2Q and Half Year Results 2013



Cautionary Statement Regarding 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 presentation contains certain forward-looking statements with respect to the operations, performance and financial condition of the Group. 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 presentation 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 patents, marketing exclusivity or trade marks, or the risk of failure to obtain patent protection; the risk of substantial adverse litigation/government investigation claims and insufficient insurance coverage; exchange rate fluctuations; the risk that R&D will not yield new products that achieve commercial success; the risk that strategic alliances and acquisitions will be unsuccessful; the impact of competition, price controls and price reductions; taxation risks; the risk of substantial product liability claims; the impact of any failure by third parties to supply materials or services; the risk of failure to manage a crisis; the risk of delay to new product launches; the difficulties of obtaining and maintaining regulatory approvals for products; the risk of failure to observe ongoing regulatory oversight; the risk that new products do not perform as we expect; the risk of environmental liabilities; the risks associated with conducting business in emerging markets; the risk of reputational damage; the risk of product counterfeiting; the risk of failure to successfully implement planned cost reduction measures through productivity initiatives and restructuring programmes; the risk that regulatory approval processes for biosimilars could have an adverse effect on future commercial prospects; and the impact of increasing implementation and enforcement of more stringent anti-bribery and anti-corruption legislation. Nothing in this presentation should be construed as a profit forecast.



2Q and Half Year Results 2013

Pascal Soriot, Chief Executive Officer



2Q highlights

- 2Q revenue down 4% CER: ~ \$500m decline from loss of exclusivity
- Good progression of our growth platforms: \$400m (double-digit growth @ CER)
- Growing late-stage pipeline:
 - 2 NDA submissions
 - 1 Phase 3 start
 - 3 late-stage projects added via business development

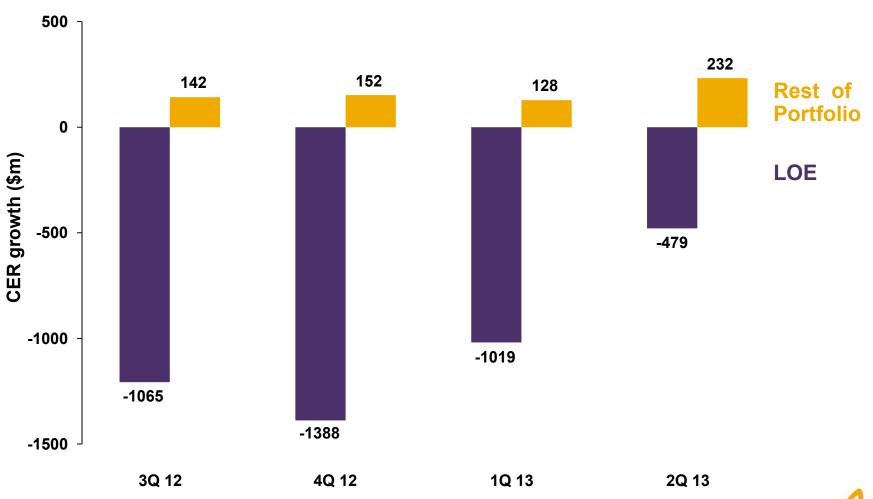


Headline results 2Q 2013

	2013 \$m	2012 \$m	Actual growth	CER growth
Revenue	6,232	6,660	-6%	-4%
Core Operating Profit	2,056	2,334	-12%	-10%
Core EPS	\$1.20	\$1.57	-23%	-21%
Restructuring	(\$0.18)	(\$0.12)		
Intangible amortisation	(\$0.27)	(\$0.13)		
Intangible impairments	(\$0.10)	(\$0.01)		
Legal provisions/other	\$0.01	(\$0.05)		
Reported EPS	\$0.66	\$1.26	-48%	-44%



