

H1 2018 Results

Presentation, conference call and webcast for investors and analysts

26 July 2018



Forward-looking statements

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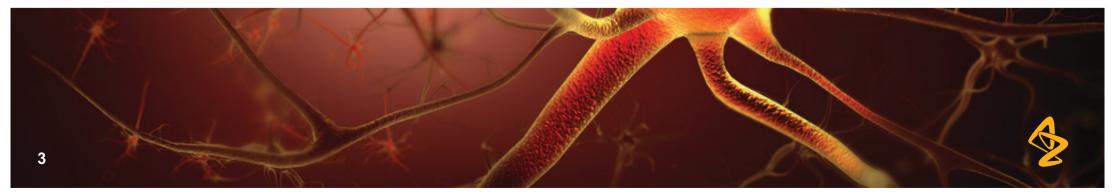
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Agenda



Overview



Oncology



CVRM, Respiratory, EMs



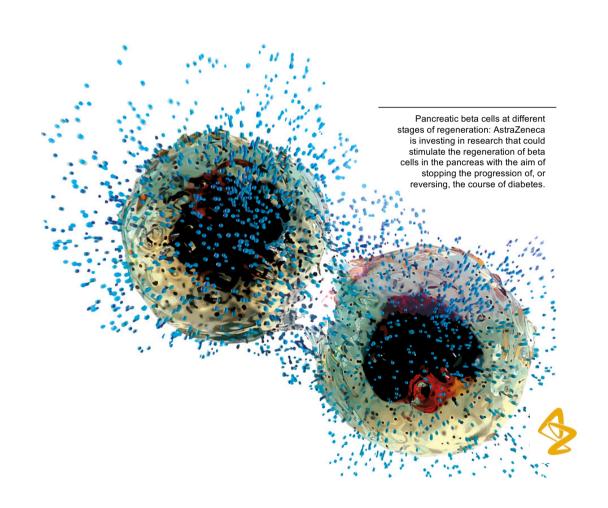
Finance



Pipeline and news flow



Closing and Q&A



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%



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



The pipeline continued to deliver

Late-stage pipeline quarterly highlights

Pipeline news

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

^{1.} Non-small cell lung cancer.



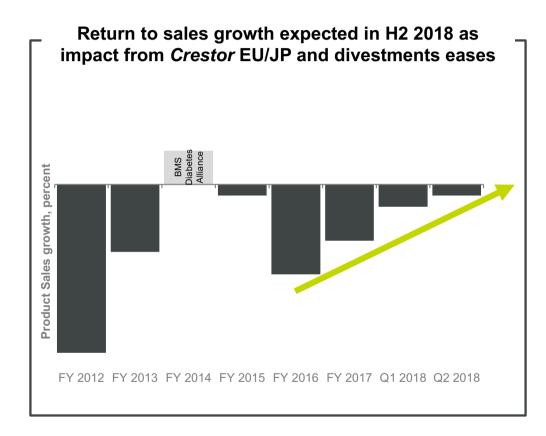
^{2.} Cardiovascular outcomes trial.

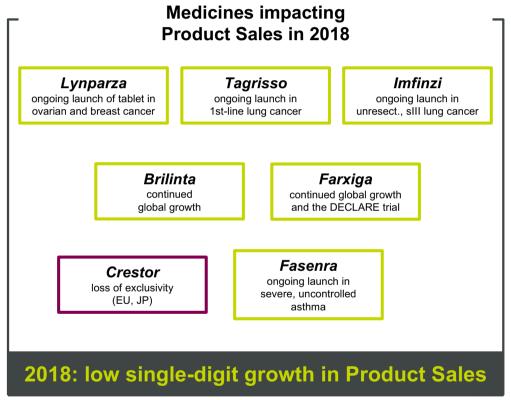
^{3.} Chronic obstructive pulmonary disease.

Status since the last results announcement on 18 May 2018.

