# 2Q and Half Year Results 2011



## 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 2011

**David Brennan, CEO** 



## **Key Developments 1H 2011**

- Pipeline
  - US FDA approves BRILINTA
  - US FDA Advisory Committee reviews dapagliflozin
- Sale of Astra Tech
  - Approx \$1.8 billion
  - Net proceeds added to share repurchases, on completion
- Market Environment



### **Headline results 1H 2011: Revenue**

	1H 2011 \$m	CER %	CER \$m
Total	16,722	-3	(489)
Crestor	3,192	+13	+368
Symbicort	1,554	+10	+143
Seroquel XR	726	+28	+154
Toprol-XL (US)	192	-55	(230)
Nexium (West Europe)	454	-29	(185)
Arimidex	414	-58	(553)
Merrem	330	-26	(110)
Casodex	271	-14	(42)



## Regional revenue performance 1H 2011

	1H 2011 \$m	CER %	CER \$m
Global Revenue	16,722	-3	(489)
US	6,596	-7	(499)
Western Europe	4,429	-8	(379)
Established RoW	2,797	+4	+100
Emerging Markets	2,900	+11	+289



### **Headline results 1H 2011**

	1H 2011 \$m	1H 2010 \$m	Actual growth	CER growth
Revenue	16,722	16,754	0%	-3%
Core R&D	(2,191)	(1,939)	+13%	+7%
Core SG&A	(4,977)	(4,583)	+9%	+5%



## **Headline results 1H 2011**

	1H 2011 \$m	1H 2010 \$m	Actual growth	CER growth
Revenue	16,722	16,754	0%	-3%
Core Operating Profit	7,000	7,507	-7%	-7%
Core EPS	\$3.96	\$3.82	+4%	+3%
Restructuring MedImmune/Merck amortisation Legal	(\$0.15) (\$0.16) (\$0.04)	(\$0.30) (\$0.14) (\$0.01)		
Reported EPS	\$3.61	\$3.37	+7%	+7%
Dividend	\$0.85	\$0.70		
Net Share Repurchases	\$2,204	\$516		



## 2Q and Half Year Results 2011

Simon Lowth, Chief Financial Officer

## **Headline results 2Q 2011**

	2Q 2011 \$m	2Q 2010 \$m	Actual growth	CER growth
Revenue	8,430	8,178	3%	-2%



## Core margin: 2Q 2011

	\$m	CER growth	% sales	Delta vs PY CER
Revenue	8,430	-2%		
Core Gross Margin	6,968	-2%	82.7	+10bps
Distribution	(88)	-8%	1.0	+10bps
Core SG&A	(2,627)	+9%	31.2	-310bps
Core Other Income	188	-2%	2.2	-
Core Pre-R&D Profit	4,441	-7%	52.7	<b>-290</b> bps
Core R&D	(1,119)	+8%	13.3	-110bps
Core Operating Profit	3,322	-10%	39.4	-400 bps



## **Headline results 2Q 2011**

	2Q 2011 \$m	2Q 2010 \$m	Actual growth	CER growth
Revenue	8,430	8,178	3%	-2%
Core Operating Profit	3,322	3,650	-9%	-10%
Core EPS	\$1.73	\$1.79	-3%	-5%



## **Headline results 2Q 2011**

	2Q 2011 \$m	2Q 2010 \$m	Actual growth	CER growth
Revenue	8,430	8,178	3%	-2%
Core Operating Profit	3,322	3,650	-9%	-10%
Core EPS	\$1.73	\$1.79	-3%	-5%
Restructuring MedImmune/Merck amortisation Legal	(\$0.08) (\$0.08) (\$0.04)	(\$0.25) (\$0.07) (\$0.01)		
Reported EPS	\$1.53	\$1.46	+5%	+3%



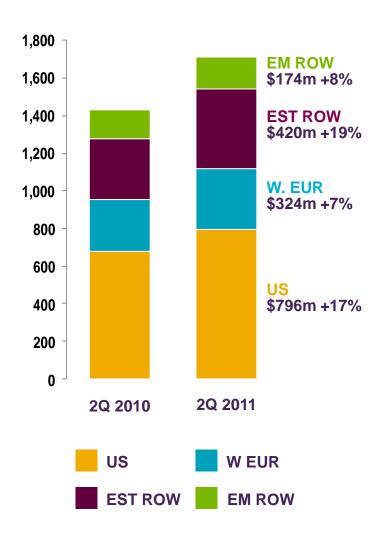
## Regional revenue performance 2Q 2011

	2Q 2011 \$m	CER %	CER \$m
Global Revenue	8,430	-2%	(139)
US	3,292	-3%	(106)
Western Europe	2,194	-9%	(208)
Established RoW	1,476	+4%	+48
Emerging Markets	1,468	+10%	+127



