

Full Year Results 2012

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Full Year Results 2012

Pascal Soriot, Chief Executive Officer

Agenda

Pascal Soriot

- Opening remarks
- Commercial performance

Briggs Morrison

- Pipeline review

Simon Lowth

- Financial performance 2012
- 2013 guidance

Pascal Soriot

- Observations from 1st 90 days
- Capital Markets day preview



Overview 2012

Achieved financial target

Revenue	\$27,973m	-15% (CER)
Core EPS	\$6.41	-9% (CER)
Reported EPS	\$4.99	-29% (CER)
Dividend	\$2.80	

Drove Brand performance

Achieved \$600m in CER revenue growth from:

- *Symbicort*
- *Onglyza*
- *Faslodex*
- *Brilinta/Brilique*
- *Iressa*
- *Seroquel XR*

Advanced the Pipeline

Europe	<i>Forxiga, Zinforo, Caprelsa</i>
US	<i>FluMist Quadrivalent</i>
Japan	<i>Symbicort SMART, Symbicort COPD, Nexium + LDA</i>

Accelerated Business Development

- Amgen collaboration** (5 clinical projects)
- Ardea acquisition** (Lesinurad in Phase III for gout)
- Expanded diabetes alliance** (Amylin portfolio)



Regional revenue performance FY 2012

	2012 \$m	CER growth	CER \$m	
Global Revenue	27,973	-15%	(4,966)	Astra Tech/Aptium (-1.7 pts)
US	10,655	-21%	(2,771)	<i>Seroquel IR (2,647)</i>
Western Europe	6,486	-19%	(1,607)	<i>(Seroquel IR, Nexium, Atacand & Merrem generics)</i>
Established ROW	5,080	-14%	(802)	
Japan	2,904	-5%	(152)	
Canada	1,090	-31%	(497)	<i>(Crestor)</i>
Other Established ROW	1,086	-12%	(153)	
Emerging Markets	5,752	+4%	214	
Emerging Europe	1,165	+2%	22	<i>(Turkey: gov't price interventions)</i> <i>Good growth in Russia</i>
China	1,512	+17%	213	
Emerging Asia Pacific	923	-3%	(25)	<i>(India: local supply issues)</i>
Other Emerging ROW	2,152	--	4	<i>(Mexico: generics and difficult market)</i> <i>(Brazil: generics for Crestor/Seroquel IR)</i>



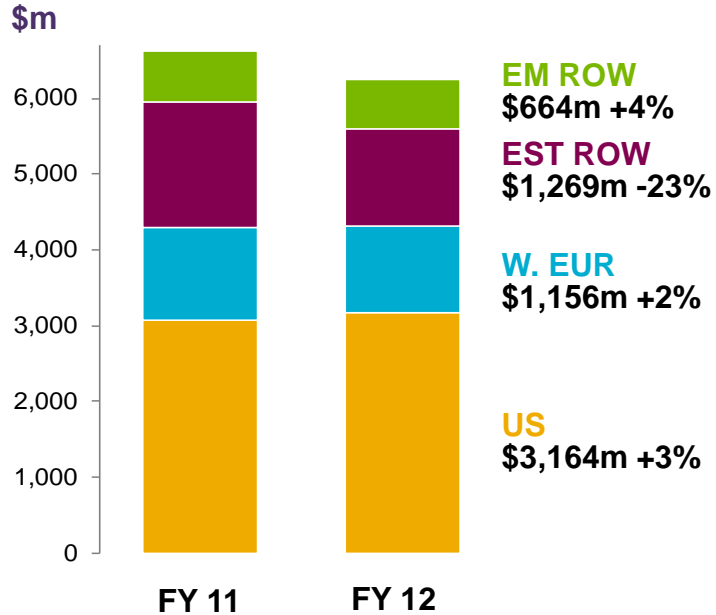
Brand revenue performance FY 2012

	2012 \$m	CER growth	CER \$m	
Global Revenue	27,973	-15%	(4,966)	
<i>Crestor</i>	6,253	-4%	(246)	Excluding Canada +2%
<i>Symbicort</i>	3,194	+5%	157	
<i>Seroquel XR</i>	1,509	+4%	64	
<i>Faslodex</i>	654	+24%	131	
<i>Iressa</i>	611	+12%	69	
<i>Onglyza</i>	323	+53%	112	
<i>Byetta/Bydureon</i>	111	n/m	111	
<i>Brilinta/Brilique</i>	89	+348%	73	
<i>Nexium</i>	3,944	-10%	(422)	
<i>Seroquel IR</i>	1,294	-70%	(3,026)	
<i>Atacand</i>	1,009	-27%	(394)	
<i>Merrem</i>	396	-29%	(167)	



Crestor

2012 Sales: \$6,253m -4% CER

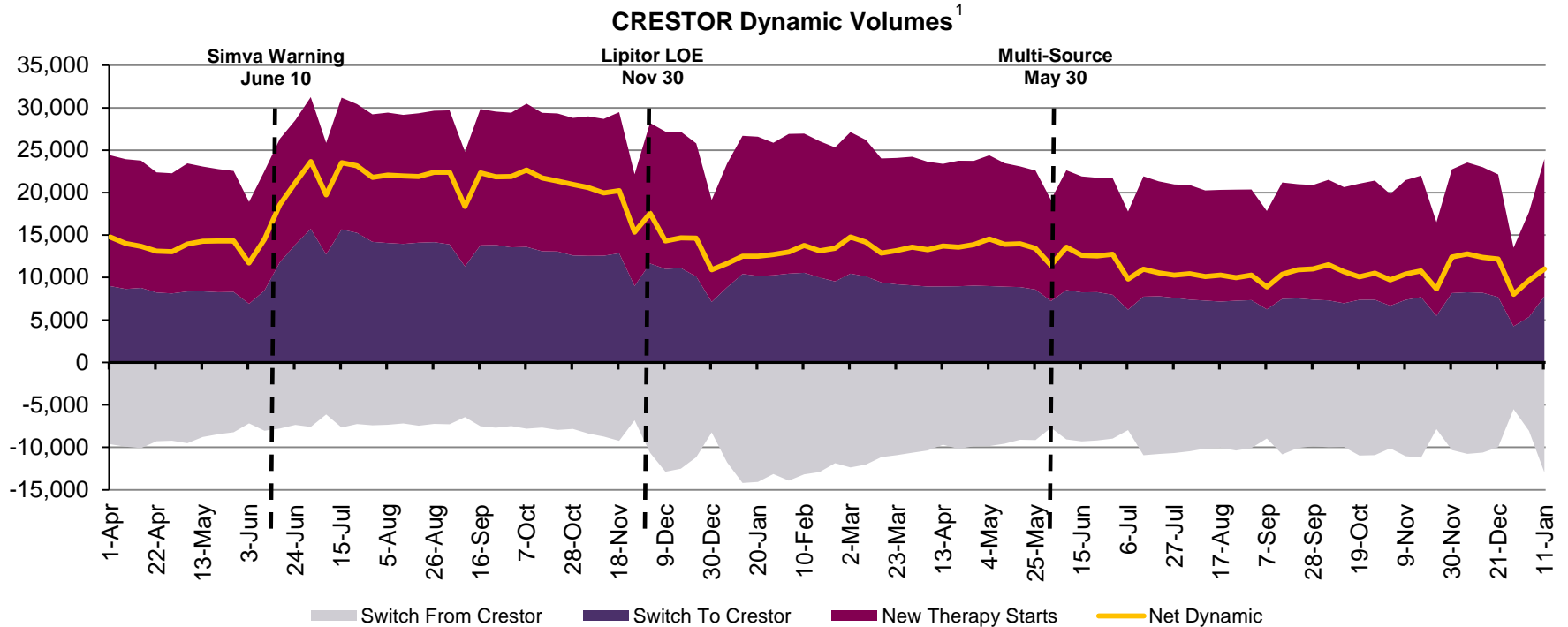


US

- FY TRx's -1.4%
- Resilient performance post generic atorvastatin



Crestor US: net dynamic volume trend



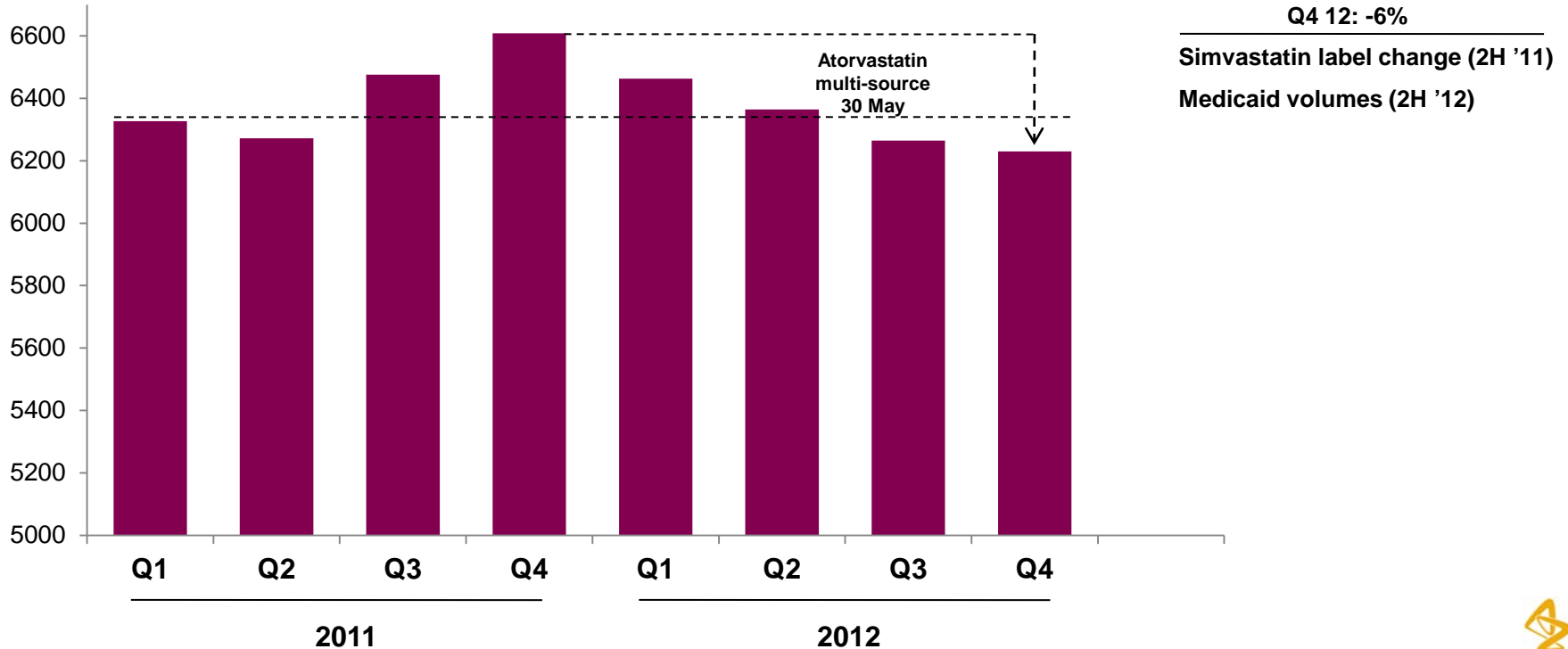
Source: IMS NPA Market Dynamics, Data Week ending 01/11/13

