

H1 and Q2 2025 Results

Conference call and webcast for investors and analysts

29 July 2025



Forward-looking statements

This document contains certain forward-looking statements with respect to the operations, performance and financial condition of the Group, including, among other things, statements about expected revenues, margins, earnings per share or other financial or other measures. Although the Group believes its 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 document and the Group undertakes no obligation to update these forward-looking statements. The Group identifies 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 the Group's control, include, among other things: the risk of failure or delay in delivery of pipeline or launch of new medicines; the risk of failure to meet regulatory or ethical requirements for medicine development or approval; the risk of failures or delays in the quality or execution of the Group's commercial strategies; the risk of pricing, affordability, access and competitive pressures; the risk of failure to maintain supply of compliant, quality medicines; the risk of illegal trade in the Group's medicines; the impact of reliance on third-party goods and services; the risk of failure in information technology or cybersecurity; the risk of failure of critical processes; the risk of failure to collect and manage data and artificial intelligence in line with legal and regulatory requirements and strategic objectives; the risk of failure to attract, develop, engage and retain a diverse, talented and capable workforce; the risk of failure to meet our sustainability targets, regulatory requirements and stakeholder expectations with respect to the environment; the risk of the safety and efficacy of marketed medicines being questioned; the risk of adverse outcome of litigation and/or governmental investigations; intellectual property risks related to the Group's products; the risk of failure to achieve strategic plans or meet targets or expectations; the risk of geopolitical and/or macroeconomic volatility disrupting the operation of our global business; the risk of failure in internal control, financial reporting or the occurrence of fraud; and the risk of unexpected deterioration in the Group's financial position. Nothing in this document, or any related presentation/webcast, should be construed as a profit forecast.



H1 and Q2 2025 Results

Conference call agenda

CEO Opening Remarks	Pascal Soriot Chief Executive Officer	
Financial Results	Aradhana Sarin Chief Financial Officer	
Oncology Haematology	Dave Fredrickson EVP, Oncology Haematology Business	Susan Galbraith EVP, Oncology Haematology R&D
BioPharmaceuticals	Ruud Dobber EVP, BioPharmaceuticals Business	Sharon Barr EVP, BioPharmaceuticals R&D
Rare Disease	Marc Dunoyer Chief Executive Officer, Alexion	
CEO Closing Remarks, Q&A	Pascal Soriot Chief Executive Officer	



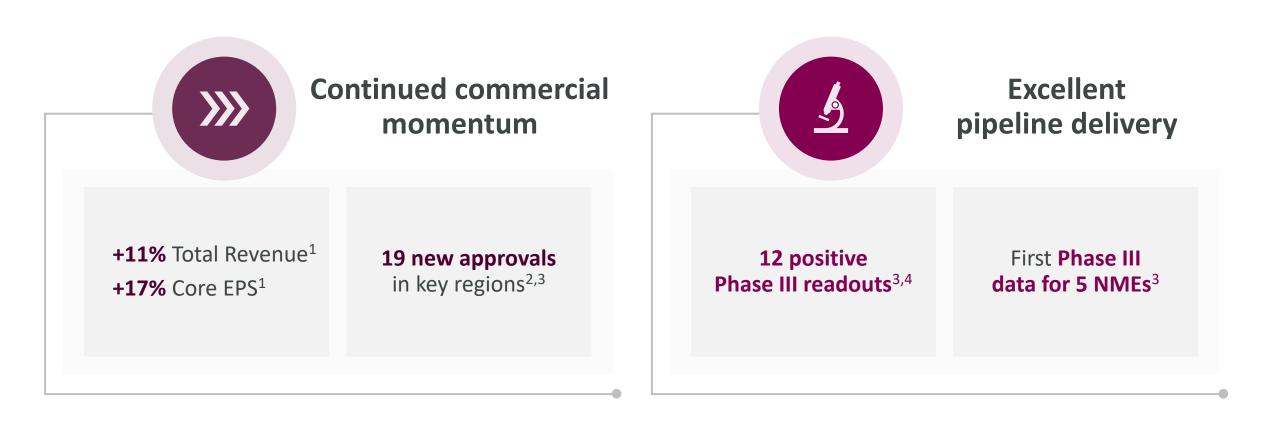


CEO Opening Remarks

Pascal Soriot
CHIEF EXECUTIVE OFFICER



Strong commercial and pipeline delivery in H1 2025

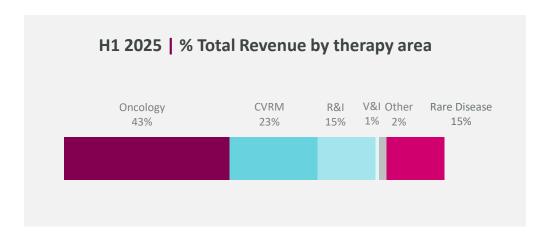


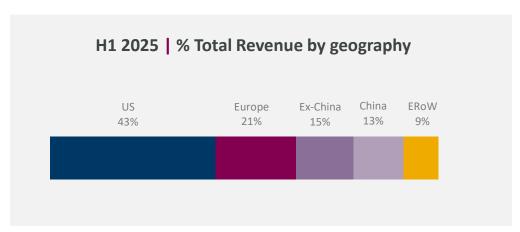


All growth rates at CER. 1. Growth rates relative to H1 2024 performance. 2. Key regions – US, EU, Japan, China. 3. Since FY 2024 Results to 29 July 2025. 4. Includes DESTINY-Breast09, DESTINY-Breast11, DESTINY-Gastric04, POTOMAC, SERENA-6, KALOS, LOGOS, NATRON, AZALEA, BaxHTN, CALYPSO, PREVAIL.

Appendix: Glossary.

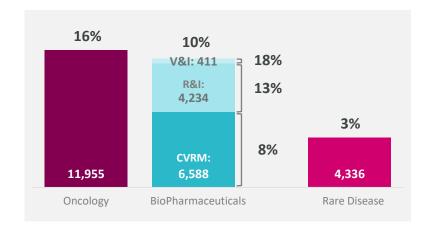
H1 2025 – growth supported by diverse, broad-based business





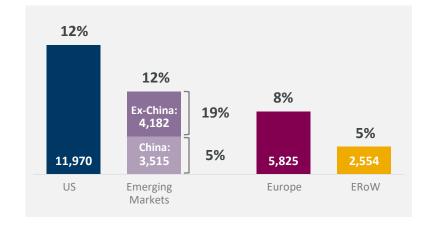
Strength across therapy areas

H1 2025 | Total Revenue (\$m)



Growth across geographies

H1 2025 | Total Revenue (\$m)





