



Q1 2025 Results

Conference call and webcast for investors
and analysts

29 April 2025



Forward-looking statements

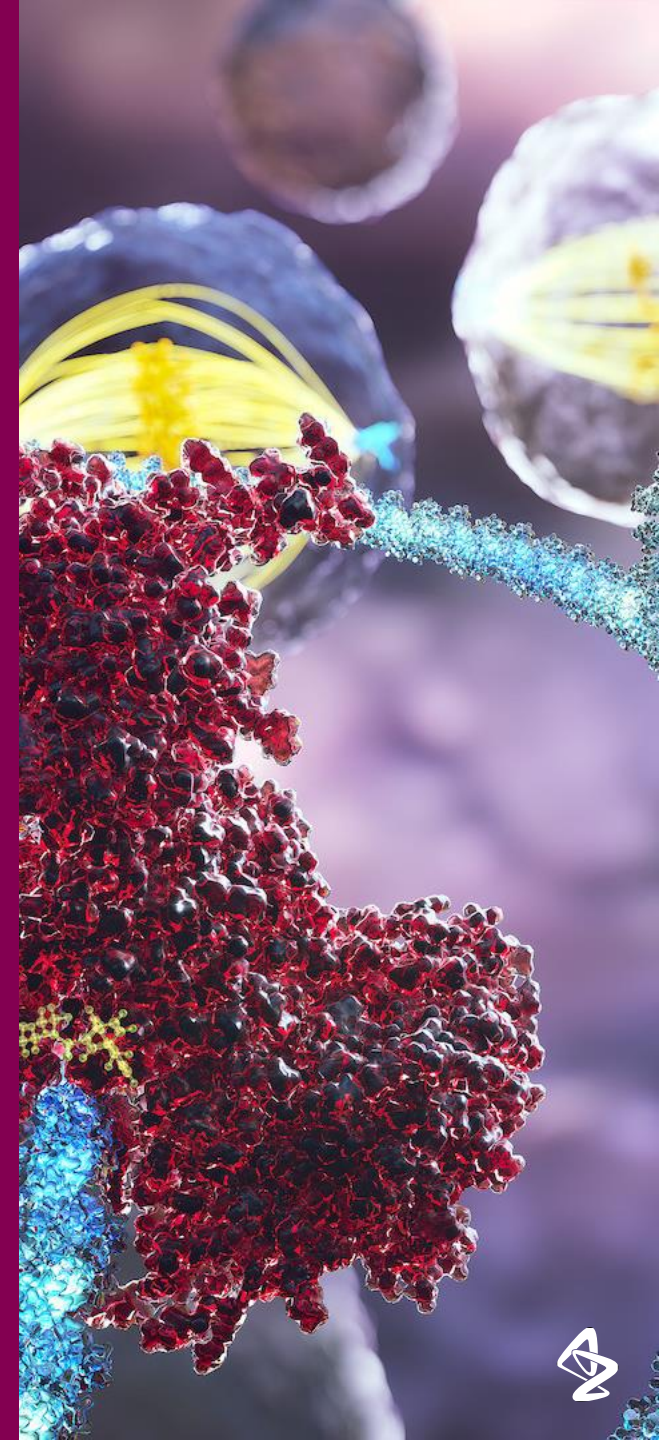
This document contains certain forward-looking statements with respect to the operations, performance and financial condition of the Group, including, among other things, statements about expected revenues, margins, earnings per share or other financial or other measures. Although the Group believes 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 AI 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; the risk of unexpected deterioration in the Group’s financial position; the risk of foreign exchange rate movements impacting our financial condition or results of operations; and the impact that global and/or geopolitical events may have or continue to have on these risks, on the Group’s ability to continue to mitigate these risks, and on the Group’s operations, financial results or financial condition. Nothing in this document, or any related presentation/webcast, should be construed as a profit forecast.