Revenue impact from loss of exclusivity is moderating



5 key growth platforms



1. Brilinta



2. Diabetes



3. Emerging Markets



4. Respiratory



5. Japan



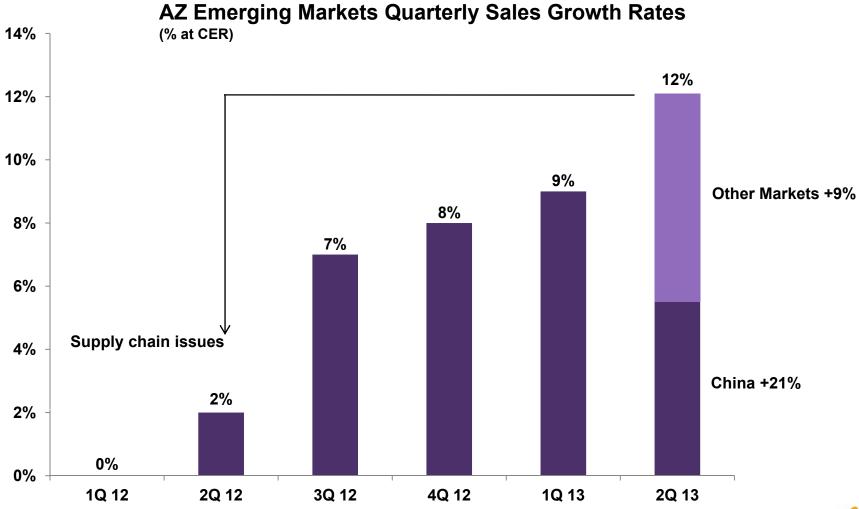
Regional revenue performance 2Q 2013

2013	CER	CER
\$m	growth	\$m

Global Revenue	6,232	-4%	(247)	
US	2,252	-4%	(87)	Ex-Seroquel IR & Toprol-XL: +4%
Europe	1,546	-13%	(231)	Seroquel IR, Seroquel XR, Atacand & Nexium LOE
Established ROW	1,059	-6%	(83)	
Japan	657	+10%	72	Nexium, Crestor , & Symbicort
Canada	162	-42%	(121)	Crestor (70)
Other Established ROW	240	-12%	(34)	Crestor pricing and Seroquel IR
Emerging Markets	1,375	+12%	154	China (+21%); Other Markets (+9%)
China	431	+21%	72	

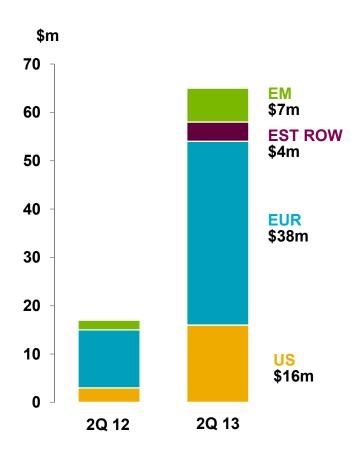


Good performance in Emerging Markets



Brilinta/Brilique

2Q 2013 Sales: \$65m



US

- US total TRx's increased 33% vs. prior quarter
 - Gained 9 points of preferred, unrestricted access in Medicare Part D plans, now >70%
 - Delivering acceleration plan, including 200 Transition of Care nurses in field 1st July
 - Increase in promotional and scientific investment in 2013 by greater than 50%

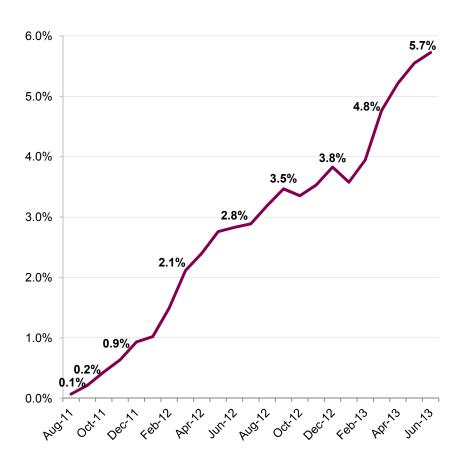
ROW sales \$49m

 At or closing in on Number 2 in volume share of the OAP market in the UK, Germany & Italy. Steady progress in France