Excellent delivery across diverse pipeline

Key Phase III trial readouts in 2025 to date

Oncology

NME

SERENA-6

camizestrant potential new endocrine backbone in HR+ BC

MATTERHORN

Imfinzi transforming perioperative care in gastric/GEJ cancers

DESTINY-Breast09

Enhertu redefining management of 1L HER2+ mBC

POTOMAC

Imfinzi opportunity to treat earlier in bladder cancer

DESTINY-Breast11

Enhertu first move into early-stage HER2+ breast cancer

FLAURA2 OS

Tagrisso backbone SoC for 1L EGFRm NSCLC and beyond

BioPharmaceuticals

KALOS/LOGOS

Breztri broadening opportunity into uncontrolled asthma

BaxHTN

baxdrostat potential first-inclass ASI addressing hard-to-treat hypertension

Rare Disease

CARES

anselamimab improving survival in a subgroup of patients with light-chain amyloidosis

NME

NME

PREVAIL

NME

gefurulimab expanding reach in generalised myasthenia gravis

Readouts across 2025 represent combined >\$10bn opportunity1



Significant progress with transformative technologies to drive 2030+ growth

Weight management and risk factors

Establish and lead in new weight management paradigm

ADCs and Radioconjugates

Replace systemic chemotherapy and

radiotherapy

Next-gen IO bispecifics

Replace existing PD-1/PD-L1 inhibitors

Cell therapy and T-cell engagers

Develop scalable cell therapies and T-cell engagers across therapy areas Gene therapy and gene editing

Make cure possible for a range of rare diseases



3 Phase III trials initiated with laroprovstat (oPCSK9)

Multiple Phase II dose optimisation trials ongoing

AZD5004 (oGLP-1) AZD6234 (SARA) 7 AZN ADCs in clinic, including sonesitatug vedotin (CLDN18.2) in Phase III for 2L+ GC/GEJA

FPI-2265 (PSMA-targeted RC) in Phase II for pre-treated mCRPC

14 Phase III trials with rilvegostomig and volrustomig initiated

First ADC combination data at ASCO 2025

AZD0120 (BCMA/CD19)
CAR-T Phase III planned
in multiple myeloma

surovatamig (CD19/CD3) in Phase III for 1L FL

EsoBiotec acquired

Preclinical and Phase I development ongoing across multiple platforms

sAAVy and AAV capsid

TALEN technology

ADCs/RCs, next-gen IO and cell therapy/TCE progressed to Phase III





Financial Results

Aradhana SarinCHIEF FINANCIAL OFFICER



H1 and Q2 2025 – Reported profit and loss

	H1 2025 \$m	CER change %	% Total Revenue	Q2 2025 \$m	CER change %	% Total Revenue
- Product Sales	26,670	10	95	13,795	10	95
- Alliance Revenue	1,293	38	5	654	35	5
Product Revenue	27,963	11	100	14,449	11	100
- Collaboration Revenue	82	66	-	8	>2x	-
Total Revenue	28,045	11	100	14,457	11	100
Gross Margin	83%	-		83%	-	
- R&D expense	(6,707)	16	24	(3,548)	16	25
- SG&A expense	(9,356)	-	33	(4,864)	(2)	34
Total operating expense ¹	(16,341)	6	58	(8,555)	5	59
Other operating income and expense	192	53	1	79	33	1
Operating profit	7,182	24	26	3,508	32	24
Tax rate	18%			22%		
Reported EPS	\$3.46	32		\$1.58	31	



Due to rounding, the sum of the dollar values and percentages may not agree to totals. Absolute values at actual exchange rates; changes at CER.

1. Total operating expense includes distribution, R&D and SG&A expenses.

Appendix: Glossary.

H1 and Q2 2025 - Core profit and loss

	H1 2025 \$m	CER change %	% Total Revenue	Q2 2025 \$m	CER change %	% Total Revenue
- Product Sales	26,670	10	95	13,795	10	95
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Product Revenue	27,963	11	100	14,449	11	100
- Collaboration Revenue	82	66	-	8	>2x	-
Total Revenue	28,045	11	100	14,457	11	100
Gross Margin	83%	-		82%	-	
- R&D expense	(6,541)	17	23	(3,453)	18	24
- SG&A expense	(7,259)	3	26	(3,802)	1	26
Total operating expense ¹	(14,078)	9	50	(7,401)	9	51
Other operating income and expense	186	51	1	71	23	-
Operating profit	9,387	13	33	4,584	14	32
Tax rate	18%			21%		
Core EPS	\$4.66	17		\$2.17	12	

Due to rounding, the sum of the dollar values and percentages may not agree to totals. Absolute values at actual exchange rates; changes at CER.

1. Total operating expense includes distribution, R&D and SG&A expenses.

Appendix: Glossary.



Investing to support our long-term growth ambition

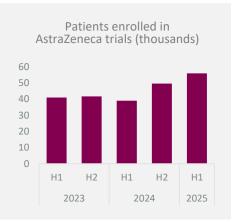
Driving operating leverage through SG&A efficiency

Total Revenue vs. Core R&D and Core SG&A as percentage of Total Revenue



Prioritising R&D

- Accelerating high-priority assets
- Investing in transformative technologies to underpin long-term growth



Optimising SG&A

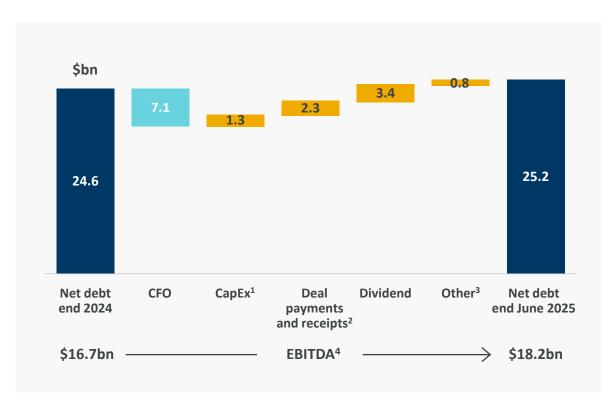
- Disease area focus
- Disciplined and strategic launch investments
- Improved productivity with digital and AI solutions



FY 2025 guidance reiterated

Net cash inflow from operating activities increased by 27% in H1 2025

Net debt/EBITDA 1.4x



FY 2025 guidance (CER)

