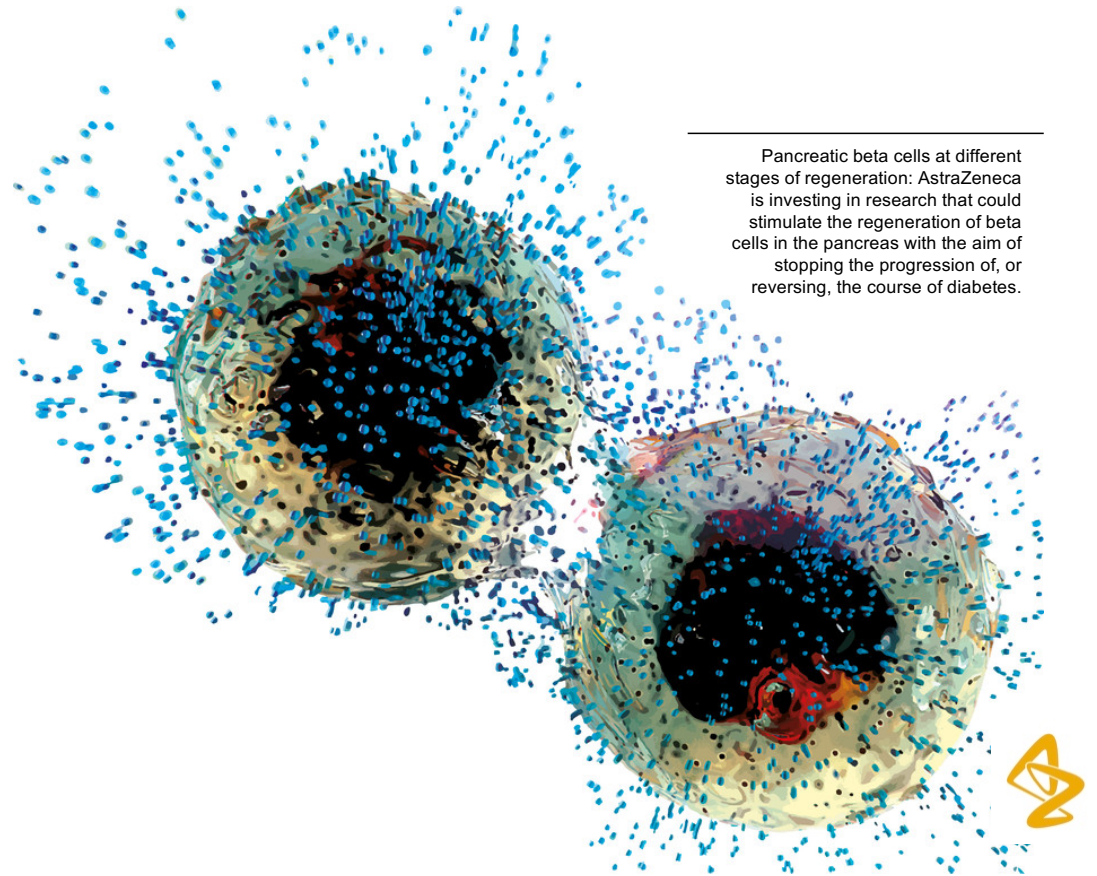
Finance



Pipeline and news flow



Closing and Q&A



Pancreatic beta cells at different stages of regeneration: AstraZeneca is investing in research that could stimulate the regeneration of beta cells in the pancreas with the aim of stopping the progression of, or reversing, the course of diabetes.

Reported Profit & Loss

	H1 2018 \$m	% change	% Total Revenue	Q2 2018 \$m	% change	% Total Revenue
Total Revenue	10,333	(5)	100	5,155	(1)	100
- Product Sales	10,015	(2)	97	5,030	(1)	98
- Externalisation Revenue	318	(54)	3	125	14	2
Gross Margin	78.6%	(3) pp ¹	-	79.9%	(2) pp	-
Operating Expenses ²	7,814	(1)	76	3,997	2	78
- R&D Expenses	2,641	(9)	26	1,362	(1)	26
- SG&A Expenses	5,008	3	49	2,551	4	50
Other Operating Inc. & Exp.	1,086	28	11	617	2	12
Tax Rate	19.2%	-	-	22.6%	-	-
EPS	\$0.54	(34)		\$0.27	(38)	

1. Percentage points. 2. Includes Distribution Expense.

Absolute values at actual exchange rates; changes at CER.