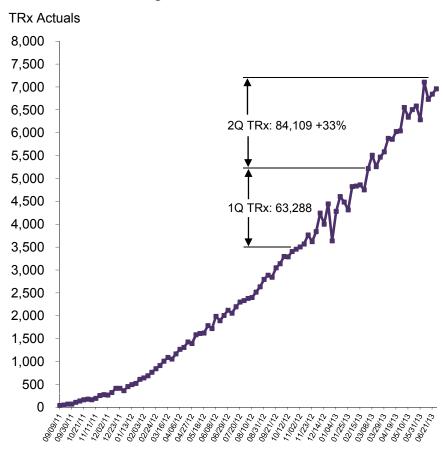


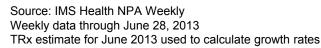
Brilinta US: Steady growth

Increasing New to Brand share in OAP market



TRx growth +33% 2Q vs 1Q 2013

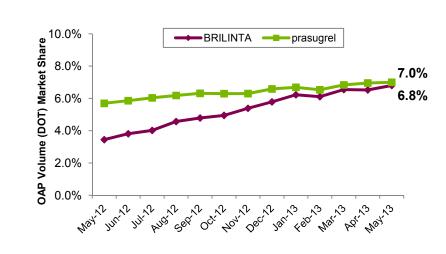




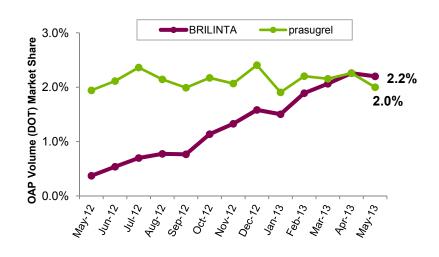


Brilinta: ROW Performance

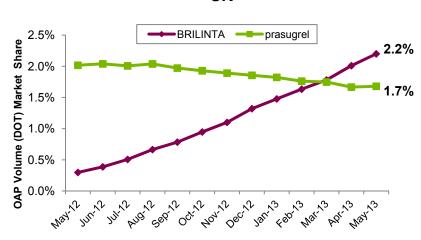
OAP Volume Market Share – All Channels Germany



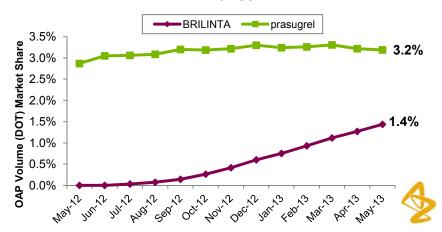
OAP Volume Market Share – All Channels Italy



OAP Volume Market Share – All Channels UK

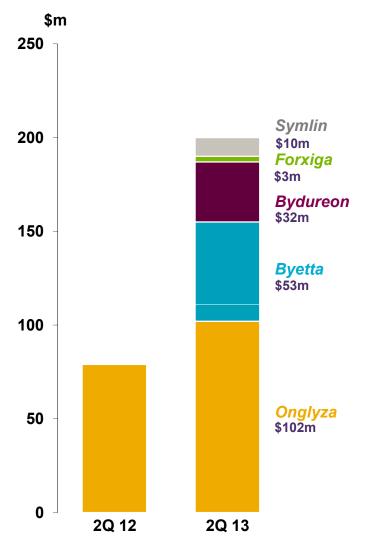






Diabetes Franchise

2Q 2013 Revenue: \$200m



Onglyza franchise

- US revenue: +29%
 - DPP4 Market growth: +6% TRx
 - Franchise TRx share 16.0% in June; stabilised
 - (-10bps from March) post formulary changes in 1Q
- ROW revenue +23%
- SAVOR top line results