1 - Retail dynamic volumes only; does not include mail order or long term care business



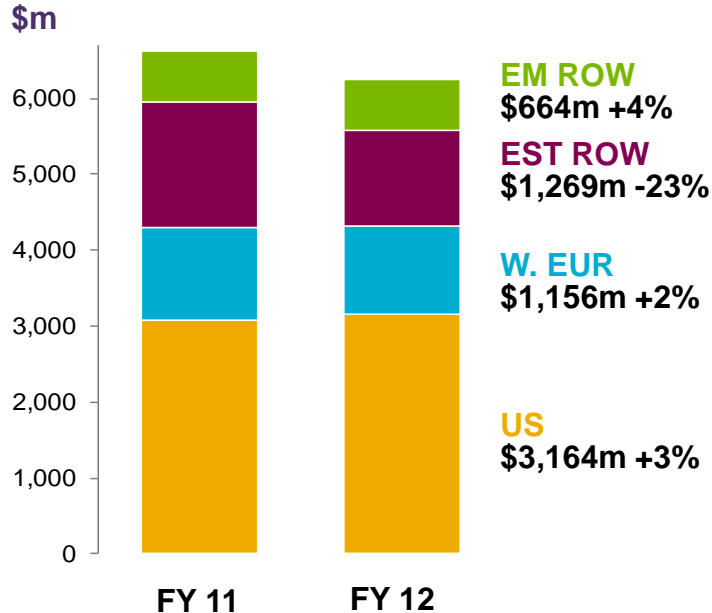
Crestor: US TRx trend

Resilient performance post multi-source generic atorvastatin



Crestor

2012 Sales: \$6,253m -4% CER



US

- FY TRx's -1.4%
- Resilient performance post generic atorvastatin

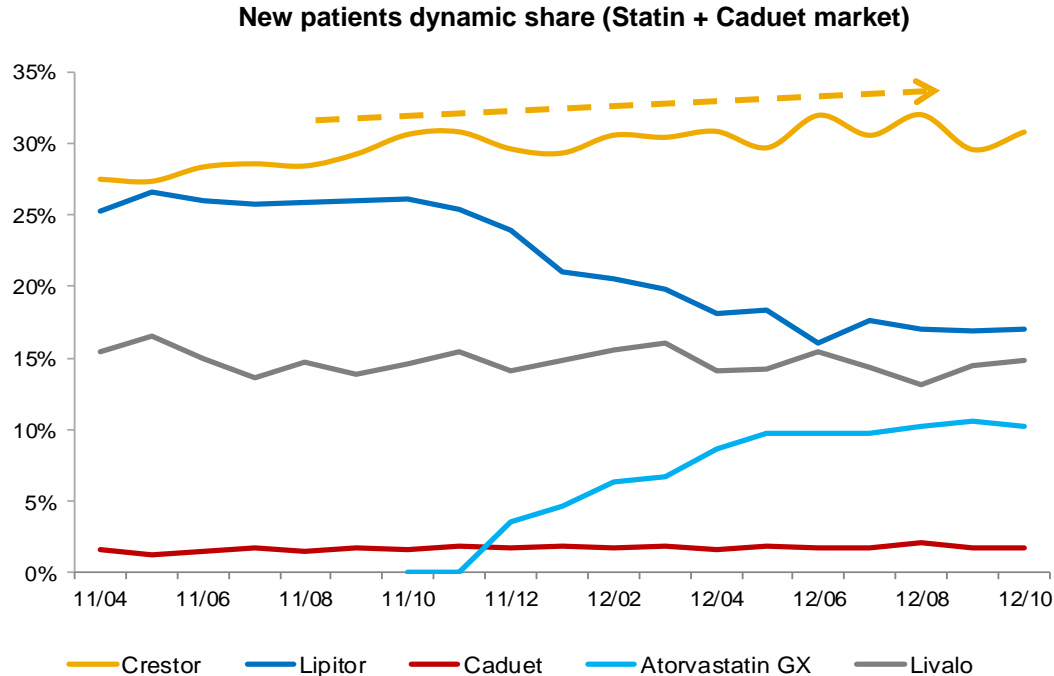
RoW

- RoW sales \$3,089m -9%
 - Adj for LOE in Canada, FY unchanged
- Crestor #1 statin in Japan
- Emerging Markets +4%
 - Adj for LOE Brazil/Mexico, +14%



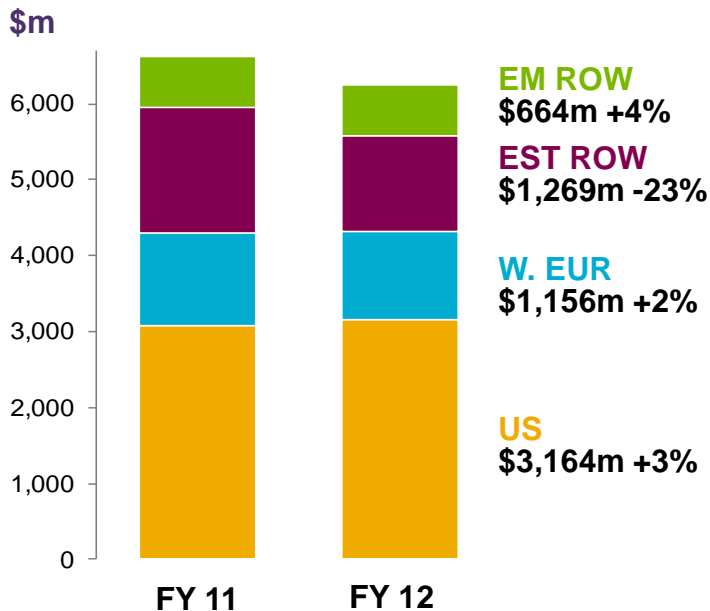
Crestor: Japan

Increasing share of new patients post generic atorvastatin launch



Crestor

2012 Sales: \$6,253m -4% CER



US

- FY TRx's -1.4%
- Resilient performance post generic atorvastatin

RoW

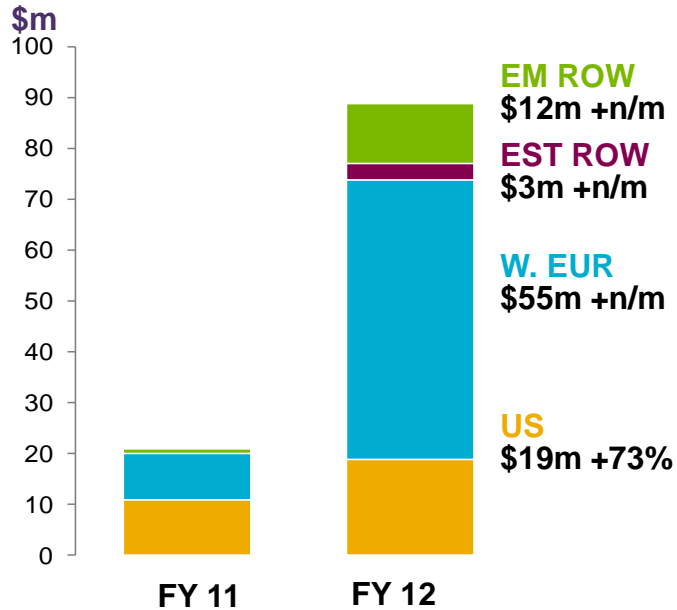
- RoW sales \$3,089m -9%
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Brilinta