Gross Margin reflects Gross Profit derived from Product Sales, divided by Product Sales.



Core Profit & Loss

	H1 2018 \$m	% change	% Total Revenue	Q2 2018 \$m	% change	% Total Revenue
Total Revenue	10,333	(5)	100	5,155	(1)	100
- Product Sales	10,015	(2)	97	5,030	(1)	98
- Externalisation Revenue	318	(54)	3	125	14	2
Gross Margin	80.0%	(3) pp	-	81.3%	(2) pp	-
Operating Expenses ¹	6,877	2	67	3,528	5	68
- R&D Expenses	2,558	(5)	25	1,318	1	26
- SG&A Expenses	4,154	7	40	2,126	8	41
Other Operating Inc. & Exp.	704	(27)	7	580	(8)	11
Tax Rate	18.8%	-	-	19.5%	-	-
EPS	\$1.17	(39)		\$0.69	(26)	

1. Includes Distribution Expense.

Absolute values at actual exchange rates; changes at CER.

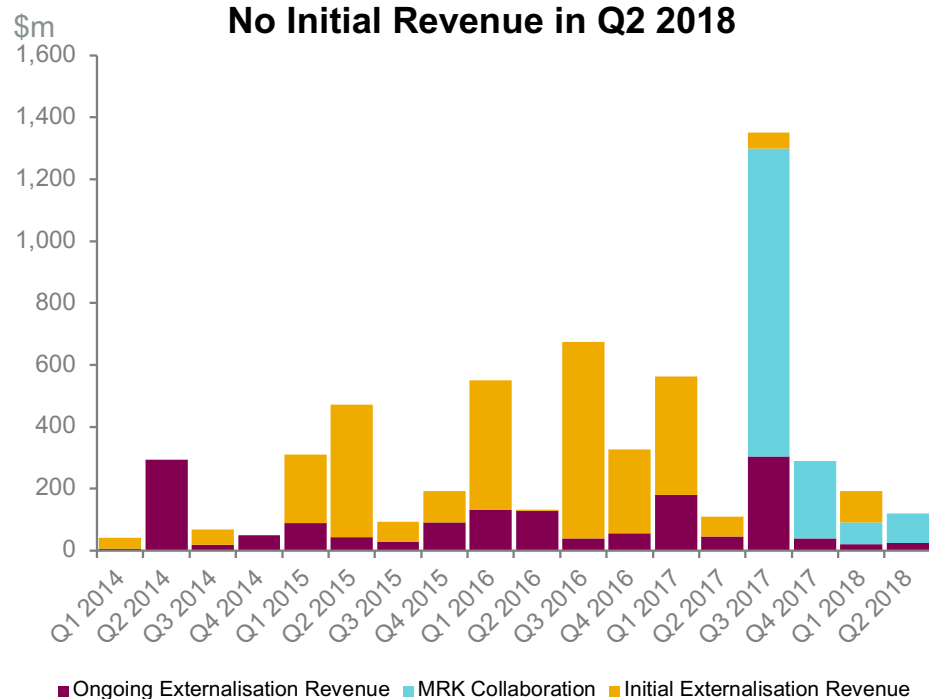
Gross Margin reflects Gross Profit derived from Product Sales, divided by Product Sales.