2018: return to sales growth on track

The sales momentum continued to improve

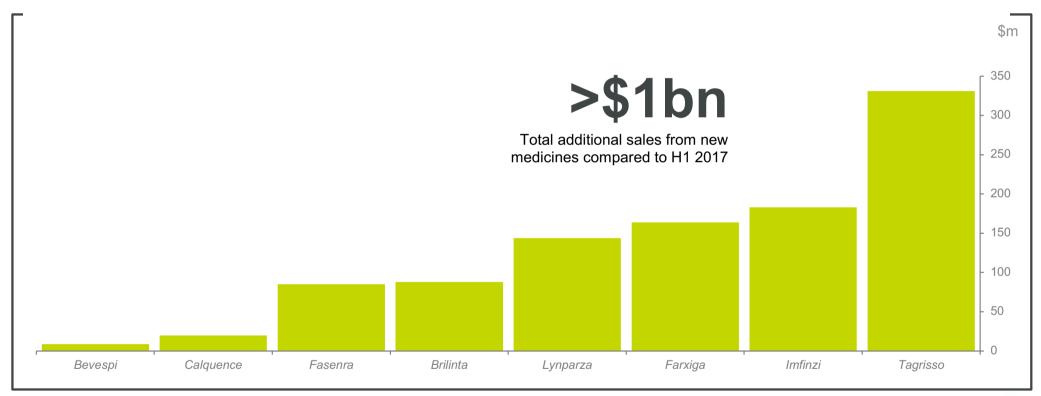






Product Sales: new medicines continued forward

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





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
DODON	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



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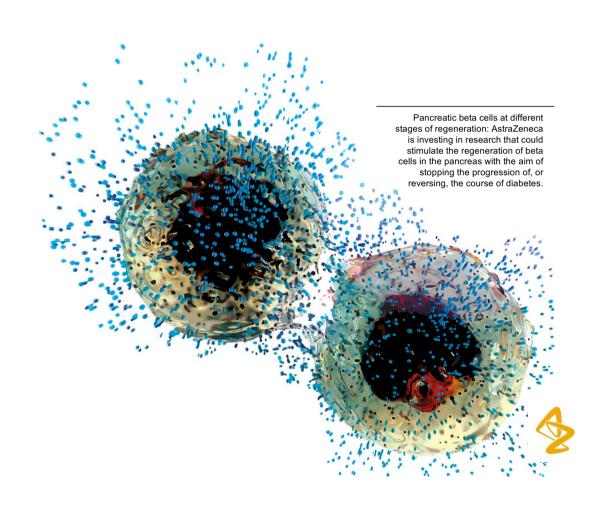
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Pipeline and news flow

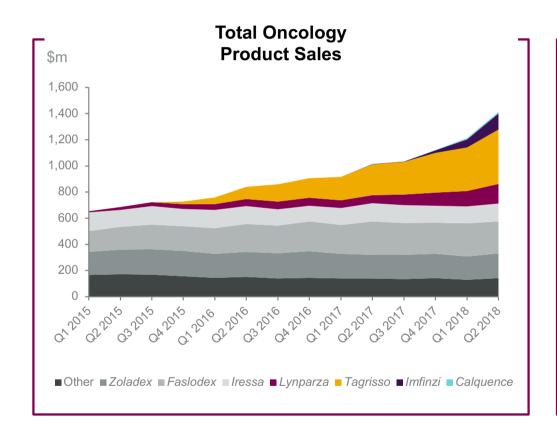


Closing and Q&A



Oncology

New medicines continued to drive strong performance



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

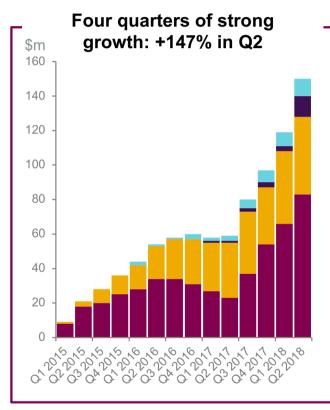
- Oncology +37%; now 27% of total Product Sales
- New medicines contributed \$0.7bn in additional sales vs. H1 2017
 - Lynparza: accelerated growth globally; promising launch in Japan
 - Tagrisso: sustained very high growth; increasing use in 2nd line; encouraging start in the 1st-line setting
 - Imfinzi: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