Byetta/Bydureon

- US revenue: \$63 million
 - Steady growth for *Bydureon* TRx, but *Byetta* share decline continues
- ROW revenue: \$22 million

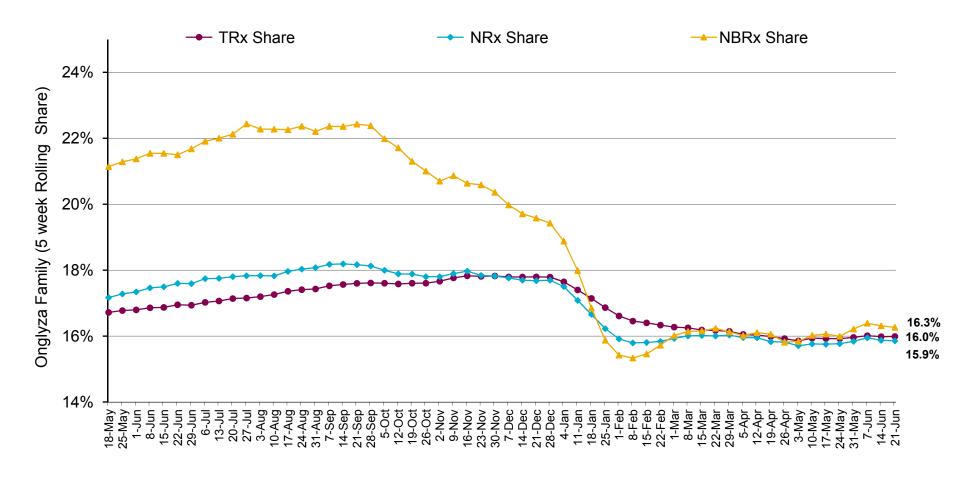
Forxiga

- Revenue \$4 million in 1H 13
- Good initial launch uptake in Germany, UK
- US NDA resubmitted; PDUFA date 11 January 2014



Source: IMS HEALTH MDART

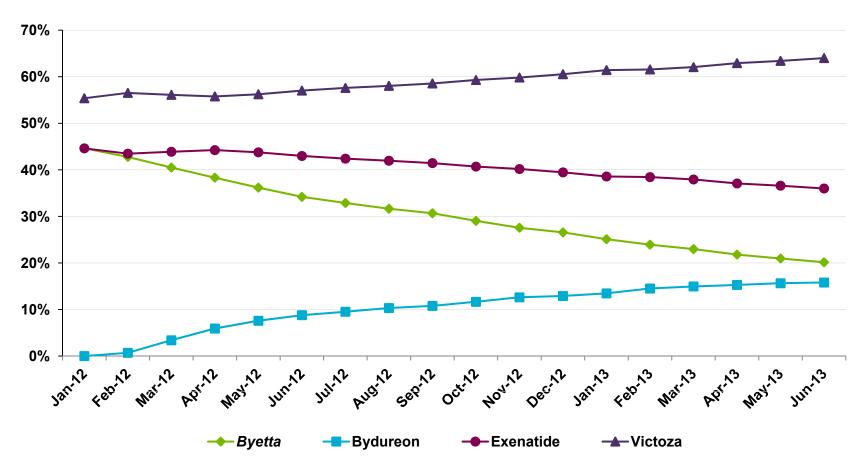
Onglyza: US franchise market share stabilised