Total Revenue

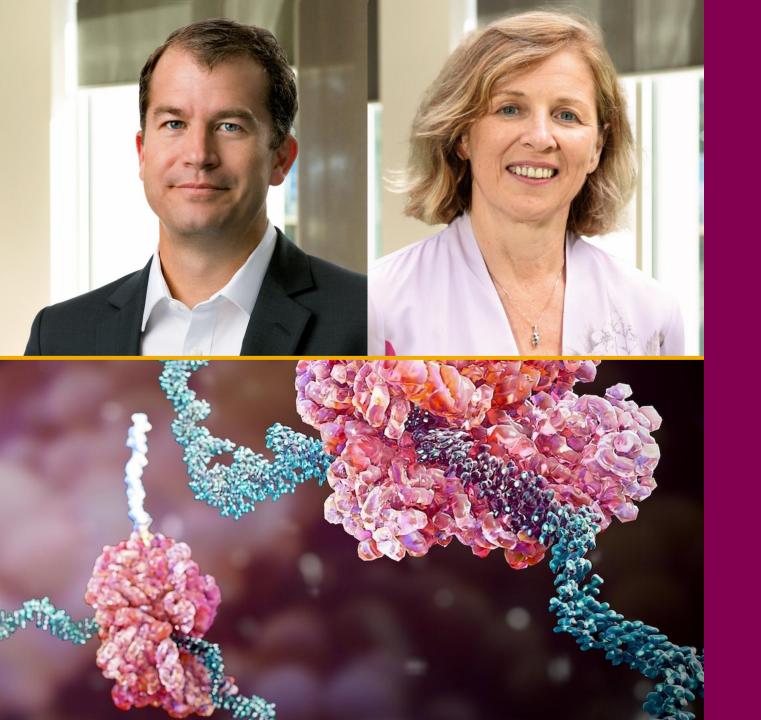
anticipated to increase by a high single-digit percentage

Core EPS

anticipated to increase by a low double-digit percentage

- Core tax rate expected to be between 18-22%
- Anticipated FX impact⁵ neutral on Total Revenue and Core EPS





Dave Fredrickson

ONCOLOGY HAEMATOLOGY BUSINESS

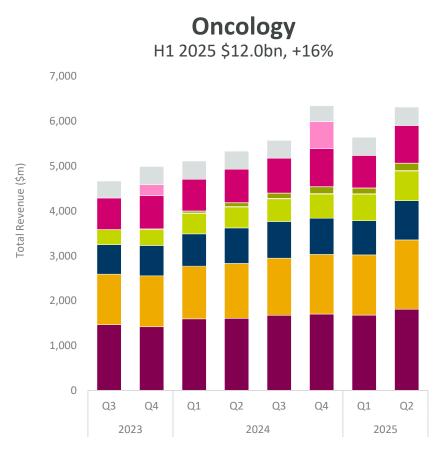
Susan Galbraith

ONCOLOGY HAEMATOLOGY R&D



Oncology – H1 and Q2 2025

Total Revenue +16% in H1 2025 driven by strong global demand across medicines



Tagrisso Imfinzi + Imjudo Calquence Enhertu Truqap Lynparza (PR) Lynparza (CR) Others¹

Q2 2025: key dynamics

- Tagrisso +12%, increasing demand across indications, leading 1L combination share
- Calquence +10%, extending BTKi leadership in CLL across major markets
- Lynparza PR +11%, sustained PARPi leadership
- Truqap +84%, continued demand growth in 2L biomarker-altered population
- *Imfinzi* +26%, strong demand growth with encouraging uptake for new launches (lung cancer ADRIATIC, AEGEAN; bladder cancer NIAGARA)
- Imjudo +18%, robust HCC and 1L NSCLC demand across major markets
- Enhertu +42%, accelerating DESTINY-Breast06 launch uptake, further CN adoption post-NRDL
- Datroway \$11m, positive early launch uptake in HR+ HER2- breast cancer

Key regulatory approvals:

 US (Datroway 2L+ EGFRm NSCLC²), EU (Calquence ECHO, AMPLIFY, Imfinzi NIAGARA), JP (Tagrisso LAURA), CN (Imfinzi ADRIATRIC, Orpathys + Tagrisso SACHI)



Oncology – high-value readouts year to date

Practice-defining data at ASCO 2025 and new high-level results across multiple tumour types

Transforming treatment across breast and gastric cancer at ASCO 2025

Establishing new endocrine backbone in HR+ BC

SERENA-6¹ | switch to camizestrant with CDK4/6i

- 56% reduction in risk of progression or death
- <1.5% discontinuation rate due to AEs</p>
- Strong trend to PFS2, meaningful prolongation in QoL

Moving earlier in HER2+ mBC

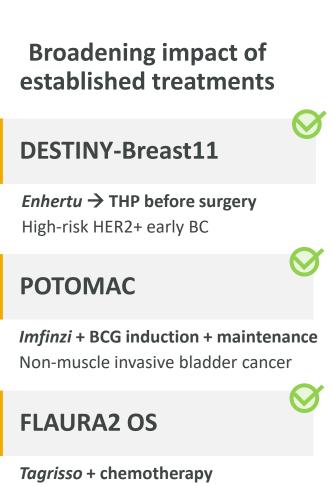
DESTINY-Breast09² | *Enhertu* + pertuzumab

- 44% reduction in risk of progression or death
- Median PFS >3 years, consistent across subgroups
- Strong trend to PFS2, early trend to OS benefit

New perioperative regimen for gastric/GEJ cancers

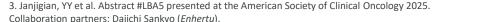
MATTERHORN³ | *Imfinzi* + FLOT

- Median EFS not yet met
- Two-thirds event-free at two years
- Strong trend to OS benefit with HR 0.78



11 metastatic FGFRm NSCLC

^{1.} Turner, NC et al. Abstract #LBA4 presented at the American Society of Clinical Oncology 2025. 2. Tolaney, SM et al. Abstract #LBA1008 presented at the American Society of Clinical Oncology 2025.