Lynparza

Expanding benefits to more patients





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





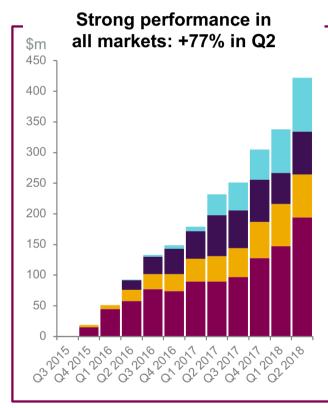




Lung cancer: Tagrisso



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

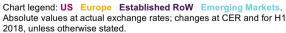


Approved in >75 countries worldwide

- 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

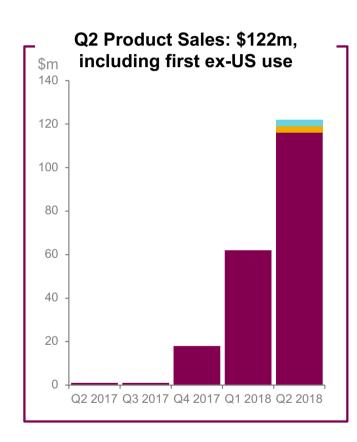




Lung cancer: Imfinzi

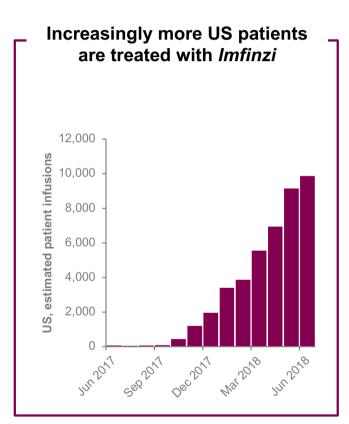


Continued fast uptake in unresectable, Stage III NSCLC (PACIFIC)



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

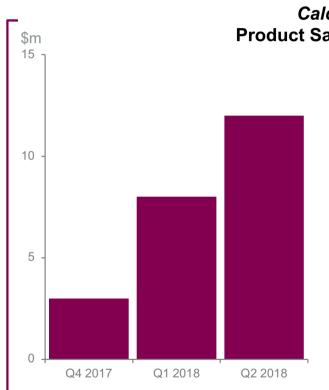






Haematology: Calquence and moxetumomab

Emerging franchise; initially in smaller indications



Calquence Product Sales highlights CALQUEN (gralabutifully many

- Product Sales \$20m, US only
- Encouraging early uptake
 Maintained ~1/4 of new-patient
 starts in approved indication
- Expanding patient benefit
 First ex-US regulatory decision expected in H2 2018
- Lifecycle plans underway in larger indications
 First Phase III data in chronic lymphocytic leukaemia in H2 2019

Moxetumomab pasudotox under US priority review

- First AstraZeneca/MedImmune immunotoxin
- US priority regulatory review with Q3 2018 PDUFA/action date
- Intended indication is 3rd-line hairy cell leukaemia
- Small indication with ~1,000 new US patients per year



New CVRM

Brilinta and Farxiga delivered strong results

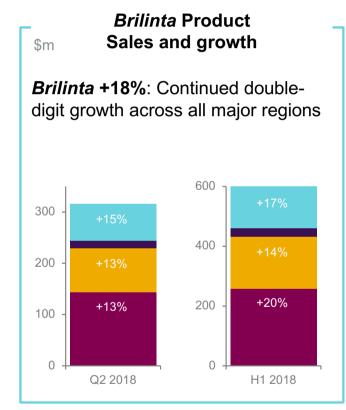


Chart legend: US Europe Established RoW Emerging Markets. Absolute values at actual exchange rates; changes at CER and for H1 2018, unless otherwise stated.

Diabetes Product Sales and growth



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

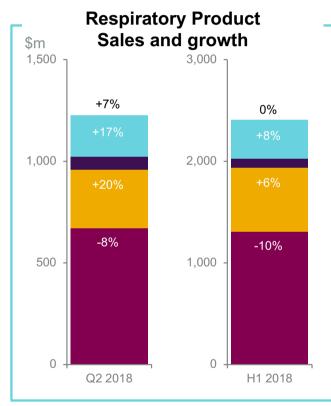
Chart legend: Farxiga Onglyza Bydureon Byetta Other.

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



Respiratory

Improving performance; Fasenra and Pulmicort offsetting Symbicort



US competitive; new medicines, Emerging Markets encouraging

US -10%

 Symbicort (-21%); relatively stable volumes in continued pricecompetitive 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)

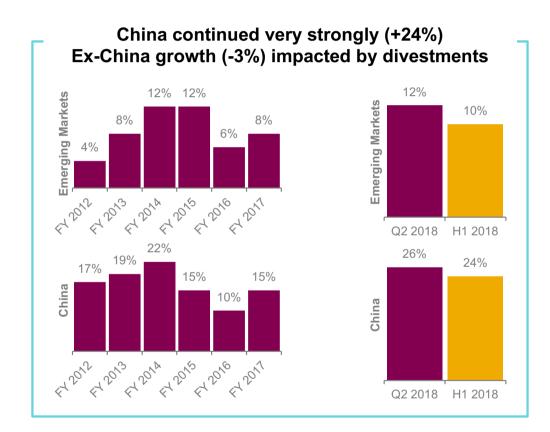
Launched

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



Emerging Markets

China continued strongly



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 secondbiggest 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)