Bydureon/Byetta: US TRx trends

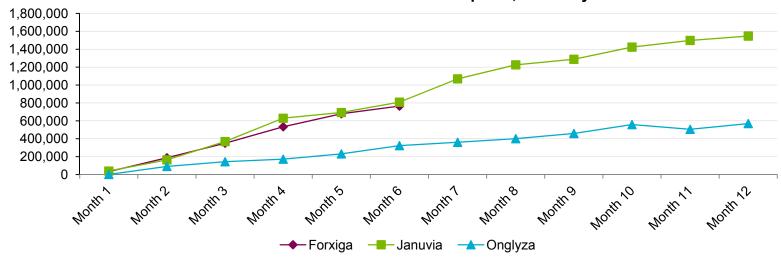
GLP-1 Market TRx



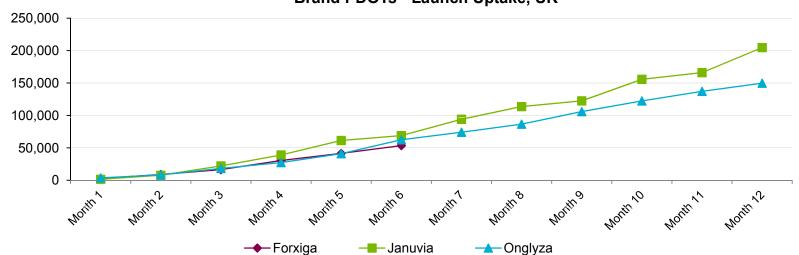


Forxiga launch uptake – Germany and UK





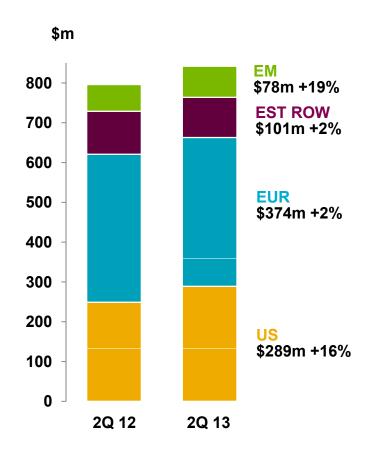
Brand PDOTs - Launch Uptake, UK





Symbicort

2Q 2013 Sales: \$842m +8% CER



US

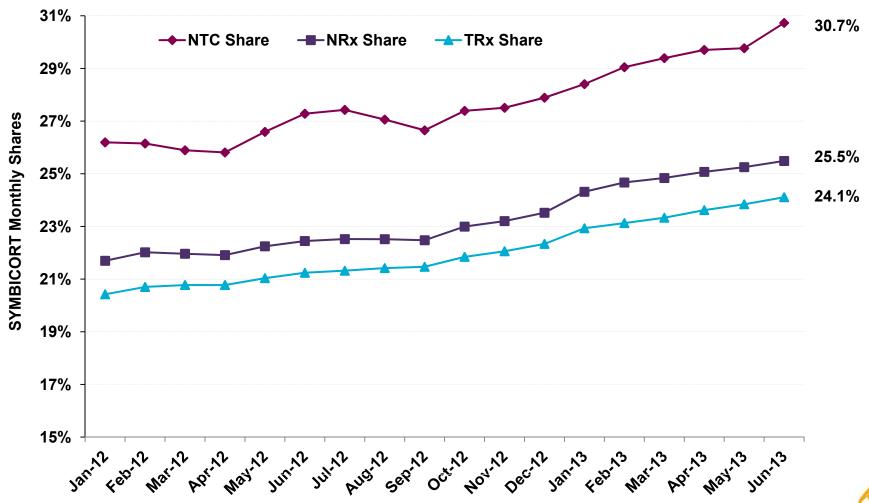
- Symbicort TRx's +16% vs market +2%
- TRx share at 24.1% in June 2013, +1.8 pts since December 2012
- New patient share 30.7% in June; new high

ROW

- Symbicort sales in ROW were \$553m, +4%
- Japan still gaining share on Adoair, reaching 42.3% (May)
 - SMART and COPD approvals