BioPharmaceuticals

Ruud Dobber

BIOPHARMACEUTICALS BUSINESS

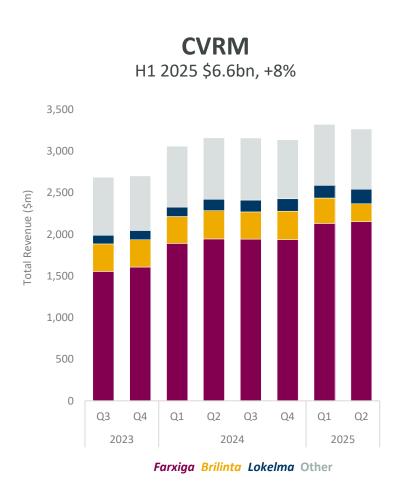
Sharon Barr

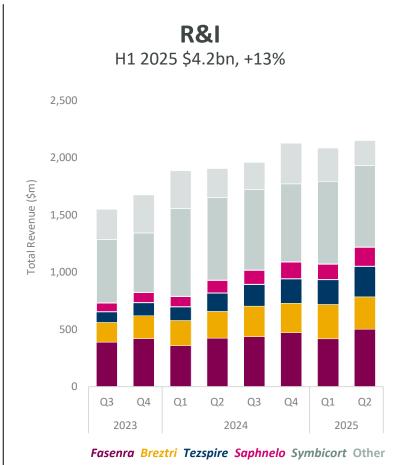
BIOPHARMACEUTICALS R&D



BioPharmaceuticals – H1 and Q2 2025

Total Revenue +10% in H1 2025 driven by strong momentum in key medicines





Q2 2025: key dynamics

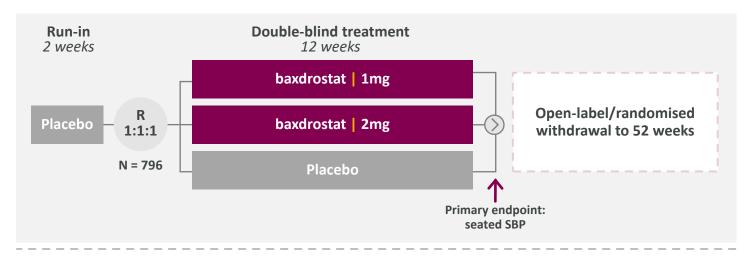
- Farxiga +10%, global demand growth mainly driven by CKD and HF
- Lokelma +27%, market leader in growing K+ binder class in hyperkalaemia
- Brilinta (38%), generics entry
- Fasenra +18%, sustained IL-5 leadership in asthma, EGPA launches
- Tezspire +65%, continued launch momentum
- Breztri +20%, fastest growing medicine in expanding FDC triple class in COPD
- Saphnelo +48%, increasing penetration in i.v. segment of SLE
- **V&I** +54%, *Beyfortus* >3x



BioPharmaceuticals – first Phase III data for baxdrostat

Potential first-in-class NME; \$5bn+ PYR opportunity across monotherapy and combinations

Phase III BaxHTN in uncontrolled or resistant hypertension



Primary and all secondary endpoints met



Clinically meaningful reduction in SBP

Favourable safety profile

Addressing high unmet need

Data to be presented at ESC 2025 | 30 August

Broad Phase III development

Monotherapy	
Bax24 24-hour SBP control	H2 2025
BaxAsia u/r HTN in Asian patients	H1 2026
BaxPA primary aldosteronism	Initiating
Combination with dapagliflozin	
BaxDUO-Arctic CKD and HTN	>2026
BaxDUO-Pacific CKD and HTN outcomes	>2026
Prevent-HF HF prevention	>2026





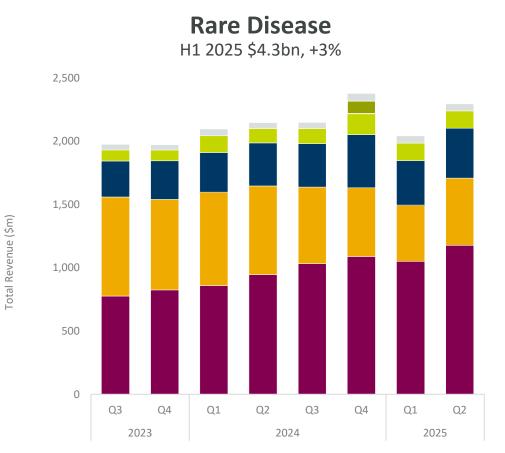
Rare Disease

Marc Dunoyer CHIEF EXECUTIVE OFFICER, ALEXION



Rare Disease – H1 and Q2 2025

Total Revenue \$4.3bn in H1 2025 driven by patient demand across the portfolio



Ultomiris¹ Soliris Strensig Koselugo (PS) Koselugo (CR) Other²

Q2 2025: key dynamics

C5 Franchise

- Ultomiris +23%, demand growth across indications, including within the competitive gMG and PNH markets
- Soliris (22%), continued successful conversion to Ultomiris across indications and additional impact from biosimilars in EU, partially offset by order timing in certain tender markets

Beyond Complement

- Strensiq +15%, continued global demand from patients with HPP
- Koselugo +18%, continued global demand from patients with NF1-PN



Rare Disease – Phase III readouts for two NMEs

Reinforcing innovation and leadership in Rare Disease

PREVAIL | gefurulimab in gMG

Dual-binding nanobody targeting C5, self-administration s.c. QW¹

- All primary and secondary endpoints met
- Statistically significant and clinically meaningful improvement in MG-ADL
- Rapid, complete and sustained complement inhibition, translating to improvements in patient outcomes

Global gMG market dynamics

<20%

Patients treated with branded medicines, expected increase to 50% in 5 years²

~40%

Patients on self-admin. medicines by 2030²

Potential to position gefurulimab first-line post-immunosuppressant therapy and steroids

CARES | anselamimab in light-chain amyloidosisSpecific anti-fibril

- Composite primary endpoint not met in Mayo Stage IIIa and IIIb
- Highly clinically meaningful improvement in prespecified subgroup on:
 - All-cause mortality
 - **OVER SECTION** Cardiovascular hospitalisation
- Data to be shared with global health authorities



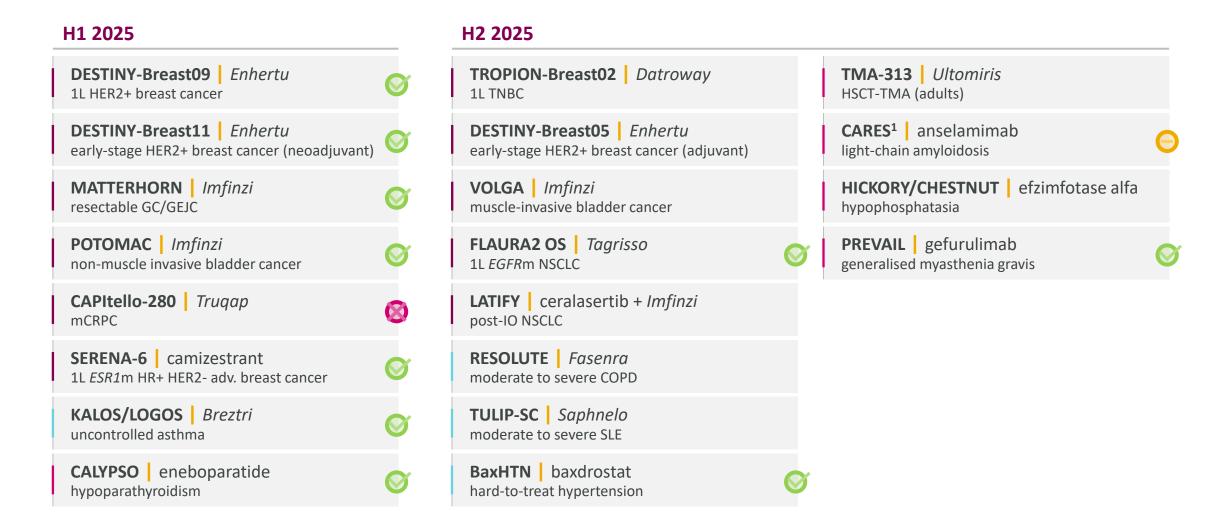


CEO Closing Remarks

Pascal Soriot
CHIEF EXECUTIVE OFFICER



Successfully delivering in unprecedented catalyst rich period





On track to deliver on 2030 ambitions supported by strong growth and pipeline momentum