Externalisation Revenue

Merck collaboration becoming a stable source of income

**Regular payments from Merck
No Initial Revenue in Q2 2018**



**Highlights from
Externalisation Revenue**

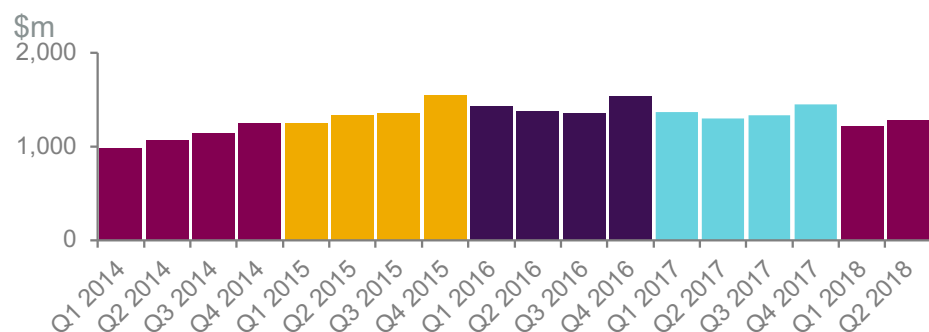
- No initial Externalisation Revenue in Q2; \$102m from partnering legacy medicines in H1
- Ongoing Externalisation Revenue \$216m, mainly from Merck collaboration (*Lynparza* milestones total \$170m, including first sales milestone). A reminder:
 - Regular milestones; approval (~1/3) and sales-related (~2/3); mono and combo therapy
 - Remaining \$500m option payments in 2018-2019

Absolute values at actual exchange rates.

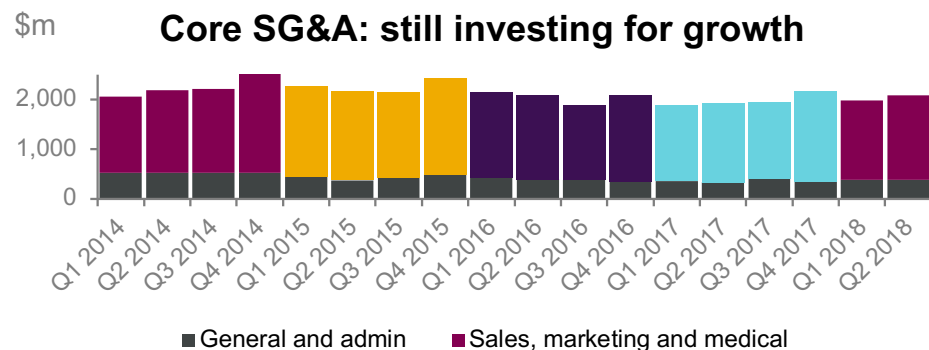


Total Core Operating Expenses increased by 2%

Core R&D: benefitting from productivity initiatives



Core SG&A: still investing for growth



Operating expenses remain in sharp focus

- **Core R&D costs declined by 5%**
 - Maintained activity level; continued benefit from productivity improvements and Merck collaboration
 - FY 2018: anticipated to be in the range of a low single-digit percentage decline to stable
- **Core SG&A costs increased by 7%**
 - Lower baseline in H1 2017; ongoing investment in launches and growth, including in China
 - FY 2018: expected to increase by a low to mid single-digit percentage

Absolute values and changes at CER and for H1 2018, unless otherwise stated.