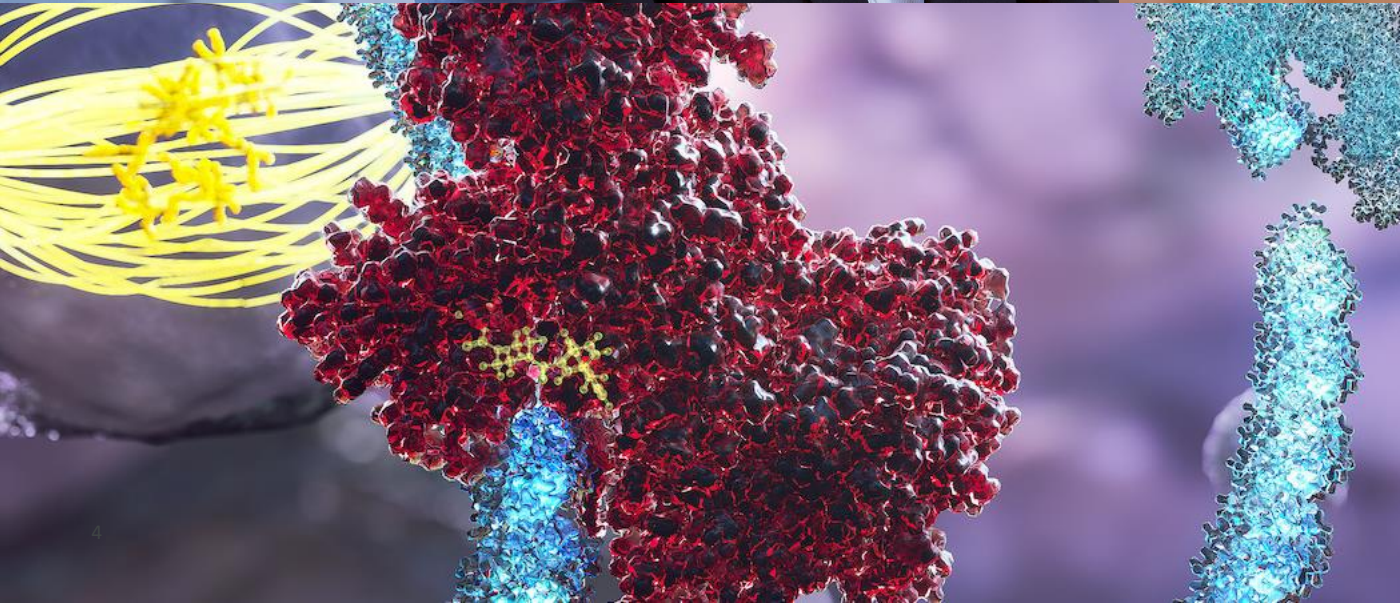


Q1 2025 Results

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 Q1 2025



Sustained commercial momentum

+10% Total Revenue¹

+21% Core EPS¹

13 new approvals
in key regions^{2,3}



Continued pipeline delivery

5 positive Phase III readouts

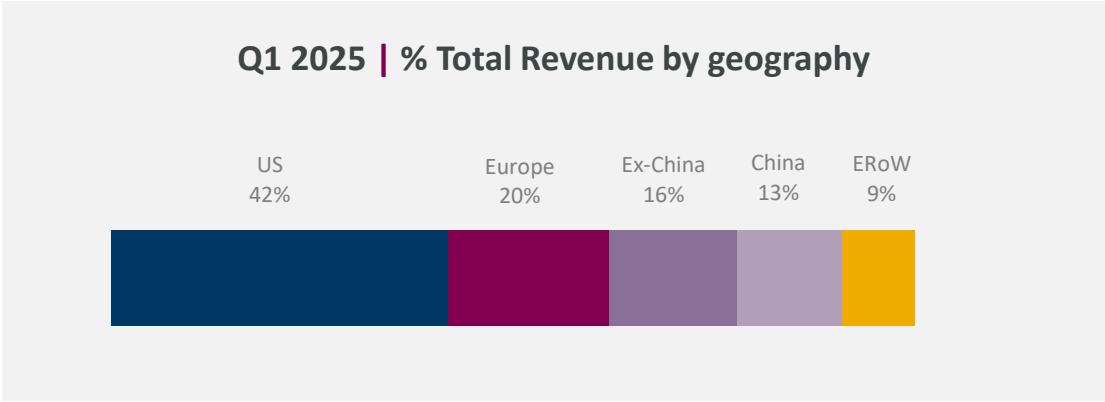
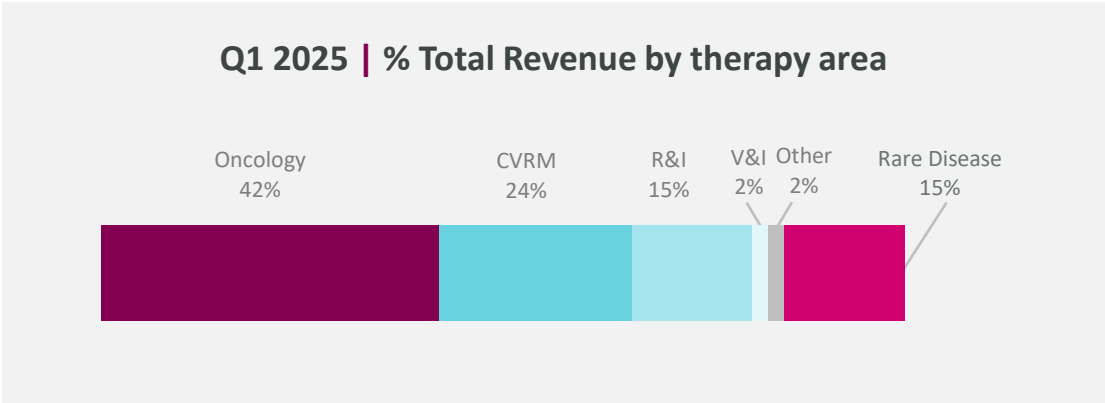
Key indication expansion