### **Crestor**

2Q 2011 Sales: \$1,714m +15%



#### US

- US TRx +2.4%
  - Statin market +1%
- Dynamic share (new & switch)
  - Crestor share up ~3 pts since simvastatin safety advisory

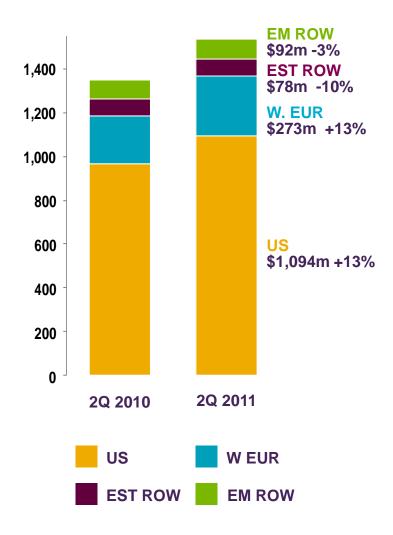
#### **RoW**

- RoW sales \$918m +12%
- Western Europe +7%
  - Volume growth double-digit
- Double-digit growth in Japan/Canada/Australia
- Generic rosuvastatin impacts
  Eastern Europe



## Seroquel

2Q 2011 Sales: \$1,537m +11%



Seroquel IR: \$1,150m +7%

Seroquel XR: \$387m +23%

#### US

- Seroquel XR TRx +19% vs market +4%
  - Seroquel XR now 16.9% of TRx & 18.7% of Revenue

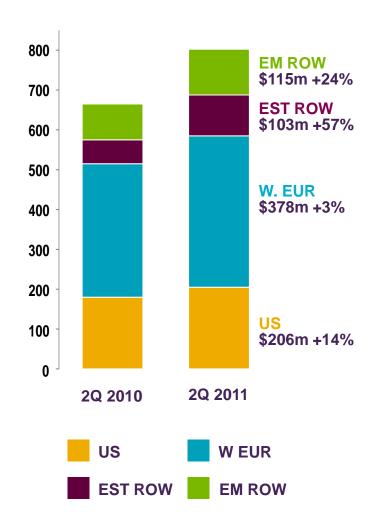
#### RoW

- Seroquel franchise sales \$443m +5%
- Seroquel XR sales +36%
  - Seroquel XR now 41% of franchise sales



## **Symbicort**

2Q 2011 Sales: \$802m +14%



#### US

- TRx +10% vs market -3%
- TRx share 19.2%
  - Up 2 pts vs June 2010
- New patient share 26.1%

#### **RoW**

• Sales \$596m +13%



### **Brand revenues 2Q 2011**

#### Onglyza™

- Global alliance revenue \$46m; US \$33m
- US franchise TRx share increased to 14.1% in June 2011
  - Up 4.1 pts since Dec 2010; Kombiglyze XR™ TRx at 2.7% share
- Franchise share of new DPP4 starts ~25%
  - Kombiglyze XR™ accounts for 36% of franchise starts

#### **BRILINTA**

- Sales of \$2m
- Protocol adoption in ~15% of 1000 target hospitals in Germany
  - Good trial rates



## Cash generation: 1H 2011

	2011 \$m	2010 \$m
Opening net cash/(debt)	3,653	535
EBITDA	7,403	7,509
Movement in working capital	(1,053)	(977)
Tax & interest paid*	(1,902)	(1,235)
Other non-cash movements	(236)	32
	4,212	5,329
Tax settlements*	(1,383)	(562)
Net cash from operating activities	2,829	4,767



<sup>\*</sup> Adjusted for Tax settlements

## Cash application: 1H 2011

	2011 \$m
Opening net cash/(debt)	3,653
Net cash from operating activities	2,829
Capex/Other investments	(629)
Dividends/Net share buy-back	(4,850)
Other movements	29
Closing net cash/(debt)	1,032
Gross debt	(9,582)
Cash/Cash equivalents and STIs	10,614