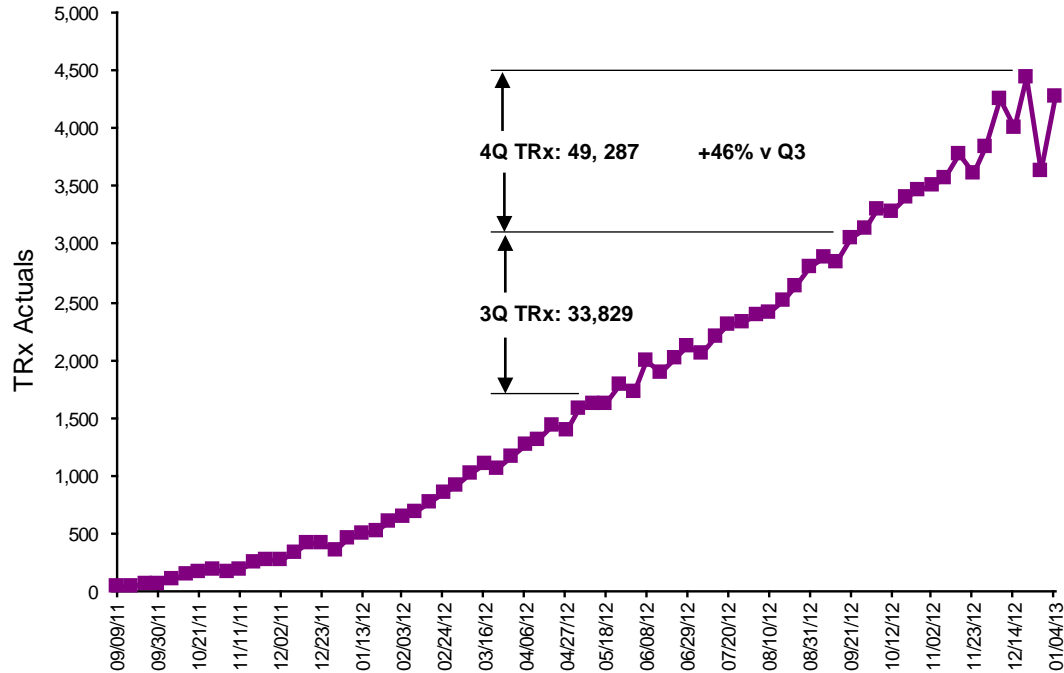
FY 2012 Sales: \$89m +348% CER

Now launched in 82 countries



Brilinta: US Launch progress

Steady increase in TRx's



On formulary in more than 60% of Top 400 hospitals

Preferred unrestricted access increased to >50% in Commercial & Medicare Part D managed care plans

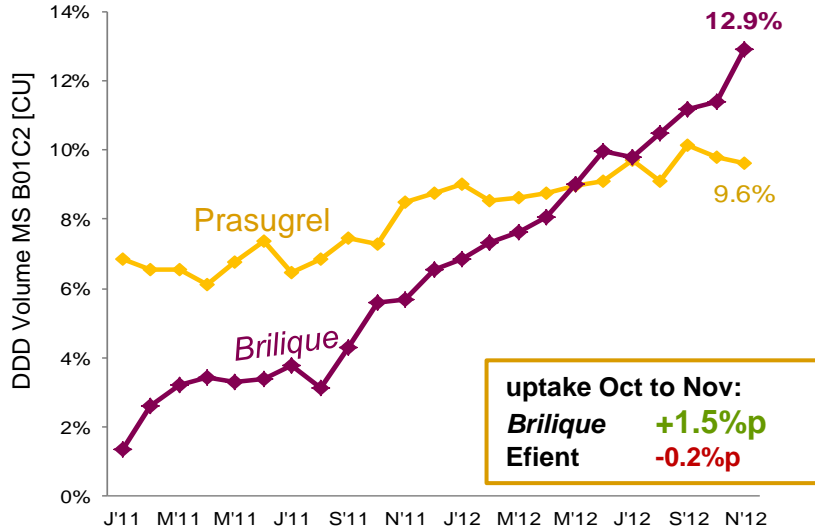


Brilique: Germany

Target hospitals: On protocol in 86%; 39% share of ACS initiations (#1)

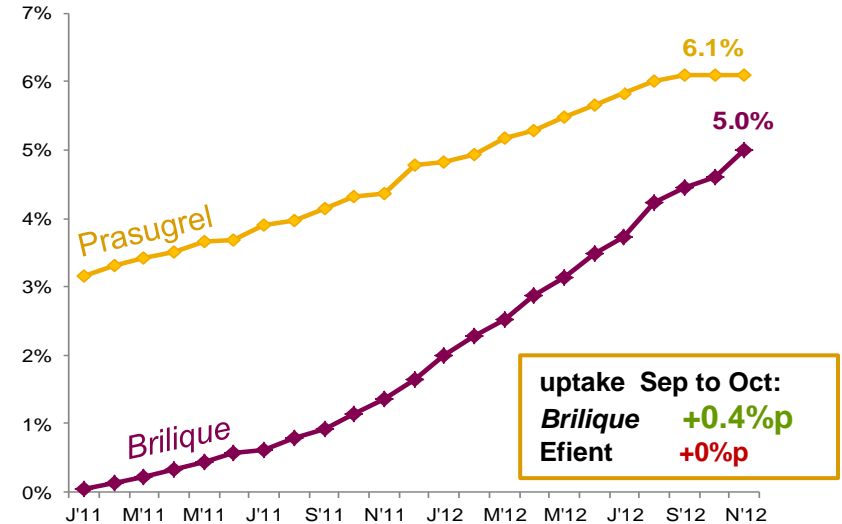
OAP Hospital Volume market share

(Daily Dosage)*



OAP Retail Volume market share

(Daily Dosage) *



*market definition: B01C2+Duoplavin+Duocover

Source: IMS DKM national (hospital) and Pharmascope national (retail)

Adjusted to daily dosage:
 Brilique: 2 Counting Units (CU) per day
 Other OAPs: 1 CU per day

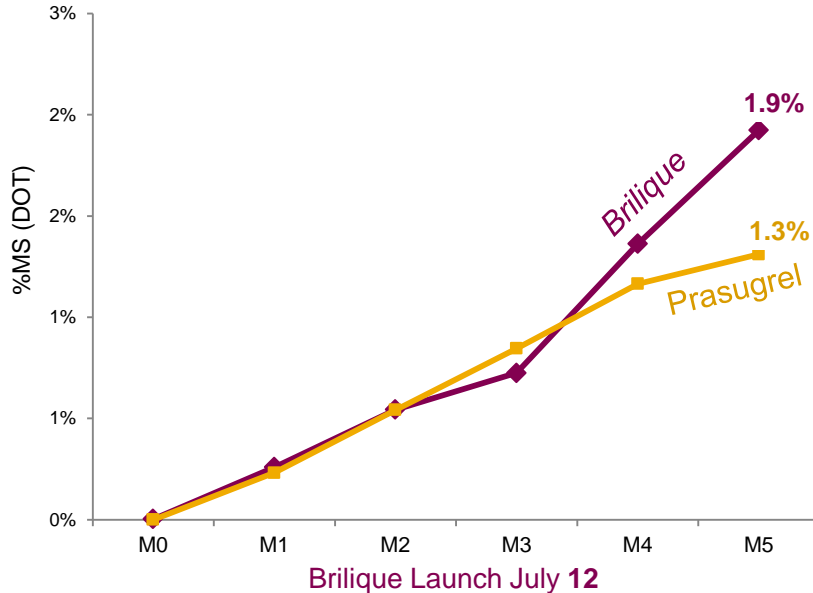


Brilique: France tracking well vs prasugrel launch

Target hospitals: On protocol in 76%; 29% share of ACS initiations (#2)

OAP Hospital Volume market share

Hospital Pharmacy Consumption⁽¹⁾

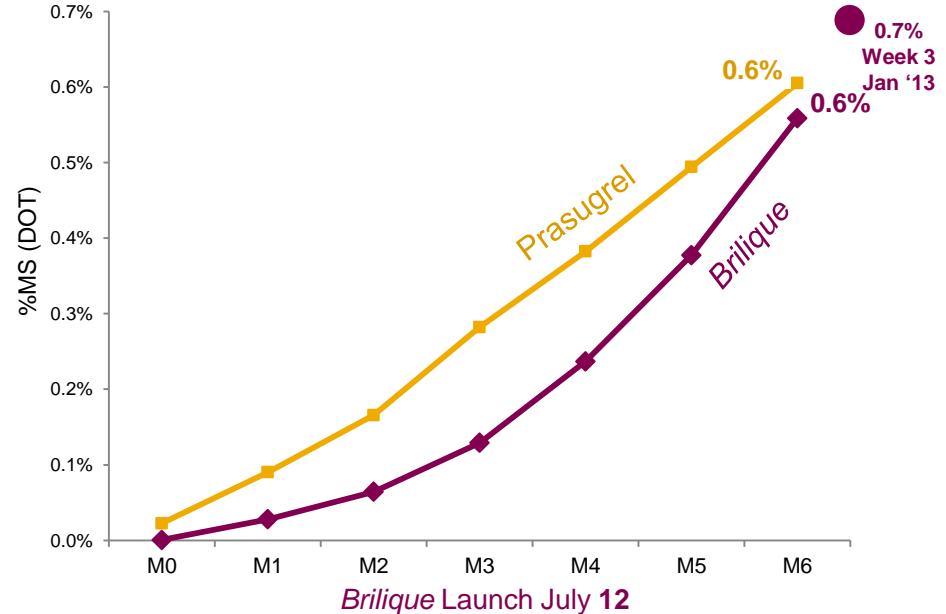


Phased on first month of sales :
M0 Efient = Dec 09 ; M0 Brilique = June 12

(1) Source: IMS EHPP November 12 (B01C2 OAP market)

OAP Retail Volume market share

sell-out ⁽²⁾



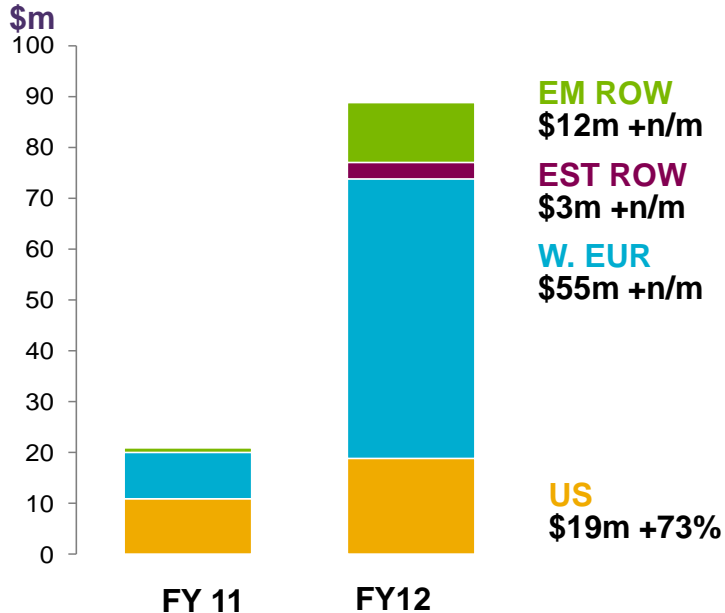
Phased on first month of sales :
M0 Efient = Jan 10 ; M0 Brilique = June 12