Imfinzi in GC/GEJC and
Enhertu in breast, gastric³

9 NMEs delivered toward
ambition of 20 by 2030
with approval of
Beyontra (acoramidis)⁴



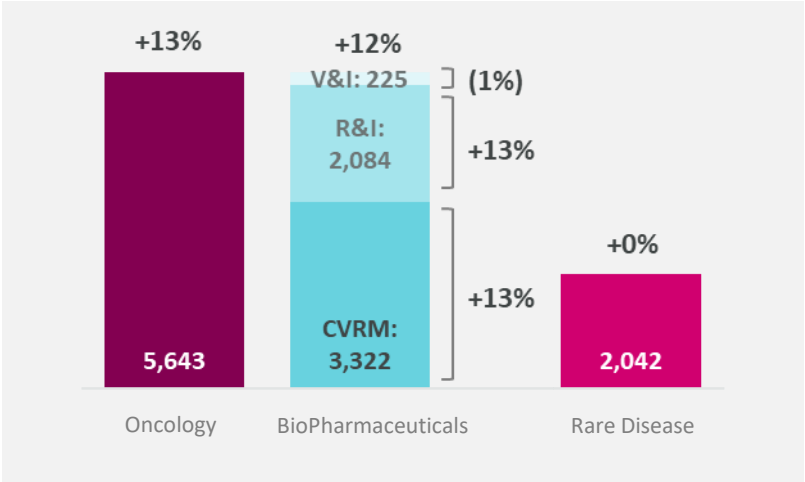
Q1 2025 – benefit from broad-based global business

Broad-based, diverse source of Total Revenue



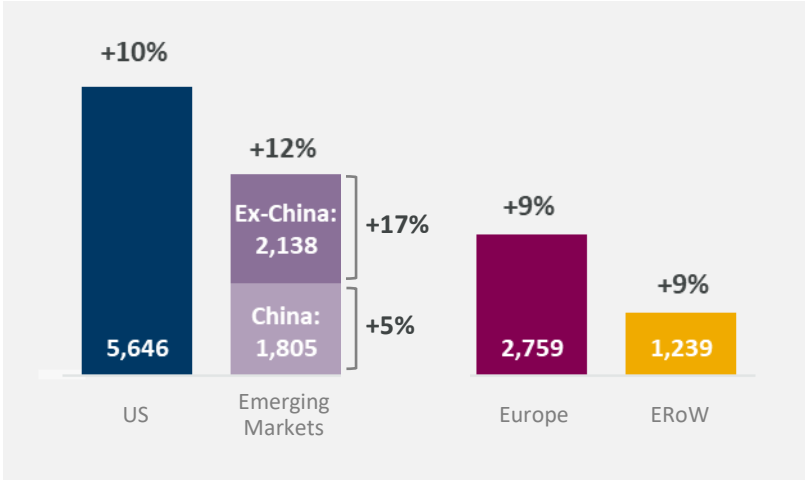
Strength across therapy areas

Q1 2025 | Total Revenue (\$m)