Symbicort: US market share trend





Achieve scientific leadership

1

FOCUS on distinctive science in 3 core TAs

Achieve scientific leadership

PRIORITISE & **ACCELERATE** our pipeline

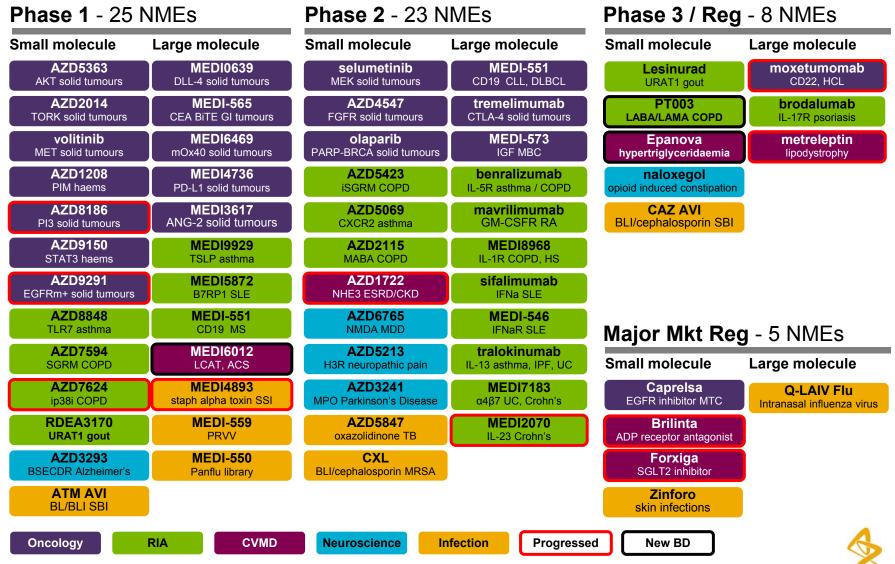
TRANSFORM our innovation culture & model

Recent pipeline news

- Forxiga (dapagliflozin) NDA resubmitted in US; 11 January 2014 PDUFA
- Moxetumomab Phase III start
- Headline results for SAVOR study
 - Onglyza met its primary safety objective of non-inferiority and did not meet the primary efficacy objective of superiority for a composite endpoint (CV death, non-fatal MI, non-fatal ischaemic stroke)
 - Presentation at ESC in September
- Metreleptin US NDA accepted; designated for priority review (PDUFA 27 Feb 2014)
- Fostamatinib will not progress to regulatory filings
- Growing portfolio of late stage assets; Omthera and Pearl acquisitions, FibroGen collaboration



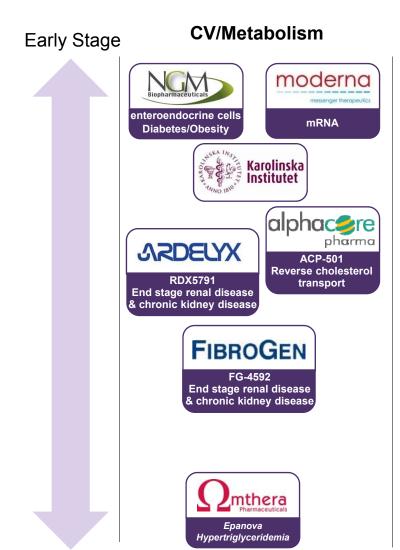
Growing late-stage pipeline



Terminated: MEDI-557, AZD1446, AZD3480, MEDI5117, AZD8330, fostamatinib, MEDI-575, MEDI7814, and MEDI4212

Pipeline table as of 30th June 2013

Business development aligned with priority TAs



messenger therapeutics mRNA THERAPEUTICS ACCURINTM nanomedicine UNIVERSITY OF CAMBRIDGE RESEARCH UK CAMBRIDGE R