#### **Shareholder returns**

- Dividends
  - Progressive dividend
  - Balance between Interim and Final
  - First Interim \$0.85
  - Approx 33 percent of FY 2010 of \$2.55
- Share Repurchases
  - 2011 target: Net \$4 billion
  - Astra Tech net proceeds
  - New 2011 estimate: \$5 billion



## **Guidance for 2011 (Core Basis)**

Revenue Flat to low single digit at CER

• Core EPS New range \$7.05 to \$7.35; up 10 cents



## 2Q and Half Year Results 2011

Simon Lowth, Chief Financial Officer

## 2Q and Half Year Results 2011

Martin Mackay, R&D President

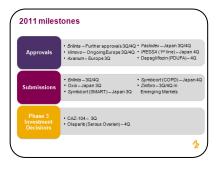
## **Agenda**



Pipeline overview



Late stage progress



Upcoming milestones



## Portfolio movement January – July 2011

#### Launched/Approved



onglyza

China

Nexium<sup>®</sup>

Japan

esomeprazole

Caprelsa

(vandetanib) Tablets

















#### **Submitted**





**USA** 



#### Dapagliflozin

Russia

Russia, Brazil

#### **New Indications**

### Renal impairment USA, Europe

#### **Phase 3 Starts**

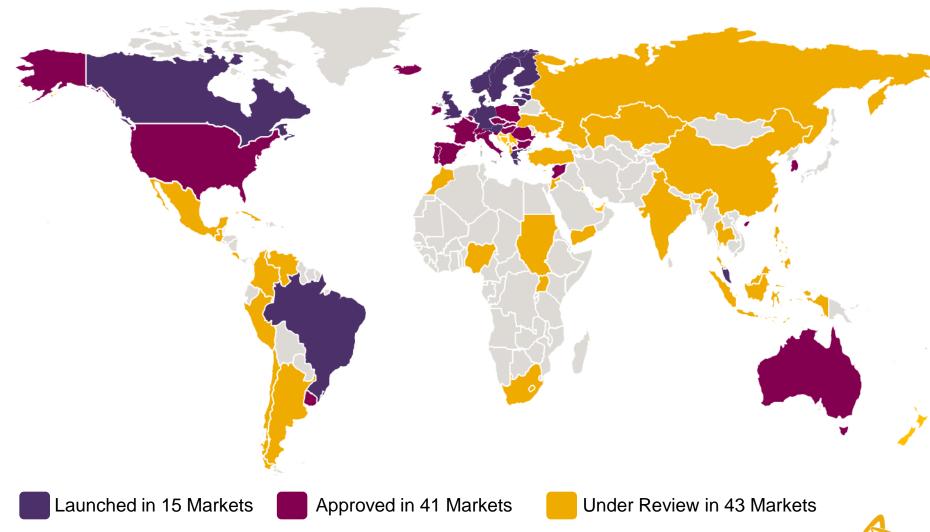
**NKTR-118** 



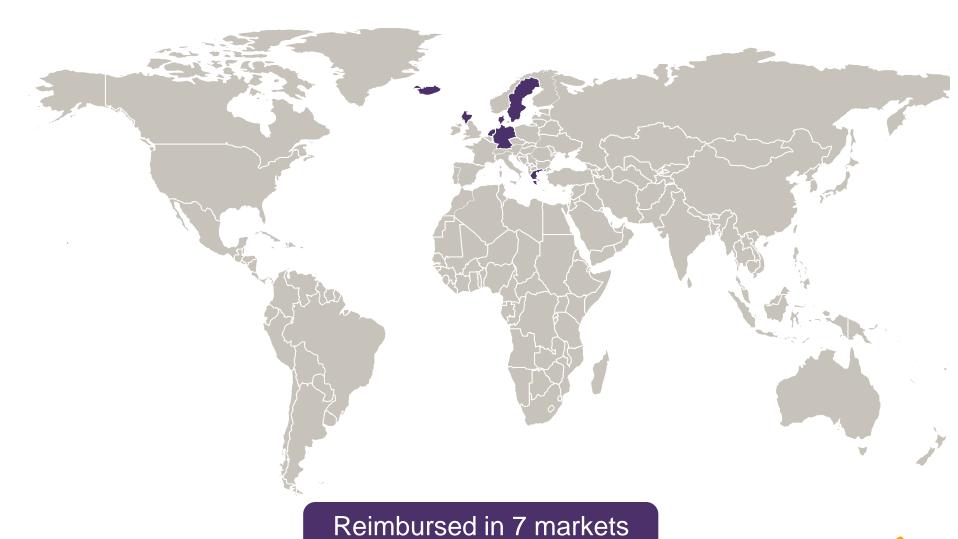




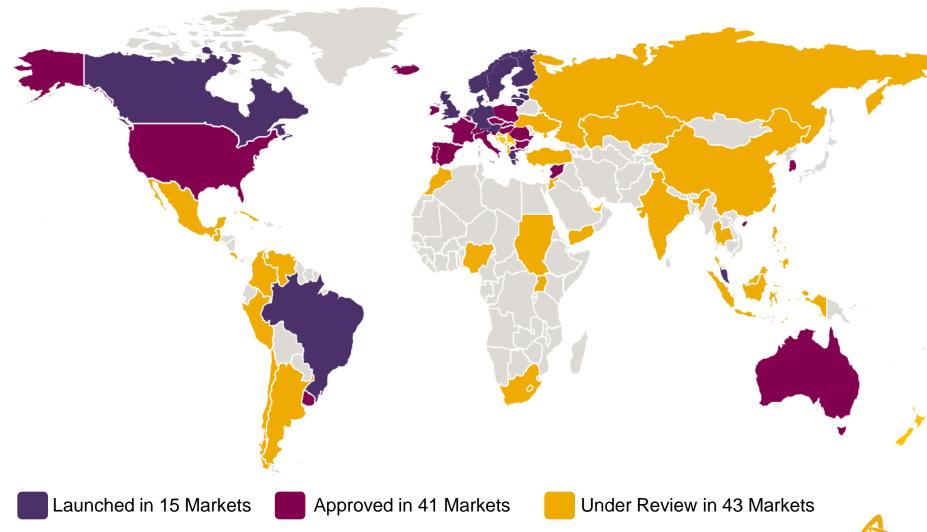
## **BRILINTA** regulatory status



## **BRILINTA** regulatory status



## **BRILINTA** regulatory status



#### BRILINTA label

#### Strong differentiating label

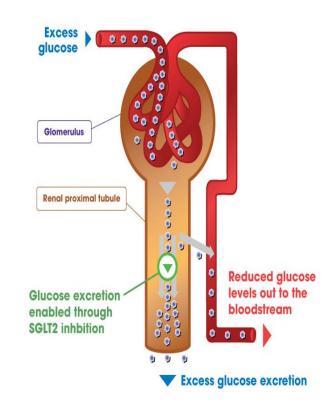
- Approved across the full continuum of ACS patients studied in PLATO
- Includes overall results of the PLATO study
