

AstraZeneca

Pascal Soriot, Executive Director, Chief Executive Officer
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14 January 2015



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:

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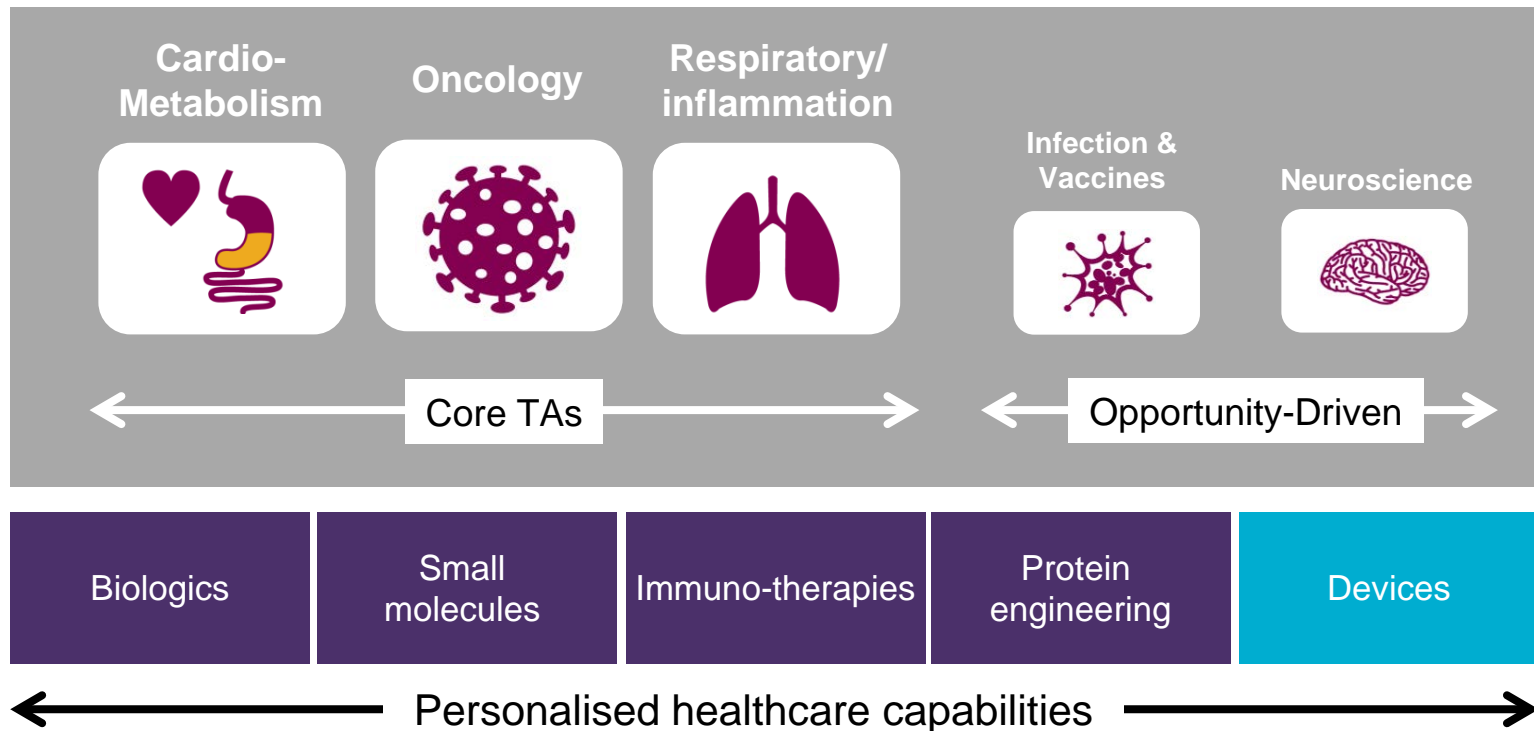
The forward-looking statements reflect knowledge and information available at the date of preparation of this presentation and AstraZeneca undertakes no obligation to update these forward-looking statements. We identify the forward-looking statements by using the words 'anticipates', 'believes', 'expects', '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.