FY 2018 guidance reiterated; unchanged capital allocation

Product Sales

A low single-digit percentage increase

Core EPS

\$3.30 to \$3.50

Capital allocation

priorities

Investment
in the business

Progressive
dividend policy

Strong, investment-
grade credit rating

Immediately earnings-accretive,
value-enhancing opportunities



Agenda



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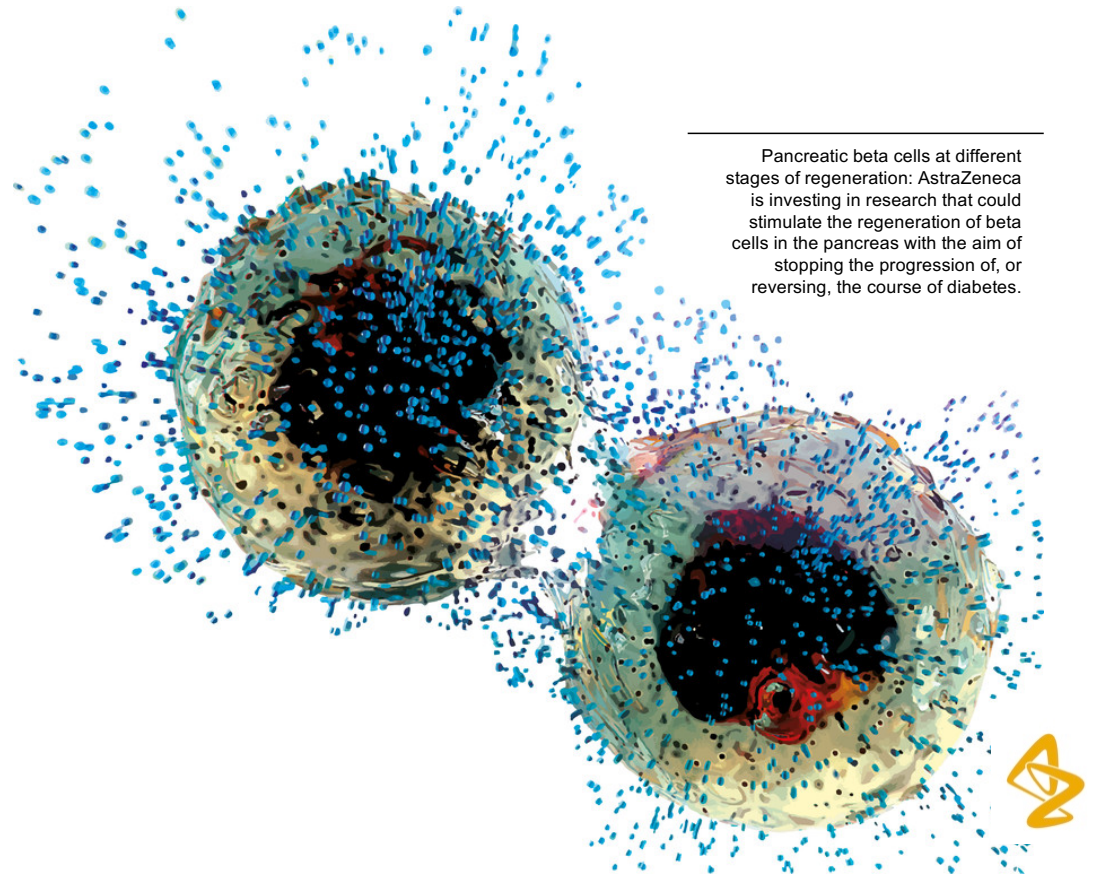
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Pipeline and news flow



Closing and Q&A



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Oncology: continued pipeline success

Next wave of medicines highlighted at ASCO

Regulatory and other development progress

- **Approvals**
 - *Lynparza* - breast cancer (JP)
 - *Tagrisso* - lung cancer 1L (EU)
 - *Imfinzi* - unresectable, Stage III NSCLC (JP)
- **Major Phase III data readouts**
 - *Lynparza* - ovarian cancer 1L - met primary endpoint
 - *Imfinzi* - unresectable, Stage III NSCLC - met primary OS endpoint
 - selumetinib - thyroid cancer - did not meet primary endpoint

American Society of Clinical Oncology (ASCO) Annual Meeting 2018

- ***Lynparza***
Study 08 randomised Phase II trial in prostate cancer
- **selumetinib**
SPRINT Phase II trial in paediatric neurofibromatosis type 1
- **moxetumomab pasudotox**
Study '1053' Phase III trial in hairy cell leukaemia
- **capivasertib (AZD5363, AKT inhibitor)**
PAKT Phase II trial in triple-negative breast cancer
- ***Lynparza* + vistusertib (AZD2014, mTORC1/2 inhibitor)**
Trial in ovarian cancer and triple-negative breast cancer

Status since the last results announcement on 18 May 2018.