  - *BRILINTA* compared to clopidogrel in combined endpoint of CV death, MI or stroke. The overall results favoured *BRILINTA* when used with low maintenance doses of aspirin
  - BRILINTA is the only oral antiplatelet to demonstrate a reduction in CV death in patients with ACS compared to clopidogrel
- Kaplan-Meier survival curve and clinical study section that describes the efficacy of BRILINTA versus clopidogrel over the 12-month treatment period
- Boxed warning for bleeding risks and the impact of aspirin dose on BRILINTA effectiveness



### Dapagliflozin

#### New approach to diabetes

- Committed to the broad clinical development programme
- Efforts are focused on working with FDA between now and October 28<sup>th</sup> PDUFA date to address outstanding questions





### NKTR-118, Fostamatinib, TC-5214 & Crestor

#### Additional late stage programmes are on track

- NKTR-118 First regulatory filing planned for 2013
- Fostamatinib On track to meet US/EU filing dates in 2013
- TC-5214 First study read out in 4Q11 and all study readouts in 2Q12
- Crestor/SATURN Scientific presentation expected at American Heart Association in 4Q11











#### 2011 milestones

#### **Approvals**

- Brilinta Further approvals 3Q/4Q
  Faslodex Japan 3Q/4Q
- Vimovo Ongoing Europe 3Q/4Q
- Axanum Europe 3Q

- IRESSA (1st line) Japan 4Q
- Dapagliflozin (PDUFA) 4Q

#### **Submissions**

- Brilinta 3Q/4Q
- Oxis Japan 3Q
- Symbicort (SMART) Japan 3Q
- Symbicort (COPD) Japan 4Q
- *Zinforo* 3Q/4Q in **Emerging Markets**

#### Phase 3 Investment Decisions

- CAZ-104 3Q
- Olaparib (Serous Ovarian) 4Q



# 2Q and Half Year Results 2011