Growth across geographies

Q1 2025 | Total Revenue (\$m)



Investing in transformative technologies and global R&D footprint to support long-term growth ambitions

Value-enhancing business development in Q1 2025



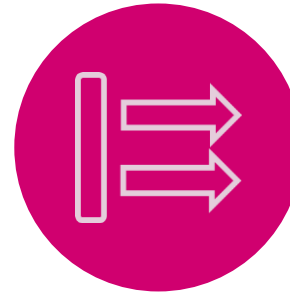
In-vivo cell therapy

- Proposed acquisition of EsoBiotec¹
- “Off-the-shelf” cell therapy platform



Novel modalities

- Multi-specific biologics with Harbour BioMed
- Macro-cyclic peptides with Syneron



Innovative delivery

- Exclusive license for ALT-B4 from Alteogen
- Subcutaneous formulations of multiple oncology assets



Cambridge, UK



Gothenburg, Sweden



Gaithersburg, US



Boston, US – Kendall Sq.



Shanghai, China



Beijing, China

**Global presence with
six strategic R&D centres**



A growing broad-based global footprint

Resilient, dual-source supply chain with investment in transformative technologies



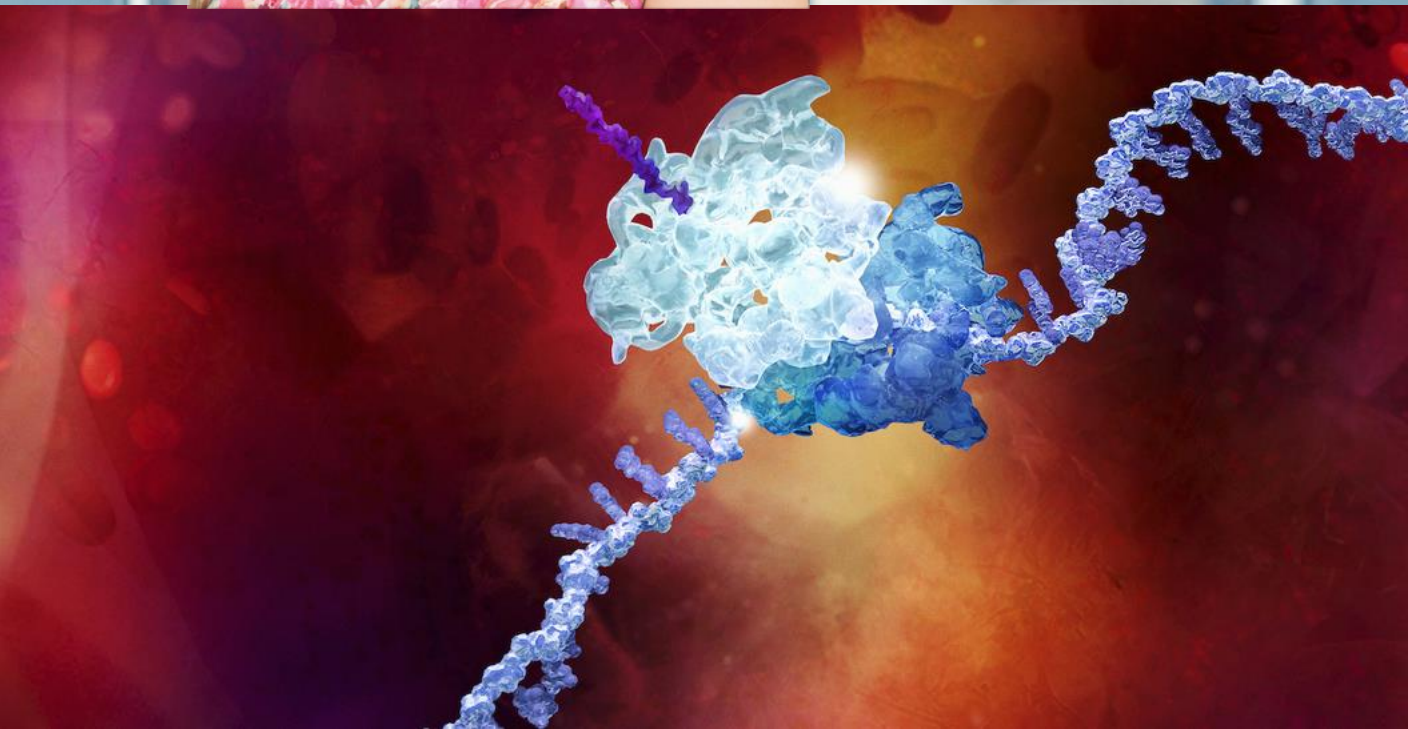
31 | production facilities

Key regions supplied by **local manufacturing**

11 | US sites

Vast majority of products **sold in US** are **manufactured in US**





Financial Results

Aradhana Sarin

CHIEF FINANCIAL OFFICER