Ambition to deliver \$80bn in Total Revenue by 2030¹

Strong growth in 2025 with global medicines demand substantially offsetting anticipated headwinds

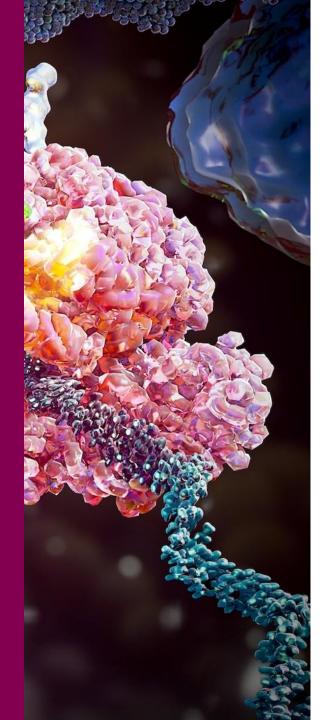
On track to deliver mid-30s% Core operating margin by 2026

Growth in SG&A slower than Total Revenue R&D to remain low 20%s of Total Revenue

Ambition to deliver at least 20 NMEs by 2030

9 NMEs launched to date
First Phase III data for 5 NMEs
in 2025 to date





Q&A Session



Pascal Soriot
CHIEF EXECUTIVE OFFICER



Aradhana Sarin
CHIEF FINANCIAL OFFICER



Marc Dunoyer CHIEF EXECUTIVE OFFICER, ALEXION



Susan Galbraith EVP, ONCOLOGY HAEMATOLOGY R&D



Dave FredricksonEVP, ONCOLOGY
HAEMATOLOGY BUSINESS



Sharon Barr EVP, BIOPHARMACEUTICALS R&D

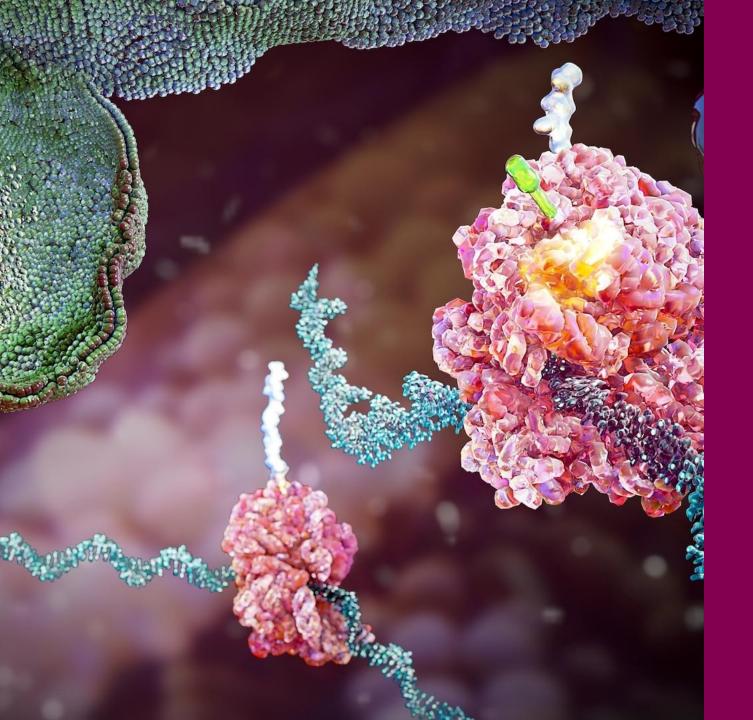


Ruud Dobber EVP, BIOPHARMACEUTICALS BUSINESS



Iskra Reic EVP, INTERNATIONAL





Appendix



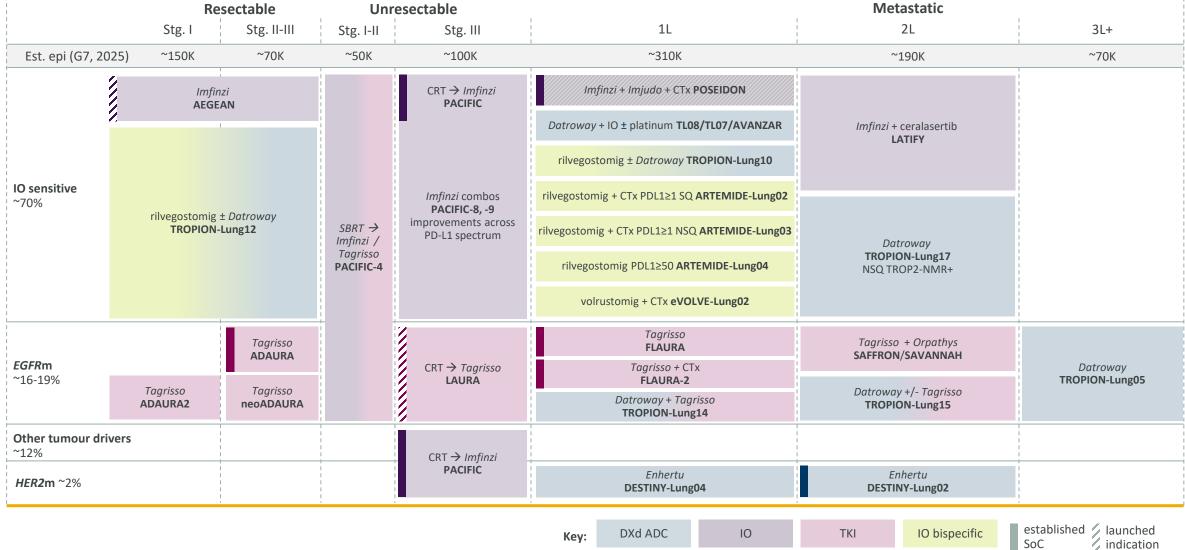
AstraZeneca P&L reference table

P&L line-item definitions

	P&L line-item definition
Product Sales	 Recognises sales from territories where Group has lead commercialisation Recognises supply of <i>Beyfortus</i> to Sanofi
Alliance Revenue	 Alliance Revenue comprises income arising from the ongoing operation of collaborative arrangements related to sales made by collaboration partners, where AstraZeneca is entitled to a share of gross profits, share of revenues or royalties, which are recurring in nature while the collaboration agreement remains in place¹
Product Revenue	The sum of Product Sales and Alliance Revenue
Collaboration Revenue	 Recognises any development or sales-based milestone received on partnered medicines as well as any upfront payments associated with business development where AstraZeneca retains a significant ongoing economic interest in the product
Total Revenue	Sum of Product Sales, Alliance Revenue and Collaboration Revenue
Gross Margin	Calculated by dividing Gross Profit by Total Revenue
Other operating income & expense	 Other operating income and expense is generated from activities outside of the Group's normal course of business, which includes Other income from divestments of or full out-license of assets and businesses including royalties and milestones where the Group does not retain a significant continued interest
Core ² Operating margin	Defined as Core Operating profit as a percentage of Total Revenue