(2) Source: IMS December 12 (B01C2 OAP market)



Brilinta

FY 2012 Sales: \$89m +348% CER



Now launched in 82 countries

US

- Steady progress in TRx growth
- Preferred, unrestricted access in Commercial and Medicare Part D plans now >50%

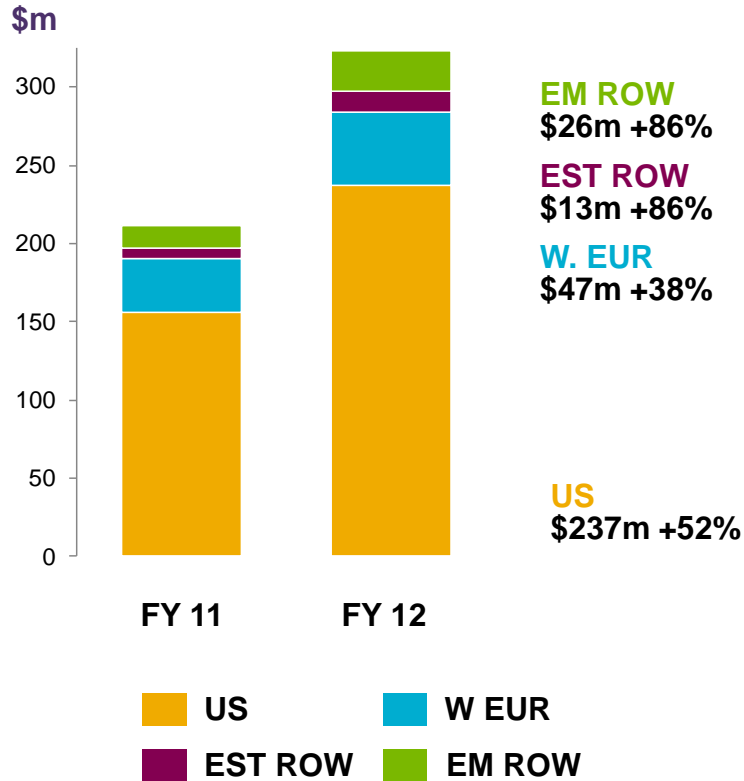
RoW

- Continued good performance in Germany
- Good launch uptake in France
- China approval Q4 2012



Onglyza Franchise

FY 2012 Global Alliance Revenue: \$323m +53% CER



US

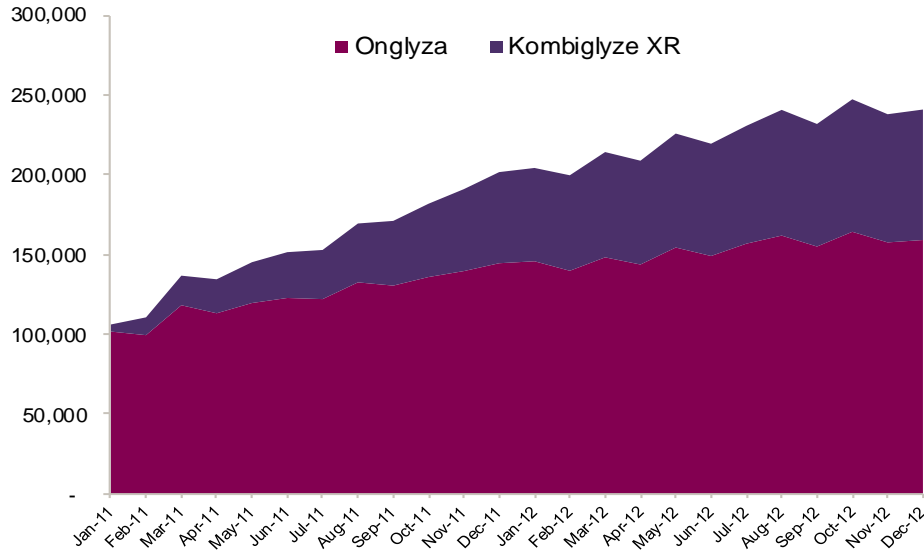
- Onglyza franchise TRx volume + 45%
 - DPP4 market growth +22%
- Onglyza franchise TRx share 17.8% in December 2012, +1.3 pts
 - Onglyza TRx share stable at 11.8%
 - *Kombiglyze XR* share 6.0%



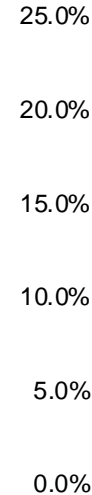
Onglyza: US TRx share increased in growing DPP4 market

Onglyza franchise TRx volume + 45%; TRx share +1.3 pts

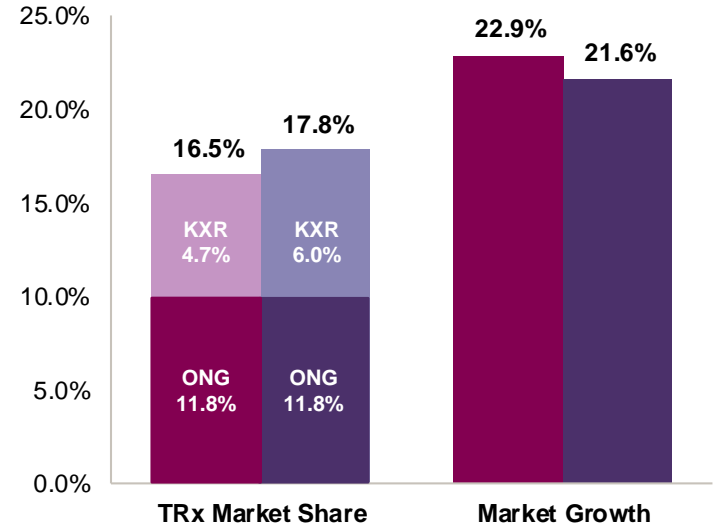
Onglyza franchise TRx volume



Onglyza franchise TRx share



DPP4 Market TRx growth

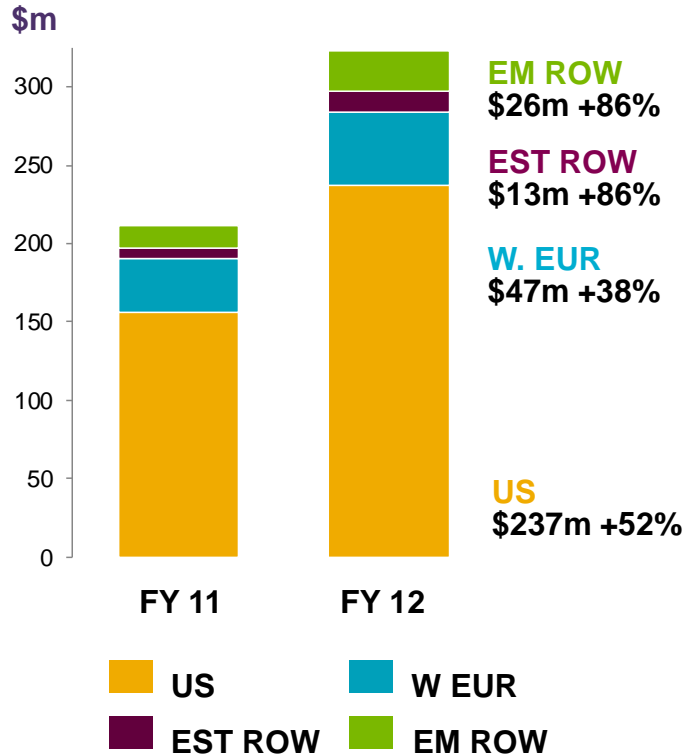


■ 2011 ■ 2012



Onglyza Franchise

FY 2012 Global Alliance Revenue: \$323m +53% CER



US

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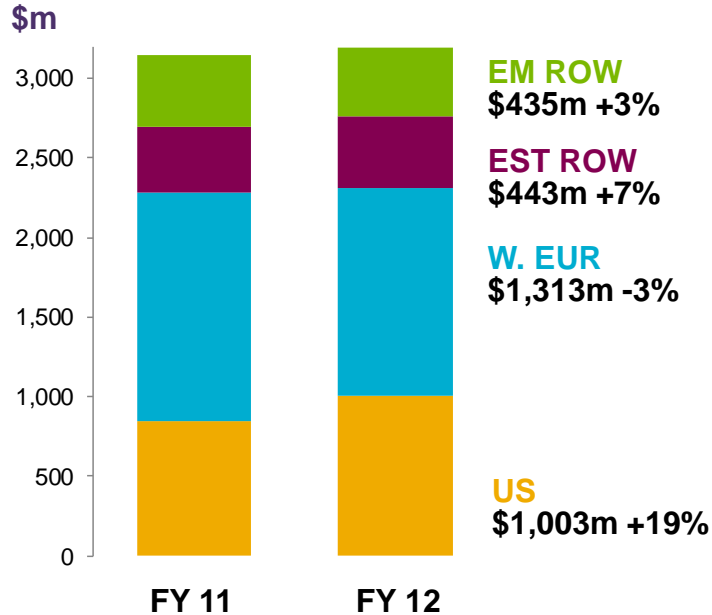
RoW

- RoW sales \$86m, +56%
- *Kombiglyze XR* launched in Brazil and Mexico
- *Komboglyze* launched in Spain, Germany and France in Q4 2012