Agenda



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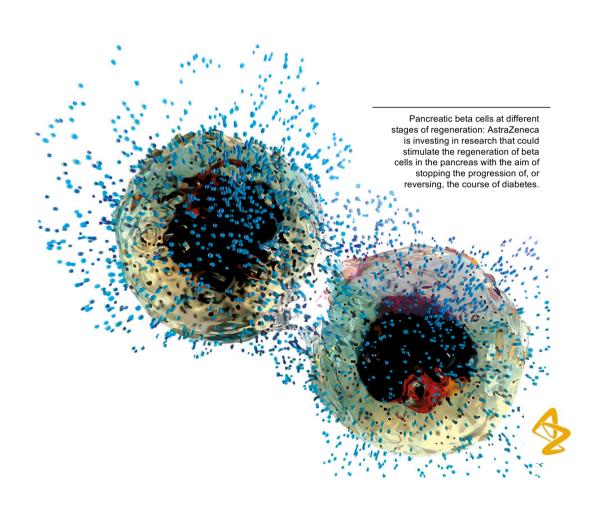
Finance



Pipeline and news flow



Closing and Q&A



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)	

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.

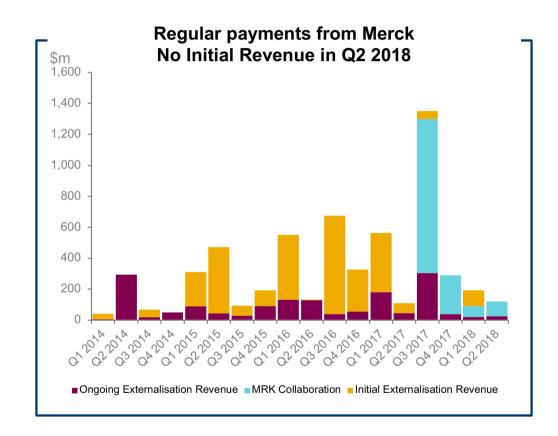


Absolute values at actual exchange rates; changes at CER.

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

Externalisation Revenue

Merck collaboration becoming a stable source of income

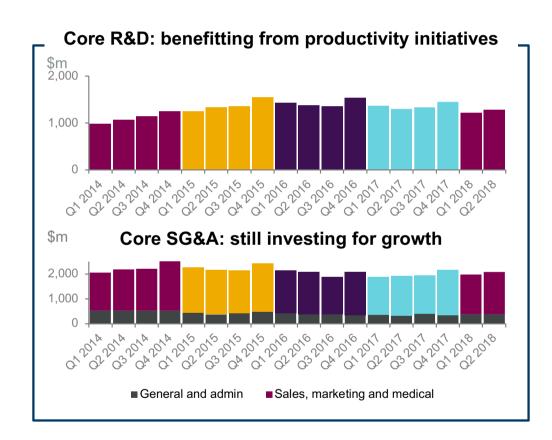


Highlights from Externalisation Revenue

- 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 salesrelated (~2/3); mono and combo therapy
 - Remaining \$500m option payments in 2018-2019



Total Core Operating Expenses increased by 2%



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 singledigit percentage



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, investmentgrade credit rating

Immediately earnings-accretive, value-enhancing opportunities



Agenda



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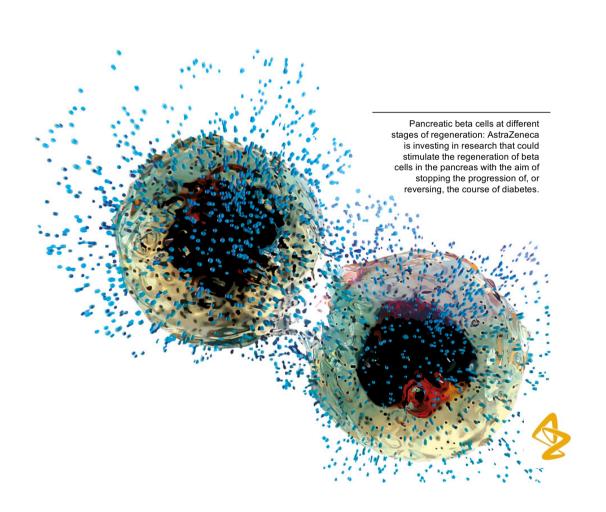
Finance



Pipeline and news flow



Closing and Q&A



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 IIINSCLC 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



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 diabetesCVOT (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