Q1 2025 – Reported profit and loss

	Q1 2025 \$m	CER change %	% Total Revenue
- Product Sales	12,875	9	95
- Alliance Revenue	639	42	5
Product Revenue	13,514	10	99
- Collaboration Revenue	74	64	1
Total Revenue	13,588	10	100
Gross Margin	84%	-	
- R&D expense	(3,159)	15	23
- SG&A expense	(4,492)	3	33
Total operating expense ¹	(7,786)	7	57
Other operating income and expense	113	71	1
Operating profit	3,674	17	27
Tax rate	14%		
Reported EPS	\$1.88	32	

¹⁰ Due to rounding, the sum of a number of 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](#).



Q1 2025 – Core profit and loss

	Q1 2025 \$m	CER change %	% Total Revenue
- Product Sales	12,875	9	95
- Alliance Revenue	639	42	5
Product Revenue	13,514	10	99
- Collaboration Revenue	74	64	1
Total Revenue	13,588	10	100
Gross Margin	84%	-	
- R&D expense	(3,088)	16	23
- SG&A expense	(3,457)	4	25
Total operating expense ¹	(6,677)	9	49
Other operating income and expense	115	78	1
Operating profit	4,803	12	35
Tax rate	16%		
Core EPS	\$2.49	21	

¹¹ Due to rounding, the sum of a number of 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.

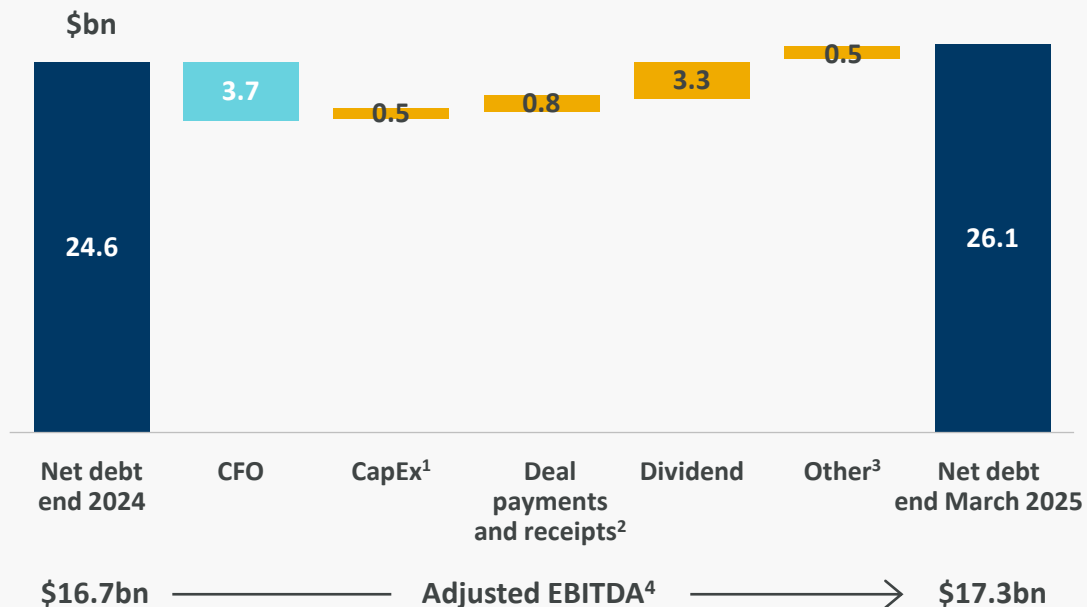
Appendix: [Glossary](#).