Symbicort

2012 Sales: \$3,194m +5% CER



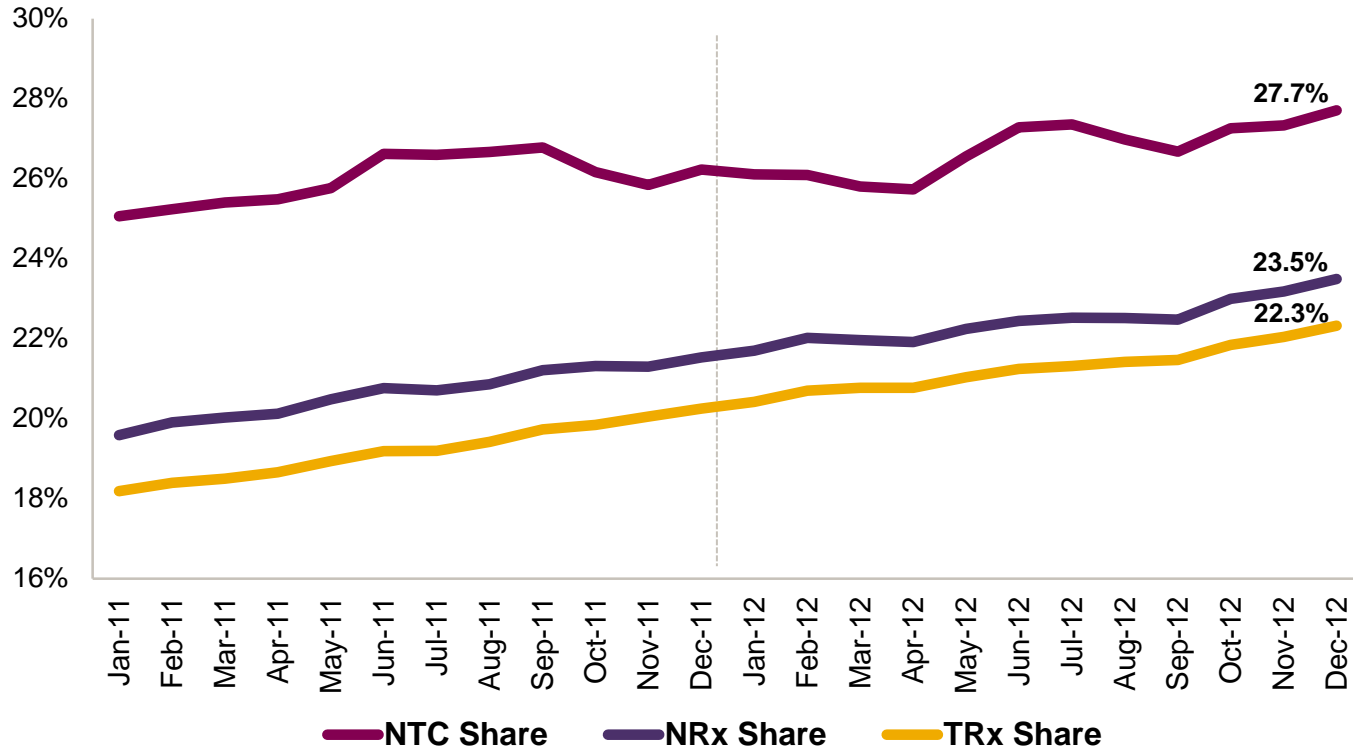
US

- US sales reach \$1bn milestone for 1st time



Symbicort: US

Steady increase in TRx and New to Combination (NTC) market share



Source for TRx, NRx figures: IMS NPA monthly – December 2012

Source for NTC figures: IMS NPA MD monthly – December 2012

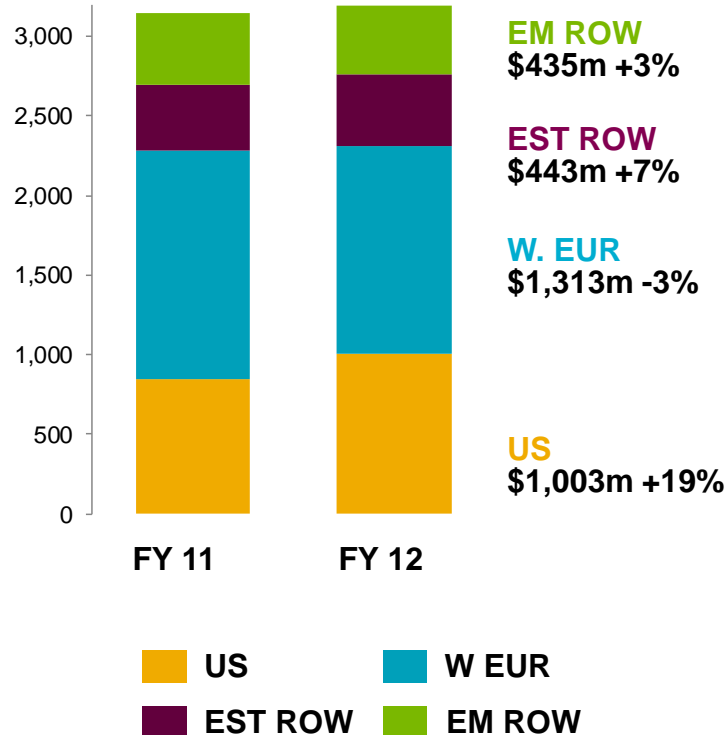
*Average share 2012 compared to average share 2011



Symbicort

2012 Sales: \$3,194m +5% CER

\$m



US

- US sales reach \$1bn milestone for 1st time
- *Symbicort* TRx's +13% vs +1% for fixed combination market
- December 2012 TRx share: 22.3% (+2 pts)
 - New patient share 27.7%

ROW

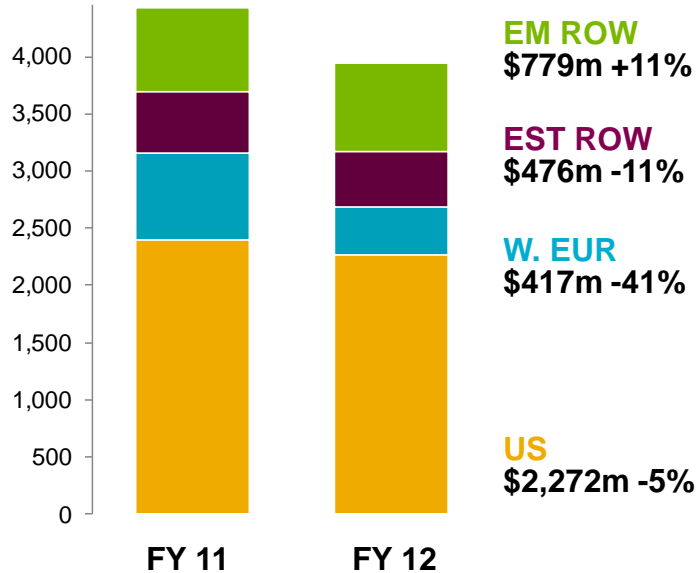
- FY sales unchanged at \$2,191m
- Good market share growth in Japan
 - *Symbicort* SMART and COPD approvals



Nexium

2012 Sales: \$3,944m, -10% CER

\$m



US

- Steady financial performance in highly generic market
- Volume decline partially offset by higher realised prices due to mix effect

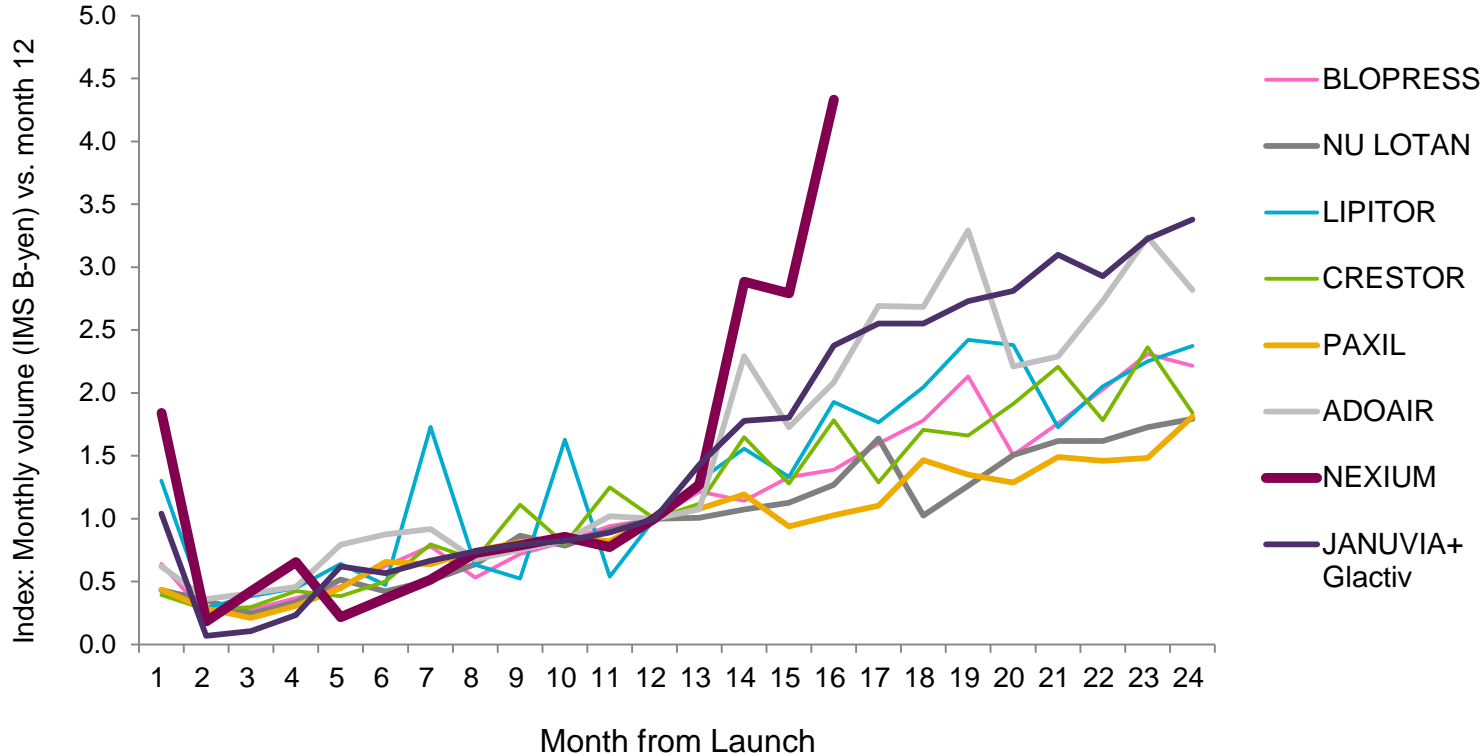
ROW

- FY sales \$1,672m, -15%
- LOE in Western Europe
- Strong performance in Japan following lifting of 2-week supply limit within Ryotanki regulations