3 preclinical projects

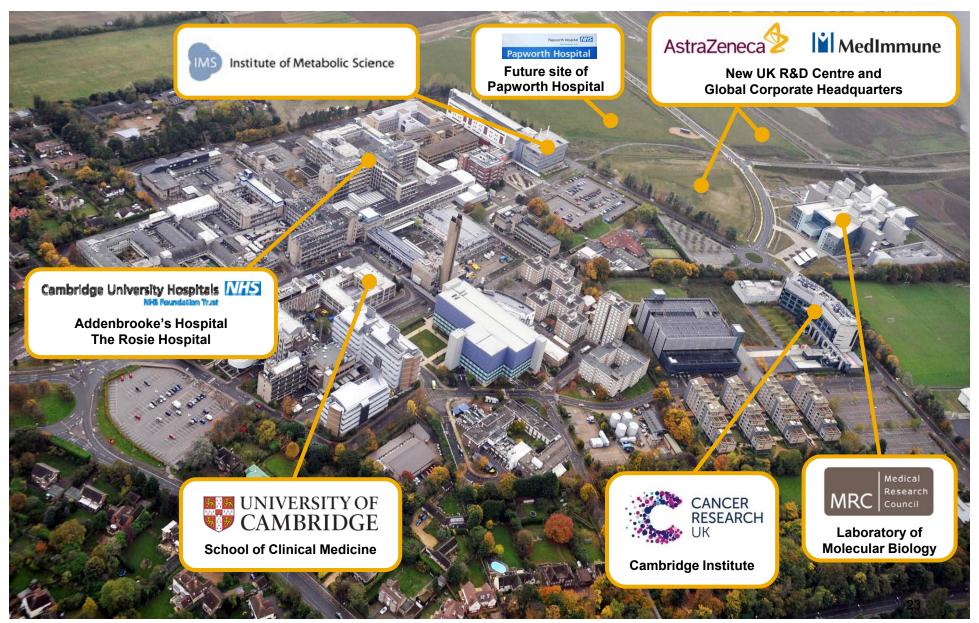
Oncology







Cambridge Biomedical Campus: A vibrant innovation hub



2Q and Half Year Results 2013

Simon Lowth, Chief Financial Officer



Core margin: 2Q 2013

	\$m	CER %	% sales	Delta vs PY CER
Revenue	6,232	-4%	-	
Core Gross Margin	5,127	-2%	82.3	+110 bps
Distribution	(76)	+2%	1.2	-10 bps
Core SG&A	(2,173)	+6%	34.9	-320 bps
Core Other Income	218	+19%	3.5	+60 bps
Core Pre-R&D Profit	3,096	-6%	49.7	-160 bps
Core R&D	(1,040)	+1%	16.7	-70 bps
Core Operating Profit	2,056	-10%	33.0	-230 bps



Phase 4 restructuring programme

	Total cost \$m	1Q 13 \$m	2Q 13 \$m
R&D	1,380	291	62
SG&A	790	240	160
COGS	130	12	86
Total	2,300*	543	308

Phasing

Costs: ~\$1,300m in 2013; balance split broadly evenly between 2014/15; small residual in 2016

Benefits: \sim \$800m by end 2016, around $\frac{1}{2}$ by end 2014



^{*} Of which cash costs are ~\$1,700m

1H 2013: Cash flow/distributions

Cash generated from operating activities \$3.8 billion (1H 2012 \$2.8 billion)

- Lower tax and interest payments partially offset the lower operating profit in 2013 (which included higher non-cash costs)
- A lump sum pension contribution drove higher outflows in the prior year

Dividends

- First interim dividend: \$0.90
 - Consistent with Board's aim to set first interim dividend at approximately 1/3 of prior full year dividend (FY 2012: \$2.80)
- Committed to progressive dividend policy
 - Board intends to maintain or grow the dividend each year
 - Cover target is 2x Core earnings over the investment cycle
 - Board recognises cover in any year is likely to vary from the 2x cover target through the investment cycle



Guidance (Core basis)

Revenue (CER)	Mid to High-single digit decline	
Core Operating Costs (CER) (R&D + SG&A)	Revised: Now a low to mid single digit increase vs 2012	
Core other income	< \$600 million (Pending resolution of Pulmicort Respules appeal)	
Reported tax rate	Around 23 percent	
Core EPS (CER)	Decline significantly > than revenue decline	



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