FY 2025 guidance reiterated

Net cash inflow from operating activities increased by 49% in Q1 2025

Net debt/Adjusted EBITDA 1.5x



FY 2025 guidance (CER)

Total Revenue

anticipated to increase by a **high single-digit** percentage

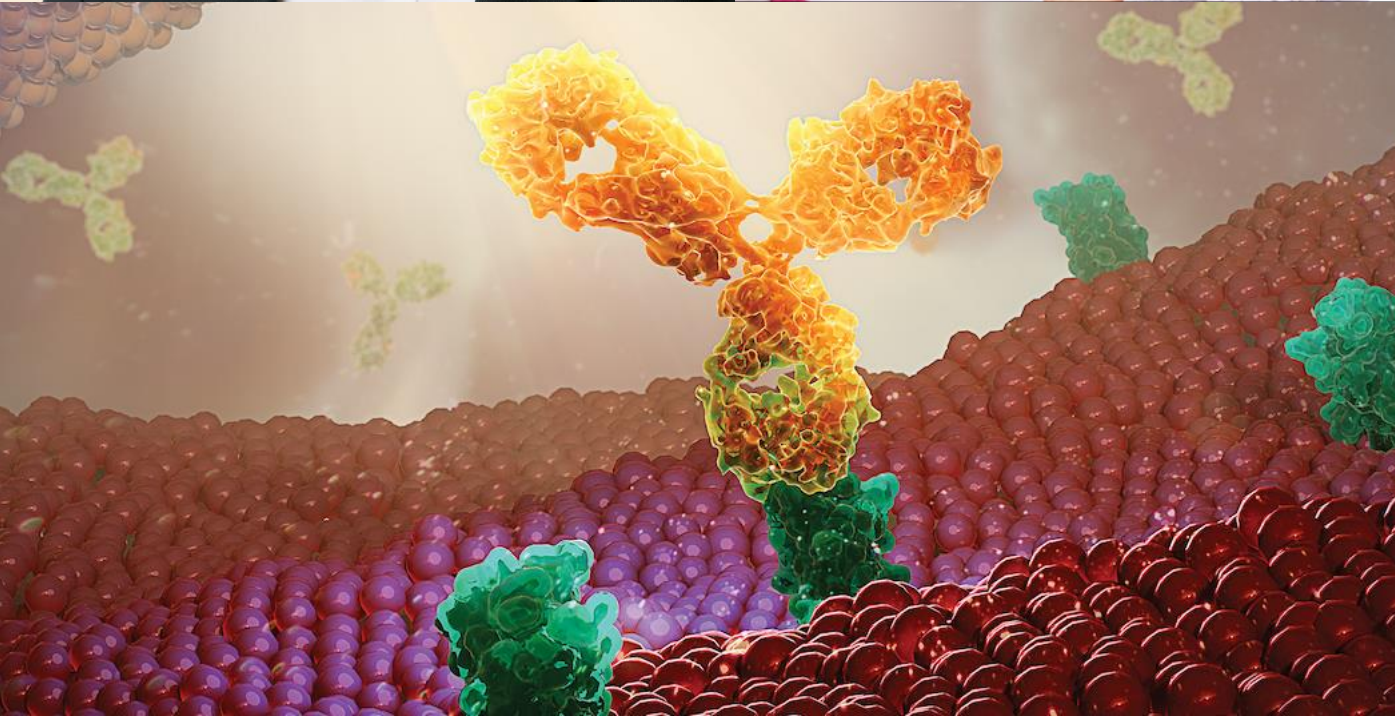
Core EPS

anticipated to increase by a **low double-digit** percentage

- Core tax rate expected to be between 18-22%
- Anticipated FX impact – low single-digit adverse impact on Total Revenue and Core EPS⁵