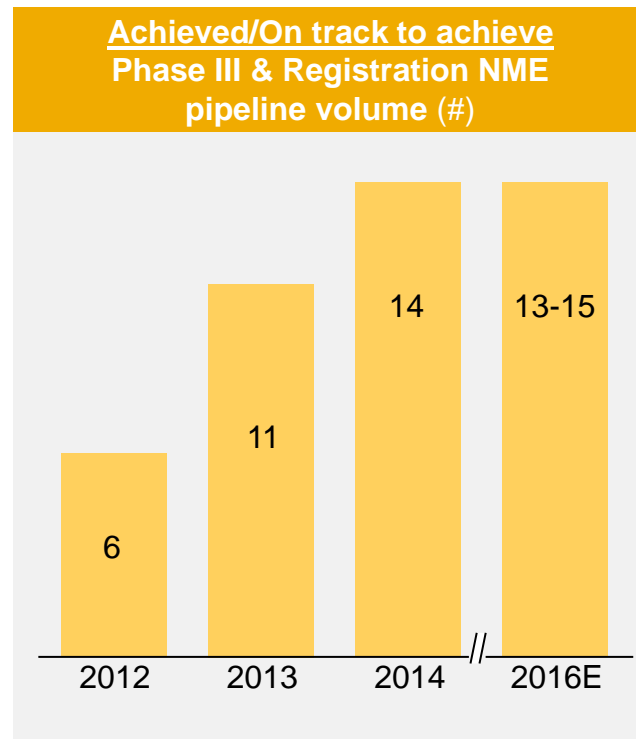
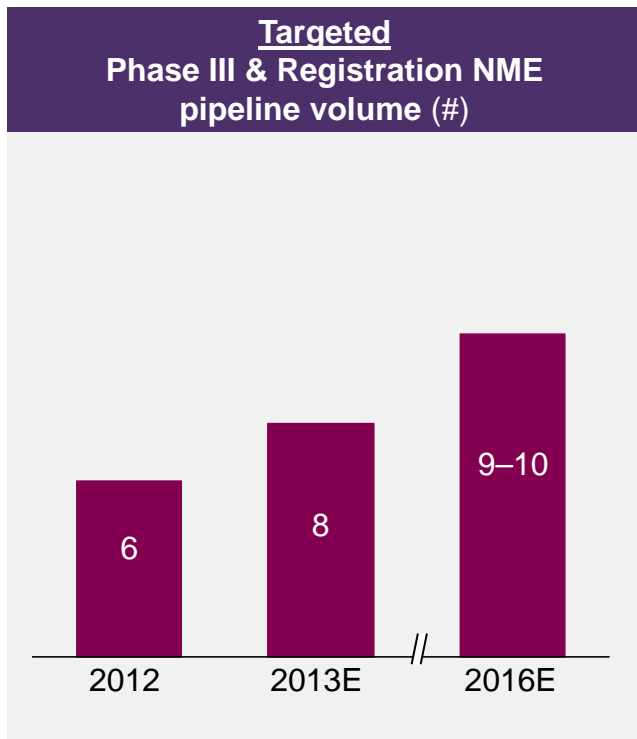
Completing the first phase of the journey



Achieve scientific leadership: R&D in 3 core therapeutic areas & across key platforms