Respiratory

A biologics portfolio that follows the science

Fasenra: distinctively targets and rapidly depletes eosinophils

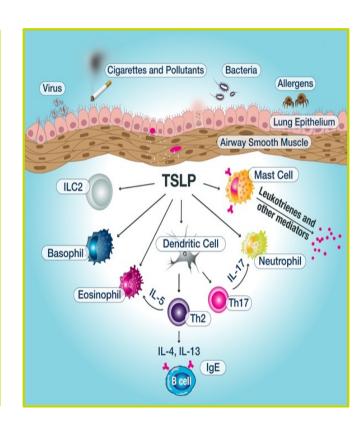
Clear patient phenotype in clinical practice

 Blood eosinophils ≥ 300 cells/µL; 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+

European Respiratory Society International Congress 2017, abstract OA3189.



Precision medicines help drive portfolio transformation

Oncology and Respiratory speciality-care medicines are in focus

Diagnostic status Approved Development Exploratory EGFRm IRESSA® (Epidermal growth aefitinib factor receptor mutation) **BRCAm HRRm** Lynparza[®] Other (BReast CAncer olaparib • III • III • (homologous recombination susceptibility gene repair mutation) mutation) EGFR and TAGRISSO* T790Mm¹ PD-L1 **TMB MFINZI** (Programmed (Tumour mutational death-ligand 1 burden) expression level) **Fasenra** 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.

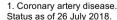
^{2.} Iressa, Lynparza, Tagrisso and Fasenra.

^{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)	<i>Lynparza</i> - breast cancer (EU)	Forxiga - type-1 diabetes (EU, JP)
Regulatory submission	Lynparza - ovarian cancer 1L	Imfinzi +/- treme - head & neck cancer 1L	Lynparza - pancreatic cancer
acceptance	Imfinzi +/- treme - lung cancer 1L (MYSTIC)	- head & neck cancer 2L	Imfinzi + treme - lung cancer 1L (NEPTUNE)
		selumetinib - neurofibromatosis type 1	Imfinzi +/- treme
	Duaklir - COPD (US)		 lung cancer 1L (POSEIDON)
	Bevespi - COPD (JP)	Farxiga - type-2 diabetes CVOT	- small-cell lung cancer
	PT010 - COPD	Lokelma - hyperkalaemia (JP) roxadustat - anaemia (US)	- bladder cancer 1L
			Calquence - chronic lymphocytic leukaemia
		anifrolumab - lupus	
			Brilinta - CAD¹/type-2 diabetes CVOT
Key Phase III data	Imfinzi +/- treme - lung cancer 1L (MYSTIC) (final OS)	Lynparza - pancreatic cancer	Lynparza - ovarian cancer (1L) (PAOLA-1)
readouts	head & neck cancer 1Lhead & neck cancer 2L	Imfinzi + treme - lung cancer 1L (NEPTUNE)	Tagrisso - lung cancer (1L) (final OS) Imfinzi +/- treme
		Brilinta - CAD/type-2 diabetes CVOT	- lung cancer 1L (POSEIDON)
	Farxiga - type-2 diabetes CVOT	••	- small-cell lung cancer
	roxadustat - anaemia		- bladder cancer 1L
			Calquence - chronic lymphocytic leukaemia
	anifrolumab - lupus		





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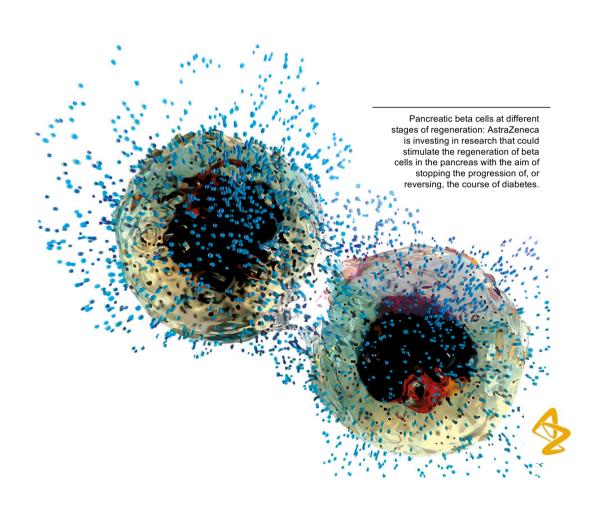
Finance



Pipeline and news flow



Closing and Q&A



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





Encouraging launches underpin 2018 return to growth Financials on track - commercial execution - guidance reiterated

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H1 2018 Results

Presentation, conference call and webcast for investors and analysts

26 July 2018