CVRM

Lokelma US approval; Farxiga and MEDI0382 in focus at ADA 2018

Regulatory and other development progress

- **Approval**
 - *Lokelma* - hyperkalaemia (US)
- **Positive CHMP opinion**
 - *Bydureon BCise* - type-2 diabetes; new device (EU)
- **Regulatory submission and/or acceptances**
 - *Farxiga* - type-1 diabetes (JP)
 - *Forxiga* combo w/*Onglyza* and metformin - type-2 diabetes (EU)
 - *Bydureon* - type-2 diabetes CVOT (US)

American Diabetes Association (ADA) Scientific Sessions 2018

- **MEDI0382**
 - Phase IIa trial showed significantly-improved glycaemic control and reduced body weight, compared to placebo¹
 - Phase IIb trial underway, with data anticipated in H1 2019
- **Farxiga**
 - *Farxiga* + a DPP-4 inhibitor (*Onglyza*) was
 - 1) non-inferior on HbA1c reduction vs. insulin glargine with or without SUs²
 - 2) achieved significant reduction in HbA1c vs. glimepiride in patients inadequately controlled on metformin³
 - presentation of updated CVD-REAL study real-world evidence data on cardiovascular outcomes



Status since the last results announcement on 18 May 2018.

1. ADA 2018, abstract 1067-P.

2. ADA 2018, abstract 260-OR.

3. ADA 2018, abstract 261-OR.



Respiratory

A biologics portfolio that follows the science

Fasenra: distinctively targets and rapidly depletes eosinophils

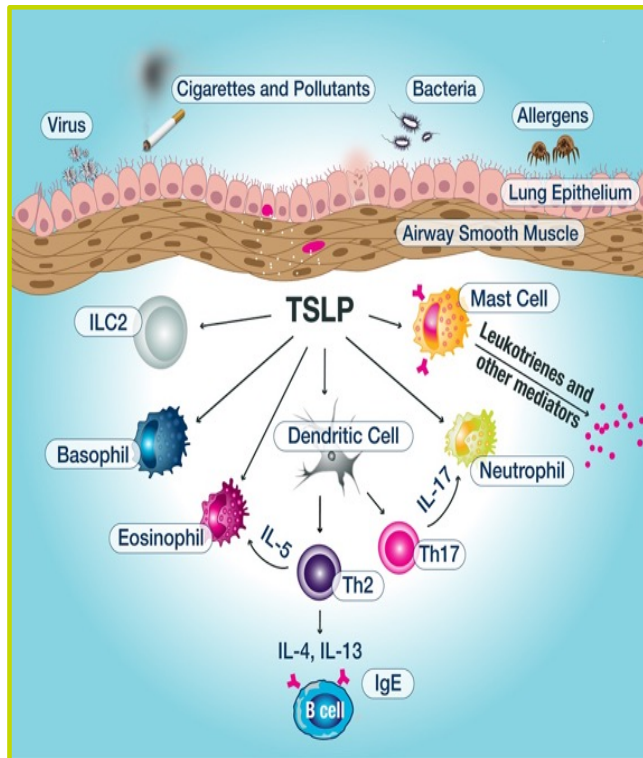
Clear patient phenotype in clinical practice

- Blood eosinophils ≥ 300 cells/ μ L; frequent exacerbator ≥ 3 exacerbations/year; chronic oral corticoid steroids; and nasal polyposis

Lifecycle management

- Home administration and OSTRO Phase III trial initiated in nasal polyposis

Phase III SOLANA (severe, uncontrolled asthma) data anticipated in H2 2018



Tezepelumab: potential best-in-disease Phase II efficacy

- Severe, uncontrolled asthma
- Reduced exacerbation rates 61% - 71%
- Improvements in lung function, asthma control and quality of life
- Unprecedented reductions of key type-2 biomarkers: blood eosinophils, fractional-exhaled nitric oxide and immunoglobulin E