Achieve scientific leadership: Late-stage pipeline volume well ahead of plan



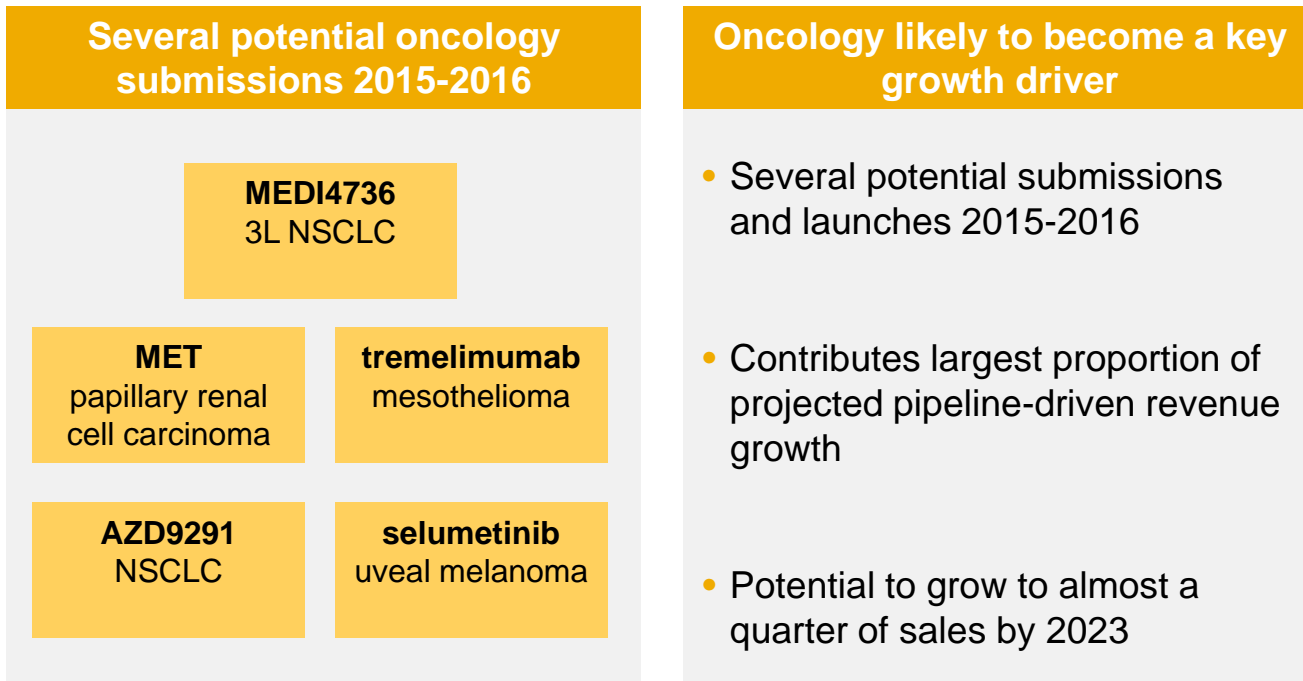
Return to growth: Upcoming submissions drive longer-term growth

LE submission opportunities				MEDI4736 + tremelimumab 2L SCCHN
			Faslodex 1L metastatic breast cancer	MEDI4736 2L SCCHN
			Brilinta stroke	Lynparza BRCAm metastatic breast cancer
		saxa/dapa FDC type 2 diabetes	brodalumab* psoriatic arthritis	Lynparza BRCAm PSR ovarian cancer (SOLO-2)
	Brilinta prior MI	Bydureon autoinjector	lesinurad FDC gout	Caprelsa differentiated thyroid cancer
NME submission opportunities		CAZ AVI serious infections	roxadustat CKD / ESRD (China)	MET papillary renal cell carcinoma
	brodalumab* psoriasis	selumetinib uveal melanoma	benralizumab severe asthma	tremelimumab mesothelioma
	PT003 (LAMA/LABA) COPD	AZD9291 NSCLC	PT001 (LAMA) COPD	MEDI4736 3L NSCLC
2015			2016	

*Partner Amgen to manage regulatory submission



Return to growth: Oncology will become sixth growth platform



Building a different shape of business



Late-stage pipeline: Key news flow through 2015

	Compound	Indication	Potential milestone
RIA	brodalumab	psoriasis	Regulatory submission
	PT003 (LAMA/LABA)	COPD	Phase III results & regulatory submission
	anifrolumab	SLE	Phase II presentation
	lesinurad	gout	Regulatory submission
CVMD	<i>Brilinta</i>	prior MI (PEGASUS)	Phase III results
	saxa/dapa FDC	type 2 diabetes	Regulatory submission
Oncology	<i>Lynparza</i>	PSR BRCAm ovarian cancer	Approval Phase III topline results (SOLO-2)
	AZD9291	2 nd line NSCLC	Regulatory submission
	MEDI4736 (PD-L1)	3 rd line NSCLC	Phase II/potential registration topline results
	MEDI4736 (PD-L1) / tremelimumab	NSCLC	Phase I presentation (ASCO)
	cediranib	ovarian cancer	Further analysis (ICON6)
Neuroscience	<i>Movantik/Moventig</i>	opioid-induced constipation	US de-scheduling & launch; EU approval



Key messages

- ✓ Strategy implementation on track
- ✓ Late-stage pipeline progressing ahead of plans
- ✓ Accelerating return to growth and ambition to become a >\$45bn company by 2023
- ✓ Building a sustainable, durable and more profitable business
- ✓ Strong 2015 news flow expected



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