Nexium Benchmark

Monthly Volume Trend from Launch



Full Year Results 2012

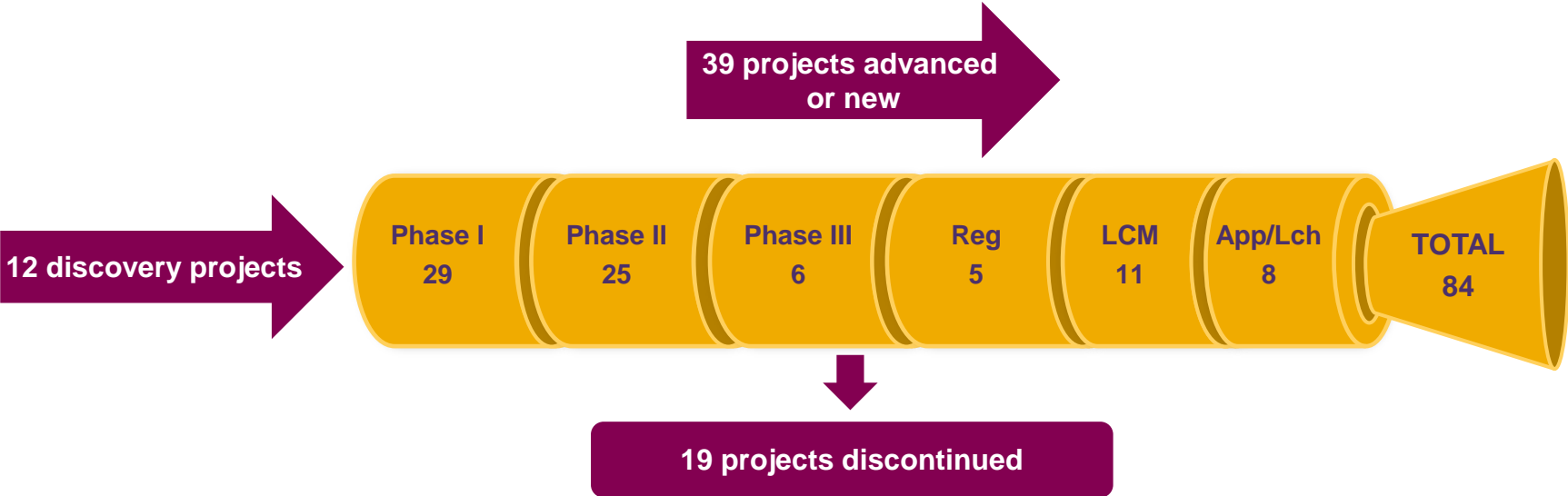
Pipeline review 2012

**Briggs Morrison, Executive Vice President
Global Medicines Development**

Driving strategic priorities



Pipeline movement since 2011



R&D pipeline – small and large molecule

	moxetumomab pasudotox (CD22) haematological malignancies
	MEDI0639 (DLL-4) solid tumours
AZD5363 (AKT) solid tumours	MEDI3617 (Ang2) solid tumours
AZD2014 (mTOR1&2) solid tumours	MEDI-565 (CEA BiTE) solid tumours
voltinib (MET) solid tumours	MEDI6469 (mOx40) solid tumours
AZD1208 (PIM) haematological malignancies	MEDI4736 (PD-L1) solid tumours
AZD9150 (STAT3) haematological malignancies	MEDI4212 (IgE) asthma
AZD8330 (MEK) solid tumours	MEDI7814 (C5/C5a) COPD
AZD8848 (ITLR7) asthma	MEDI2070 (IL-23) Crohn's
AZD7594 (ISGRM) COPD	MEDI9929 (TSLP) asthma
AZD1772 (NHE3) end stage renal disease	MEDI5872 (B7RP1) SLE
AZD1446 (a4b2 NNR) Alzheimer's	MEDI5117 (IL-6) rheumatoid arthritis
AZD3293 (Beta secretase) Alzheimer's	MEDI-557 RSV prophylaxis
ATM AVI (BL/BLI) targetted bacterial infection	MEDI-559 RSV prophylaxis
RDEA3170 (URAT1) gout	MEDI-550 (PIV) pandemic influenza prophylaxis

Small

Large

selumetinib (MEK) solid tumours	
AZD4547 (FGFR) solid tumours	MEDI-575 (PDGFRα) NSCLC
olaparib (PARP) gBRCA ovarian cancer	MEDI-551 (CD19) haematological malignancies
AZD5423 (ISGRM) COPD	tremelimumab (CTLA-4) solid tumours
AZD5069 (CXCR2) asthma	MEDI-573 (IGF) MBC
AZD2115 (MABA) COPD	benralizumab (IL-5R) asthma/COPD
AZD3241 (MPO) Parkinson's disease	mavrilimumab (GM-CSFR) rheumatoid arthritis
AZD6765 (NMDA) MDD	MEDI8968 (IL-1R) COPD
AZD3480 (a4b2 NNR) Alzheimer's	sifalimumab (IFNα) SLE
AZD5213 (H3R) Alzheimer's	tralokinumab (IL-13) asthma/IPF
CXL (BLI /cephalosporin) MRSA	MEDI-546 (IFNα) SLE
AZD5847 (Oxazolidine antibacterial inhibitor)TB	MEDI7183 (α4β7) ulcerative colitis, Crohn's

Small

Large

Legend

Oncology
Respiratory & Inflamm. (inc. autoimmune.)
CVGI
Neuroscience
Infection (and vaccines)
AMGEN/ARDEA

Caprelsa (VEGFR/EGFR) medullary thyroid cancer
fostratinib (SYK) rheumatoid arthritis
Forxiga (SGLT2) type 2 diabetes
Briiinta (ADP) arterial thrombosis
metreleptin (leptin analogue) lipodystrophy
naloxegol (oral peripherally-acting mu-opioid antagonist) Opioid-induced constipation
Zinforo (cephalosporin) pneumonia/skin infections
CAZ AVI (BLI /cephalosporin) serious infections
lesinurad (URAT1) gout

Small

Faslodex 1 st line advanced breast cancer (FALCON)
Iressa treatment beyond progression (IMPRESS)
Symbicort Breath Actuated Inhaler asthma/COPD
Birilinta outcomes study in PAD (EUCLID)
Birilinta outcomes study MI (PEGASUS-TIMI 54)
Bydureon outcomes study (EXSCEL)
Bydureon Dual Chamber Pen
Forxiga high CV risk
Forxiga triple therapy
SaxaDapa diabetes
Onglyza outcomes study (SAVOR-TIMI 53)

FluMist Quadrivalent Q/LAIV flu vaccine
brodalumab (IL-17R) psoriasis

Large

Phase I

LCM/parallel indications in different therapy area to lead *

MEDI-551
(anti-CD19 MAb) multiple sclerosis

Phase II

fostratinib
(SYK) haematological malignancies

tralokinumab
(IL-13) ulcerative colitis

Phase III – Registration

11 further LCM opportunities

* Project count excludes 8 LCM projects under first regulatory review or approved/launched in at least one major market ; excludes Symbicort pMDI LABA postmarketing safety study



Pipeline progress in 2012

Programme	Indication	2012 achievement	Timing
Caprelsa	medullary thyroid cancer	Launch in Europe	H1
Symbicort SMART	asthma	Launch in Japan	
Nexium low dose aspirin	prevention of recurrence of peptic ulcer	Launch in Japan	
Zinforo	skin and soft tissue infections and community acquired pneumonia	Launch in Europe	H2
Symbicort Turbuhaler	chronic obstructive pulmonary disease	Launch in Japan	
Oxis	chronic obstructive pulmonary disease	Launch in Japan	
Forxiga	type 2 diabetes	Launch in Europe	
Komboglyze	type 2 diabetes	Launch in Europe	H1
FluMist Quadrivalent	flu vaccine	Approval in US	
Brilinta	acute coronary syndrome	Approval in China	
FluMist Quadrivalent	flu vaccine	Regulatory submission in Europe	H2
Forxiga/metformin (fixed dose combination)	type 2 diabetes	Regulatory submission in Europe	
Forxiga	type 2 diabetes (add-on to dipeptidyl peptidase-4)	Regulatory submission in Europe	
Forxiga	type 2 diabetes (add-on to insulin)	Regulatory submission in Europe	
Nexium	H. pylori gastritis	Regulatory submission in Japan	
CAZ AVI	serious infections	Phase III start	H1
saxagliptin/dapagliflozin (fixed dose combi.)	type 2 diabetes	Phase III start	
Iressa	treatment beyond progression	Phase III start	
brodalumab	psoriasis	Phase III start	H2
Brilinta	peripheral artery disease	Phase III start	
Nexium	severe reflux oesophagitis	Phase III start (Japan)	
Faslodex	first line metastatic breast cancer	Phase III start	



Late stage – launch products

BRILINTA

Acute coronary syndrome

- Increased investment in PARTHENON programme
- BRILINTA included in 11 sets of global guidelines as part of the standard of care