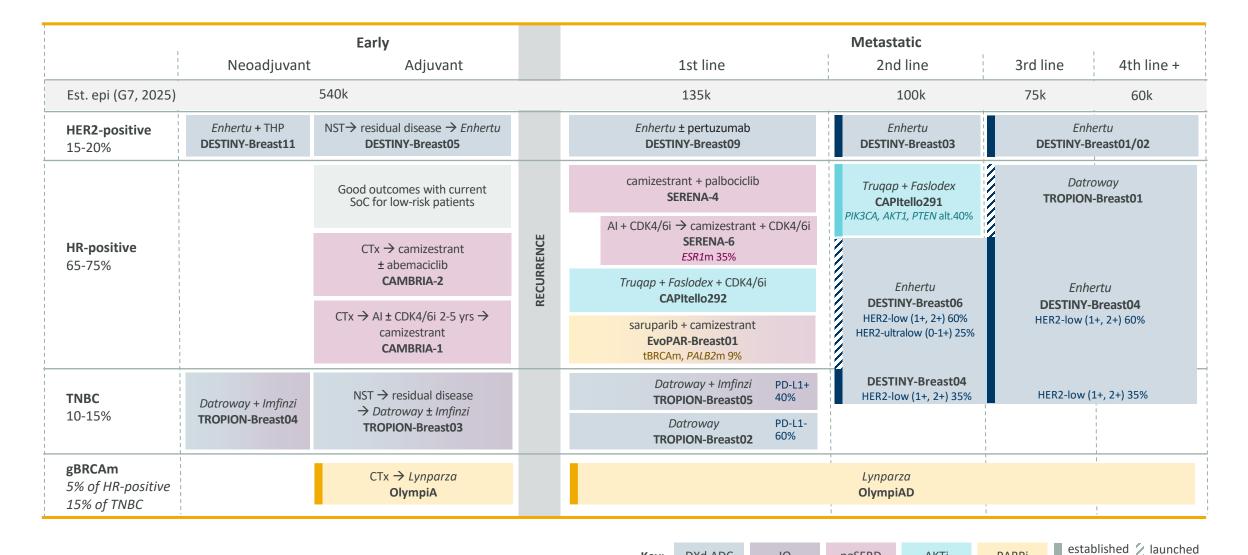


AstraZeneca in non-small cell lung cancer





AstraZeneca in breast cancer



DXd ADC

Key:

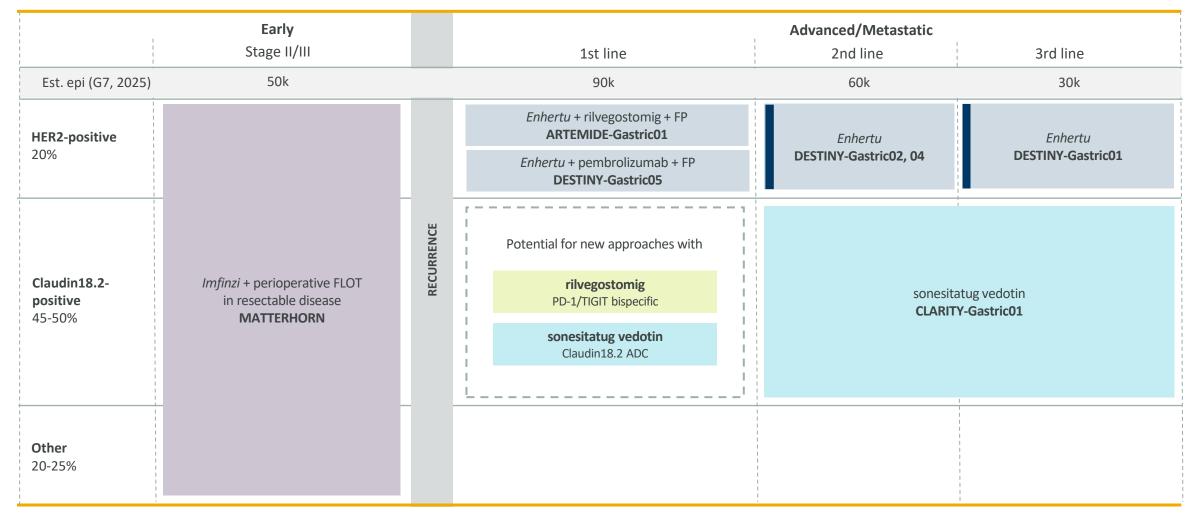
ngSERD

AKTi

PARPi



AstraZeneca in gastric cancer



Key:

DXd ADC

10

AZN ADC

IO bispecific

established // launched SoC



sonesitatug vedotin previously AZD0901.

Strengthening manufacturing and R&D footprint to support future growth



Six strategic R&D centres globally:



Cambridge, UK



Gaithersburg, US



Shanghai, China



Gothenburg, Sweden



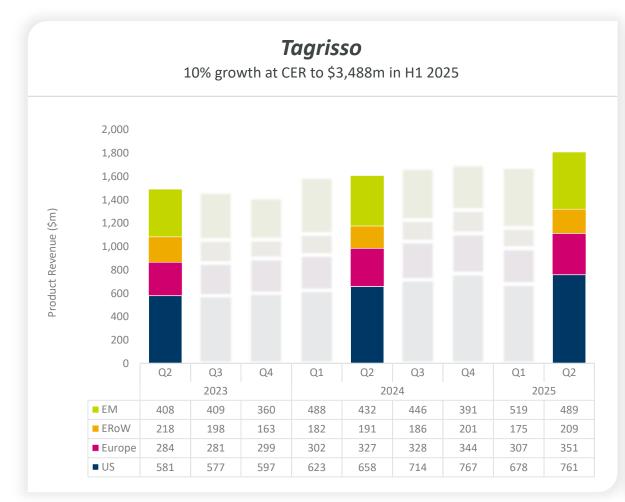
Boston, US - Kendall Sq.