**Phase III programme PATHFINDER
first data anticipated 2019+**



Precision medicines help drive portfolio transformation

Oncology and Respiratory speciality-care medicines are in focus

Diagnostic status			
	Approved	Development	Exploratory
 IRESSA® gefitinib	EGFRm (Epidermal growth factor receptor mutation)		
 Lynparza® olaparib	BRCAm (BRCA Cancer susceptibility gene mutation)	HRRm (homologous recombination repair mutation)	Other
 TAGRISSO™ osimertinib	EGFR and T790Mm ¹		
 IMFINZI™ durvalumab Injection for Intravenous Use 50 mg/mL	PD-L1 (Programmed death-ligand 1 expression level)		TMB (Tumour mutational burden)
 Fasenra™ (benralizumab) Subcutaneous Injection 30 mg		Eosinophilia	

Twenty two

diagnostic test approvals
in three major markets
(US, EU, JP)

\$1.4bn

Precision-medicine
contribution to H1 2018 sales²

27%

Speciality-care medicine
contribution to H1 2018 sales³

1. Substitution of methionine (M) for threonine (T) at amino acid position 790 mutation.

2. Iressa, Lynparza, Tagrisso and Fasenra.

3. Oncology and Fasenra.



Late-stage pipeline news flow in 2018 and 2019

Unlocking and realising the potential of new medicines

	H2 2018	H1 2019	H2 2019
Regulatory decision	Tagrisso - lung cancer 1L (JP) Imfinzi - unresectable, Stage III NSCLC (EU) moxetumomab pasudotox - hairy cell leukaemia 3L (US) Bydureon autoinjector - type-2 diabetes (EU) Bevespi - COPD (EU)	Lynparza - breast cancer (EU)	Forxiga - type-1 diabetes (EU, JP)
Regulatory submission acceptance	Lynparza - ovarian cancer 1L Imfinzi +/- treme - lung cancer 1L (MYSTIC) Duaklir - COPD (US) Bevespi - COPD (JP) PT010 - COPD	Imfinzi +/- treme - head & neck cancer 1L - head & neck cancer 2L selumetinib - neurofibromatosis type 1 Farxiga - type-2 diabetes CVOT Lokelma - hyperkalaemia (JP) roxadustat - anaemia (US) anifrolumab - lupus	Lynparza - pancreatic cancer Imfinzi + treme - lung cancer 1L (NEPTUNE) Imfinzi +/- treme - lung cancer 1L (POSEIDON) - small-cell lung cancer - bladder cancer 1L Calquence - chronic lymphocytic leukaemia Brilinta - CAD ¹ /type-2 diabetes CVOT
Key Phase III data readouts	Imfinzi +/- treme - lung cancer 1L (MYSTIC) (final OS) - head & neck cancer 1L - head & neck cancer 2L Farxiga - type-2 diabetes CVOT roxadustat - anaemia anifrolumab - lupus	Lynparza - pancreatic cancer Imfinzi + treme - lung cancer 1L (NEPTUNE) Brilinta - CAD/type-2 diabetes CVOT	Lynparza - ovarian cancer (1L) (PAOLA-1) Tagrisso - lung cancer (1L) (final OS) Imfinzi +/- treme - lung cancer 1L (POSEIDON) - small-cell lung cancer - bladder cancer 1L Calquence - chronic lymphocytic leukaemia