FORXIGA

Adults with type 2 diabetes

- First in class SGLT2, now launched in UK, Germany and Denmark
- DECLARE cardiovascular outcomes trial will begin to enrol patients in Q2

ONGLYZA

Adults with type 2 diabetes

- SAVOR-TIMI 53 study – 16,500 patients completed in 19 months, two years ahead of schedule
- Complies with new FDA T2DM guidance regarding long-term CV risk

SYMBICORT

Asthma and COPD

- PATHOS real world evidence study compliments strong existing clinical database
- In Japan, SMART launched in June and COPD launched in September



Late stage – phase III

NALOXEGOL

Opioid induced constipation

- Top-line results from KODIAC-04 and KODIAC-05, and the KODIAC-07 safety extension trial
- On track for regulatory submissions in US, EU, and Canada mid-2013

CAZ AVI

Serious infections

- Innovative combination of antibiotic ceftazidime and betalactamase inhibitor avibactam
- Developed for treatment of hospitalised patients with serious Gram negative bacterial infections

BRODALUMAB

Psoriasis

- Phase III initiated in Q3, 2012
- In phase II, high level of response in patients with moderate to severe plaque psoriasis

FOSTAMATINIB

Rheumatoid arthritis

- First oral kinase inhibitor with selectivity for spleen tyrosine kinase in development for rheumatoid arthritis
- Phase III OSKIRA programme on track to report in Q2, with anticipated filings in the US and EU in Q4

LESINURAD

Gout

- Phase III programme started in early 2012, all four studies are actively enrolling
- Mechanism of action and tolerability profile could fundamentally change the treatment of gout



What's ahead – potential NME phase III starts 2013-14

2013	2014	
benralizumab – asthma	AZD6765 – depression	ATM AVI – serious infections
olaparib – solid tumours	sifalimumab – systemic lupus erythematosus	AZD4547 – solid tumours
moxetumomab pasudotox – hairy cell leukaemia	mavrilimumab – rheumatoid arthritis	AZD1722 – end stage renal disease
selumetinib – non-small cell lung cancer	MEDI-551 – haematological malignancies	MEDI7183 – ulcerative colitis
CXL – methicillin-resistant Staphylococcus aureus	MEDI-575 – non-small cell lung cancer	AZD5069 – asthma
	tralokinumab – asthma	

2013/14 phase III starts for lead indications only



Expected news flow highlights 2013

	Product	Milestone	Timing
NME	fostamatinib – rheumatoid arthritis	Phase III data	H1
	naloxegol – opioid-induced constipation	Phase III data (KODIAC-08)	
	moxetumomab pasudotox – hairy cell leukaemia	Potential phase III start	H2
	olaparib – solid tumours	Potential phase III start	
	CXL – methicillin-resistant Staphylococcus aureus	Potential phase III start	
	benralizumab – asthma	Potential phase III start	
	selumetinib – non-small cell lung cancer	Potential phase III start	
	naloxegol – opioid-induced constipation	Submission for approval in USA & Europe	
fostamatinib – rheumatoid arthritis	Submission for approval in USA & Europe		
Other	Forxiga – type 2 diabetes (triple therapy)	Submission for approval in Europe	H1
	Brilinta – acute coronary syndrome	Submission for approval in Japan	
	Forxiga – type 2 diabetes	Submission for approval in Japan & China	
	Zoladex – three month depot (breast cancer)	Submission for approval in Japan	H2
	Onglyza SAVOR-TIMI53 – outcomes study	Study results	
	brodalumab – psoriatic arthritis	Potential phase III start	
	Forxiga – type 2 diabetes	Resubmission for approval in USA	
	Onglyza SAVOR-TIMI53 – outcomes study	Submission for approval in USA & Europe	
	Bydureon Dual Chamber Pen – type 2 diabetes	Submission for approval in USA	
	FluMist Quadrivalent – flu vaccine	Launch in US	
Casodex oral dispersible tablet	Launch in Japan		



Driving strategic priorities



Full Year Results 2012

Full Year Results 2012

Simon Lowth, Chief Financial Officer

Headline results FY 2012

	2012 \$m	2011 \$m	Actual growth	CER growth
Revenue	27,973	33,591	-17%	-15%
Core Operating Profit	10,430	13,167	-21%	-18%
Core EPS	\$6.41	\$7.28	-12%	-9%
Restructuring	(\$0.94)	(\$0.63)		
MedImmune/Merck amortisation	(\$0.40)	(\$0.32)		
Intangible impairments	-	(\$0.01)		
Legal provisions & other	(\$0.08)	\$1.01*		
Reported EPS	\$4.99	\$7.33	-32%	-29%

* Includes \$1.08 gain from sale of Astra Tech



Headline results Q4 2012

	2012 \$m	2011 \$m	Actual growth	CER growth
Revenue	7,282	8,656	-16%	-15%
Core Operating Profit	2,532	2,990	-15%	-13%
Core EPS	\$1.56	\$1.61	-3%	+1%
Restructuring	(\$0.23)	(\$0.36)		
MedImmune/Merck amortisation	(\$0.11)	(\$0.08)		
Intangible impairments	-	-		
Legal provisions & other	-	(\$0.01)		
Reported EPS	\$1.22	\$1.16	+5%	+10%



Core margin: FY 2012

	\$m	CER growth	% sales	Delta vs PY CER
Revenue	27,973	-15%		
Core Gross Margin	22,716	-16%	81.2	-90bps
Distribution	(320)	-5%	1.2	-20bps
Core SG&A	(8,541)	-12%	30.5	-90bps
Core Other Income	1,027	+24%	3.7	+110bps
Core Pre-R&D Profit	14,882	-16%	53.2	-90bps
Core R&D	(4,452)	-11%	15.9	-70bps
Core Operating Profit	10,430	-18%	37.3	-160bps



Restructuring Programme: Phase 3 2012-2014

	Headcount Impact 2012-2014	Programme Cost 2012-2014 \$m	Progress to date	
			Costs \$m	Headcount
Global Supply Chain	1,350	(500)	(136)	~750
SG&A	3,750	(800)	(631)	~3,250
R&D	2,200	(800)	(791)	~2,260
Total	~ 7,300	(2,100)†	(1,558)*	~ 6,300

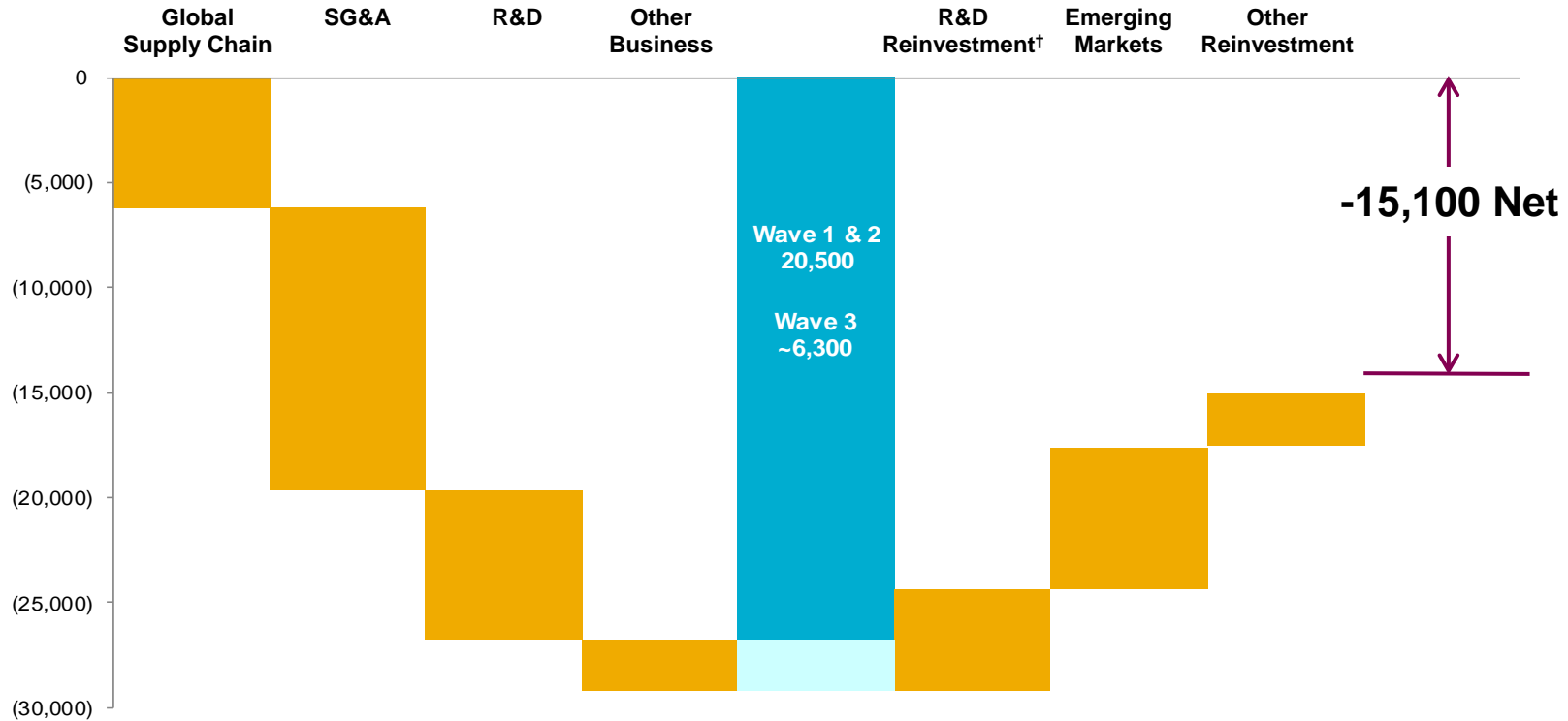
Estimated annual benefits \$1,600m by end 2014
of which ~\$350m has been realised by end 2012