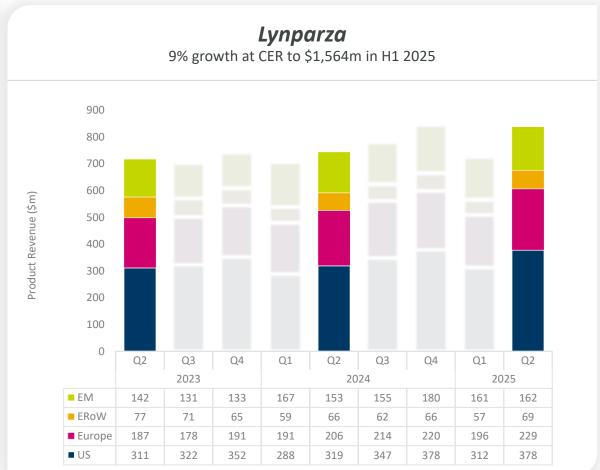


Beijing, China

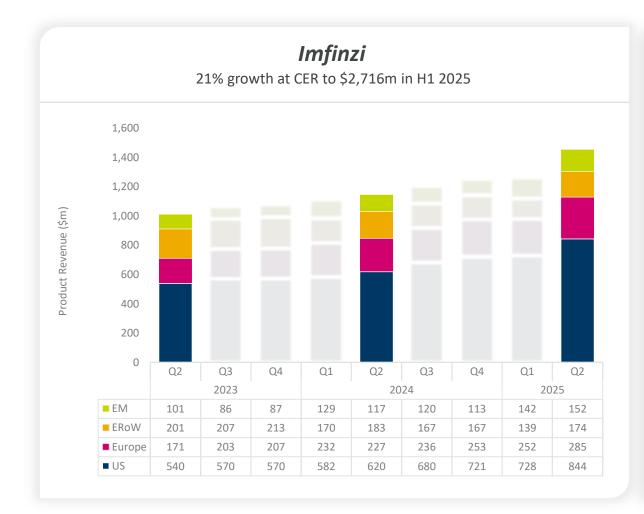
Investments support \$80bn 2030 Total Revenue¹ ambition