1. Coronary artery disease.
Status as of 26 July 2018.



Agenda



Overview



Oncology



CVRM, Respiratory, EMs



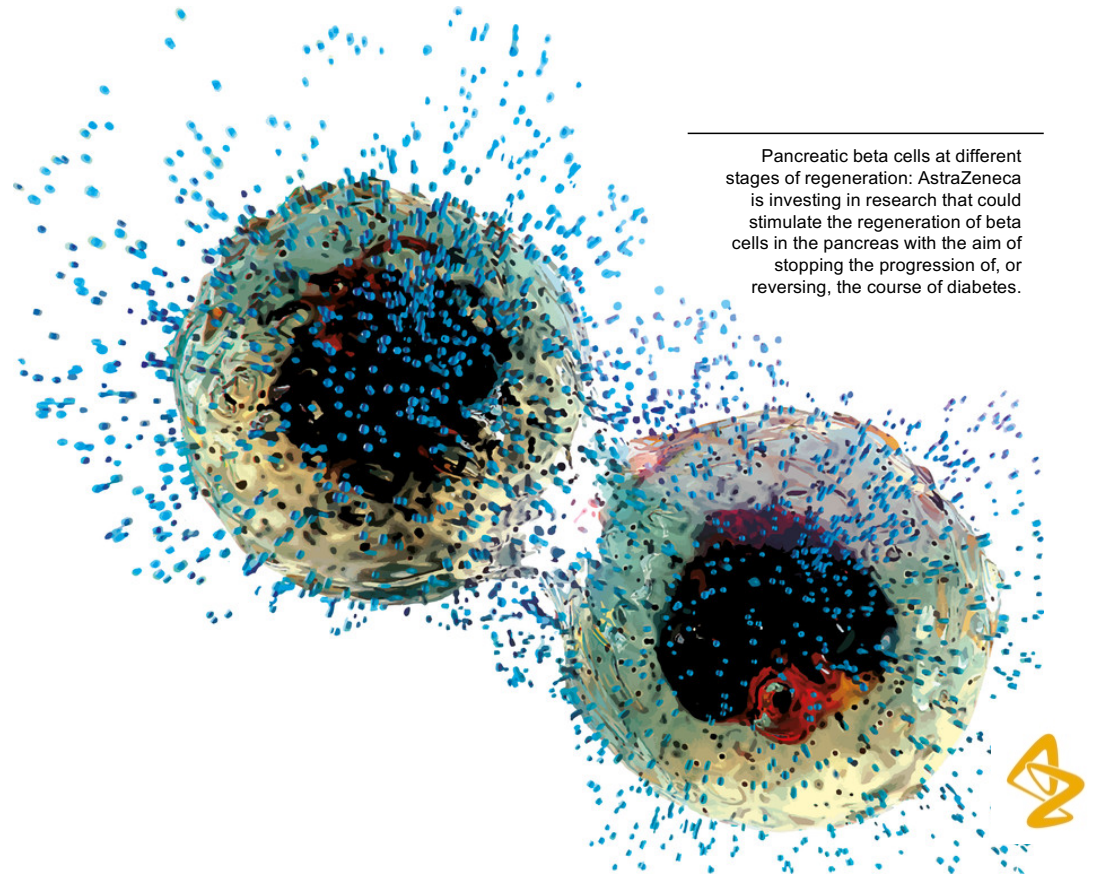
Finance



Pipeline and news flow



Closing and Q&A



Pancreatic beta cells at different stages of regeneration: AstraZeneca is investing in research that could stimulate the regeneration of beta cells in the pancreas with the aim of stopping the progression of, or reversing, the course of diabetes.

Encouraging launches underpin 2018 return to growth

Financials on track - commercial execution - guidance reiterated

- **Financials on track**
 - Product Sales improved, as anticipated
 - Very strong launches offset by the impact of *Crestor* EU/JP and divestments
 - Total Revenue impacted by lower Initial Externalisation Revenue
 - Core Operating Expenses increased by 2%; cost management in focus
- **New medicines delivered >\$1bn in additional sales and grew by 69% vs. H1 2017**
 - *Lynparza*, *Tagrisso*, *Imfinzi* all very strong
 - CVRM blockbusters *Brilinta* and *Farxiga* continued global growth
 - Respiratory improved in Q2 and *Fasenra* consolidated rapid launch
 - China maintained rapid pace
- **Pipeline continued to deliver important news flow**
- **FY 2018 guidance reiterated**

Absolute values, changes and guidance all at CER.



The background of the image is a dense network of neurons, depicted with glowing yellow and orange cell bodies and branching dendrites and axons against a dark, reddish-brown background. The neurons are interconnected, creating a complex web of light. Overlaid on this network is the text 'Q&A' in a large, white, sans-serif font. The text is centered horizontally and vertically, with the ampersand being slightly smaller than the letters. The overall effect is one of a vibrant, active neural network.

Q&A



Encouraging launches underpin 2018 return to growth

Financials on track - commercial execution - guidance reiterated

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Absolute values, changes and guidance all at CER.



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H1 2018 Results

Presentation, conference call and webcast for investors and analysts

26 July 2018