* An additional \$261m was charged during Q4 2011

† Of which \$1.7bn are cash costs



Net headcount developments: 2006-2012



Headcount Dec 31st 2006 (66,800) Headcount Dec 31st 2012 (51,700)

† Includes the acquisition of MedImmune



Cash generation: FY 2012

	2012 \$m	2011 \$m
Opening net funds	2,849	3,653
EBITDA	10,666	15,345
Movement in working capital	(706)	(897)
Tax & interest paid	(2,588)	(4,547)
Other non-cash movements	(424)	(597)
Profit on disposal of Astra Tech	-	(1,483)
Net cash from operating activities	6,948	7,821



Cash application: FY 2012

	2012 \$m
Opening net funds	2,849
Net cash from operating activities	6,948
Purchased Amylin intangible assets	(3,360)
Other net capex	(1,060)
Dividends/Net share buy-back	(5,871)
Acquisitions	(1,187)
Other movements	312
Closing net funds	(1,369)
Gross debt	(10,310)
Cash/Cash equivalents, STIs & Net Derivative Financial Instruments	8,941



Cash distributions

Dividend

- FY 2012 dividend \$2.80 per share
- Progressive dividend policy confirmed (maintain or grow)
- Basis for dividend cover revised to 2x Core earnings (new definition)
- Cover likely to vary from the 2x cover target

Share repurchases

- FY 2012: Net \$2.2bn prior to suspension
- FY 2013: No repurchases planned



Core financials: FY 2012

	<i>Previous definition</i>		<i>New definition</i>	
	\$m	% sales	\$m	% sales
Revenue	27,973		27,973	
Core Gross Margin	22,716	81.2	23,041	82.4
Distribution	(320)	1.2	(320)	1.2
Core SG&A	(8,541)	30.5	(8,389)	30.0
Core Other Income	1,027	3.7	1,068	3.8
Core Pre-R&D Profit	14,882	53.2	15,400	55.0
Core R&D	(4,452)	15.9	(4,241)	15.1
Core Operating Profit	10,430	37.3	11,159	39.9
Core EPS	\$6.41		\$6.87	



Guidance for 2013 (new Core basis)

Revenue (CER)	Mid to High-single digit decline
Core Operating Costs (CER) (R&D + SG&A)	Hold to a slight increase vs 2012
Core Other Income	< \$600 million
Reported tax rate	Around 23 percent
Core EPS (CER) <i>(new definition)</i>	Decline significantly > revenue decline

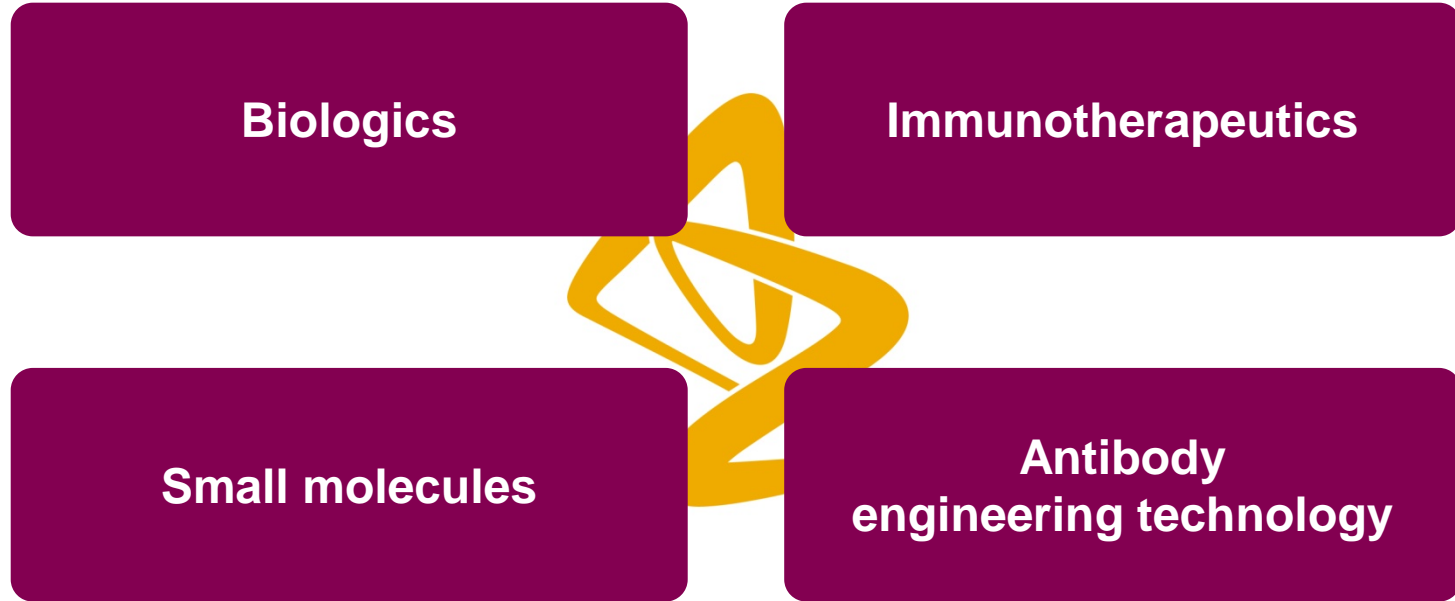


Full Year Results 2012

Full Year Results 2012

Pascal Soriot, Chief Executive Officer

Unique combination of capabilities



R&D pipeline – small and large molecule

	moxetumomab pasudotox (CD22) haematological malignancies
	MEDI0639 (DLL-4) solid tumours
AZD5363 (AKT) solid tumours	MEDI3617 (Ang2) solid tumours
AZD2014 (mTOR1&2) solid tumours	MEDI-565 (CEA BiTE) solid tumours
voltinib (MET) solid tumours	MEDI6469 (mOx40) solid tumours
AZD1208 (PIM) haematological malignancies	MEDI4736 (PD-L1) solid tumours
AZD9150 (STAT3) haematological malignancies	MEDI4212 (IgE) asthma
AZD8330 (MEK) solid tumours	MEDI7814 (C5/C5a) COPD
AZD8848 (ITLR7) asthma	MEDI2070 (IL-23) Crohn's
AZD7594 (ISGRM) COPD	MEDI9929 (TSLP) asthma
AZD1772 (NHE3) end stage renal disease	MEDI5872 (B7RP1) SLE
AZD1446 (a4b2 NNR) Alzheimer's	MEDI5117 (IL-6) rheumatoid arthritis
AZD3293 (Beta secretase) Alzheimer's	MEDI-557 RSV prophylaxis
ATM AVI (BL/BLI) targetted bacterial infection	MEDI-559 RSV prophylaxis
RDEA3170 (URAT1) gout	MEDI-550 (PIV) pandemic influenza prophylaxis

Small

Large

selumetinib (MEK) solid tumours	
AZD4547 (FGFR) solid tumours	MEDI-575 (PDGFRα) NSCLC
olaparib (PARP) gBRCA ovarian cancer	MEDI-551 (CD19) haematological malignancies
AZD5423 (ISGRM) COPD	tremelimumab (CTLA-4) solid tumours
AZD5069 (CXCR2) asthma	MEDI-573 (IGF) MBC
AZD2115 (MABA) COPD	benralizumab (IL-5R) asthma/COPD
AZD3241 (MPO) Parkinson's disease	mavrilimumab (GM-CSFR) rheumatoid arthritis
AZD6765 (NMDA) MDD	MEDI8968 (IL-1R) COPD
AZD3480 (a4b2 NNR) Alzheimer's	sifalimumab (IFNα) SLE
AZD5213 (H3R) Alzheimer's	tralokinumab (IL-13) asthma/IPF
CXL (BLI /cephalosporin) MRSA	MEDI-546 (IFNα) SLE
AZD5847 (Oxazolidine antibacterial inhibitor)TB	MEDI7183 (α4β7) ulcerative colitis, Crohn's

Small

Large

Legend

Oncology
Respiratory & Inflamm. (inc. autoimmune.)
CVGI
Neuroscience
Infection (and vaccines)
AMGEN/ARDEA

Caprelsa (VEGFR/EGFR) medullary thyroid cancer
fostratinib (SYK) rheumatoid arthritis
Forxiga (SGLT2) type 2 diabetes
Briiinta (ADP) arterial thrombosis
metreleptin (leptin analogue) lipodystrophy
naloxegol (oral peripherally-acting mu-opioid antagonist) Opioid-induced constipation
Zinforo (cephalosporin) pneumonia/skin infections
CAZ AVI (BLI /cephalosporin) serious infections
lesinurad (URAT1) gout

Small

Faslodex 1 st line advanced breast cancer (FALCON)
Iressa treatment beyond progression (IMPRESS)
Symbicort Breath Actuated Inhaler asthma/COPD
Birilinta outcomes study in PAD (EUCLID)
Birilinta outcomes study MI (PEGASUS-TIMI 54)
Bydureon outcomes study (EXSCEL)
Bydureon Dual Chamber Pen
Forxiga high CV risk
Forxiga triple therapy
SaxaDapa diabetes
Onglyza outcomes study (SAVOR-TIMI 53)

FluMist Quadrivalent Q/LAIV flu vaccine
brodalumab (IL-17R) psoriasis

Large

Phase I

LCM/parallel indications in different therapy area to lead *

MEDI-551
(anti-CD19 MAb) multiple sclerosis

Phase II

fostratinib
(SYK) haematological malignancies

tralokinumab
(IL-13) ulcerative colitis

Phase III – Registration

11 further LCM opportunities

* Project count excludes 8 LCM projects under first regulatory review or approved/launched in at least one major market ; excludes Symbicort pMDI LABA postmarketing safety study



Today: 5 key growth platforms



1. *Brilinta*



2. Diabetes



3. Emerging Markets



4. Respiratory



5. Japan



Full Year Results 2012