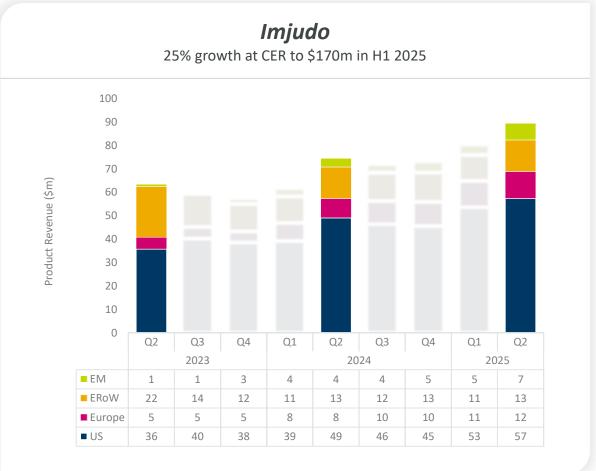




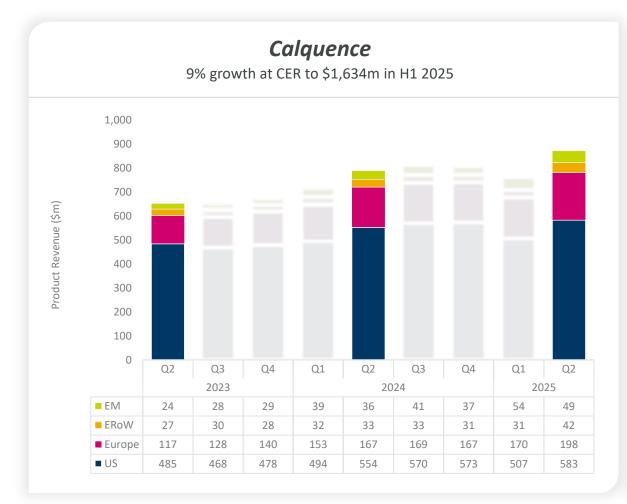


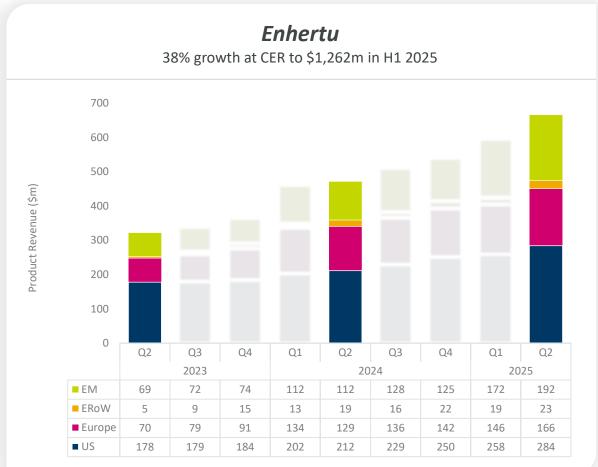














BioPharmaceuticals: Cardiovascular, Renal & Metabolism

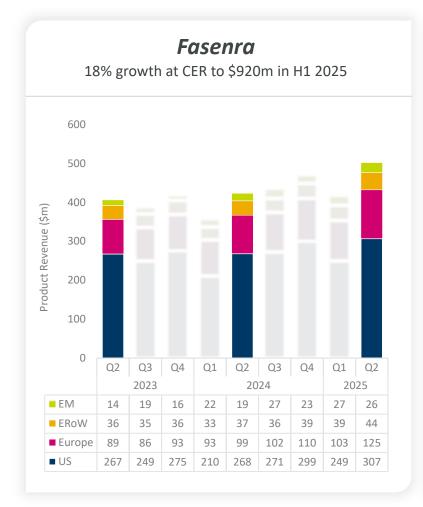




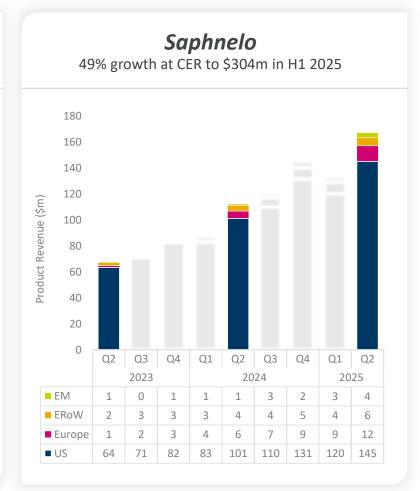




BioPharmaceuticals: Respiratory & Immunology

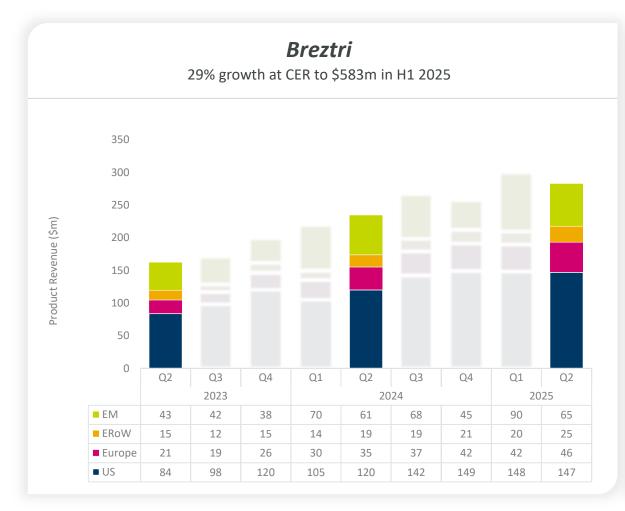


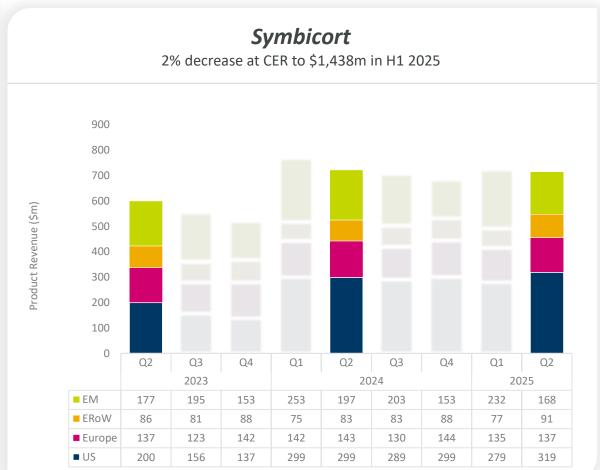






BioPharmaceuticals: Respiratory & Immunology







Rare Disease

Ultomiris 24% growth at CER to \$2,228m in H1 2025 1,400 1,200 1,000 Product Revenue (\$m) 800 600 200 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 2023 2024 2025 EM 52 152 173 166 179 131 138 143 170 173 202 209 238 235 228 270 490 | 482 | 550 | 597 | 632 | 604 | 667







Glossary

1L, 2L, 3L	first-, second-, third-line
AAV	adeno-associated virus
ADC	antibody-drug conjugate
adv.	advanced
Al	aromatase inhibitor
AKT1	AKT serine/threonine kinase 1
ASCO	American Society of Clinical Oncology
ASI	aldosterone synthase inhibitor
ВС	breast cancer
BCG	Bacillus Calmette-Guérin
BCMA	B-cell maturation antigen
ВТКі	Bruton's tyrosine kinase
C5	complement component 5
СарЕх	capital expenditure
CD19	cluster of differentiation 19
CD3	cluster of differentiation 3
CDK4/6i	cyclin-dependent kinase 4/6 inhibitor
CER	constant exchange rates
CFO	net cash inflow from operating activities
CKD	chronic kidney disease
CLDN18.2	Claudin-18.2
CLL	chronic lymphocytic leukaemia
CN	China
COPD	chronic obstructive pulmonary disease
CR	Collaboration Revenue
CRT	chemoradiotherapy
СТх	chemotherapy
CVRM	Cardiovascular, Renal and Metabolism
Dxd	deruxtecan
EBITDA	earnings before interest, tax, depreciation and amortisation
EFS	event-free survival
<i>EGFR</i> m	epidermal growth factor receptor-mutant
EGPA	eosinophilic granulomatosis with polyangiitis
epi	epidemiology
EPS	earnings per share
ERoW	Established Rest of World
ESC	European Society of Cardiology
ESR1m	estrogen receptor alpha-mutated
EU	Europe
	•

FDC	fixed-dose combination
FLOT	luorouracil, leucovorin, oxaliplatin and docetaxel
FP	fluoropyrimidine
FX	foreign exchange
gBRCAm	germline BRCA-mutated breast cancer
GC	gastric cancer
GEJ	gastroesophageal junction
GEJA	gastroesophageal junction adenocarcinoma
GEJC	gastroesophageal junction cancer
gMG	generalised myasthenia gravis
HER2-/negative	human epidermal growth factor receptor 2-negative
HER2-low	human epidermal growth factor receptor 2-low
HER2m	human epidermal growth factor receptor 2-mutant
HER2+/positive	human epidermal growth factor receptor 2-positive
HER2-ultralow	human epidermal growth factor receptor 2-ultralow
HF	heart failure
HR	hazard ratio
HR+/positive	hormone receptor-positive
HSCT-TMA	hematopoietic stem cell transplantation-associated thrombotic microangiopathy
HTN	hypertension
i.v.	intravenous
IL-5	interleukin-5
10	immuno-oncology
JP	Japan
mAb	monoclonal antibody
mBC	metastatic breast cancer
mCRPC	metastatic castration-resistant prostate cancer
mg	milligram
MG-ADL	Myasthenia Gravis Activities of Daily Living
NME	new molecular entity
NMIBC	non-muscle invasive bladder cancer
NMR+	nuclear magnetic resonance-positive
NRDL	national reimbursement drug list
NSCLC	non-small cell lung cancer
NSQ	non-squamous
NST	neoadjuvant systemic treatment
oGLP-1	oral glucagon-like peptide-1
oPCSK9	oral protein convertase subtilisin/kexin type 9
OS	overall survival

PDL1 programmed death-ligand 1 PD-L1 programmed cell death ligand 1 PFS progression free survival PFS pre-filled syringe second progression-free survival PFS2 PIK3CA phosphatidylinositol-4,5-biphosphate 3-kinase catalytic subunit PNH paroxysmal nocturnal haemoglobinuria PR Product Revenue prostate-specific membrane antigen **PSMA** phosphatase and TENsin homolog deleted on chromosome 10 PTEN PYR Peak-Year Revenue QoL quality of life QW once weekly R&D Research & Development R&I Respiratory & Immunology RC radioconjugate subcutaneous s.c. SARA selective amylin receptor agonist SBP systolic blood pressure **SBRT** stereotactic brain radiotherapy self-admin. self-administered SG&A Selling, General & Administrative SLE systemic lupus erythematosus SoC standard-of-care Stg. stage tBRCAm tumor BRCA mutation TCE T-cell engager THP docetaxel, trastuzumab and pertuzumab **TIGIT** T-cell immunoreceptor with immunoglobulin and ITIM domains TKI tyrosine kinase inhibitor TL07 TROPION-Lung07 TL08 TROPION-Lung08 **TNBC** triple negative breast cancer u/r HTN uncontrolled/resistant hypertension US **United States** V&I Vaccines & Immune Therapies YTD year-to-date

poly-ADP ribose polymerase inhibitor

PARPi



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