Due to rounding, the sum of a number of dollar values and percentages may not agree to totals. 1. Capital expenditure on tangible assets and software-related intangible assets. 2. Comprises purchase and disposal of intangible assets (excluding software-related assets, including AZ Forest), movement in profit participation liability, purchase and disposal of non-current asset investments, payments to associates and joint ventures, disposal of investments in associates and joint ventures, acquisitions of subsidiaries, net of acquired net debt and payment of contingent consideration on business combinations. The Company uses Debt issuance to finance new Business Development opportunities. 3. Comprises mainly Own shares purchased by Employee Benefit Trust. 4. Rolling 12m EBITDA. AstraZeneca credit ratings: Moody's: short-term rating P-1, long-term rating A1, outlook stable. S&P Global Ratings: short-term rating A-1, long-term rating A+, outlook stable. 5. If foreign exchange rates for April 2025 to December 2025 were to remain at the average rates seen in March 2025. Appendix: [Glossary](#).





Oncology

Dave Fredrickson
ONCOLOGY HAEMATOLOGY BUSINESS

Susan Galbraith
ONCOLOGY HAEMATOLOGY R&D

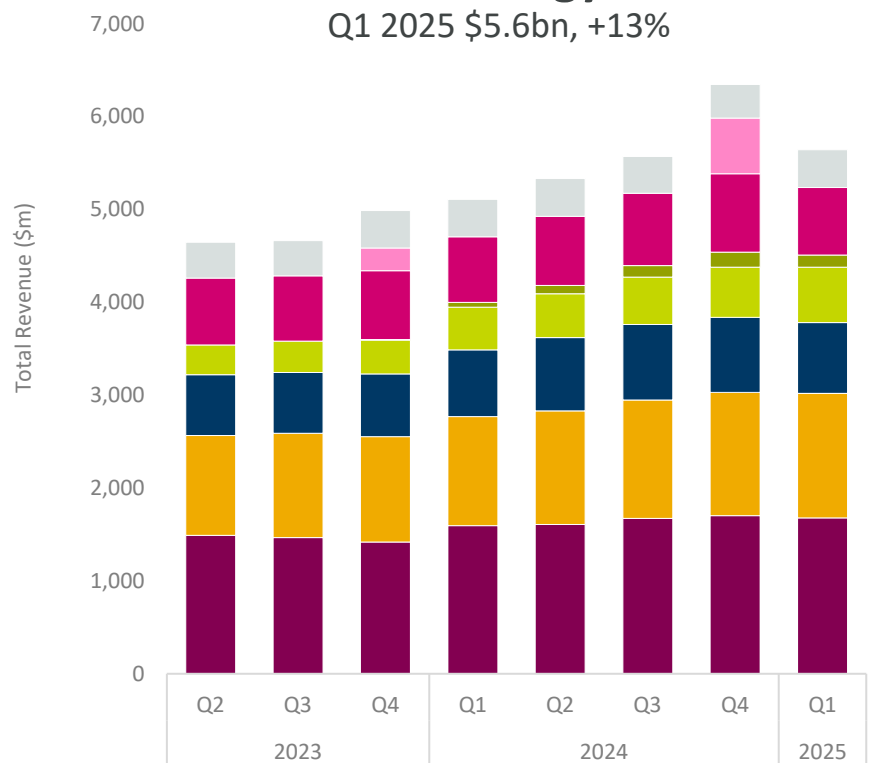


Oncology – Q1 2025

Strong demand across key regions, partly offset by oral oncolytics Part D redesign in US

Oncology

Q1 2025 \$5.6bn, +13%



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

Q1 2025: key dynamics

- **Tagrisso** +8%, increasing demand across indications, strong LAURA launch uptake
- **Calquence** +8%, continued BTKi leadership in CLL across major markets
- **Lynparza PR** +5%, sustained PARPi leadership
- **Truqap** \$132m, market leadership in 2L biomarker-altered, partly impacted by one-time blister pack destocking in US
- **Imfinzi** +16%, strong demand growth, continued JP repricing impact
- **Imjudo** +33%, durable HIMALAYA demand across major markets
- **Enhertu** +34%, strong demand in CN post-NRDL inclusion, DESTINY-Breast06 launch
- **Datroway** \$4m, positive early launch signals in HR+/HER2- breast

-
- **Key regulatory approvals:** US (*Imfinzi* NIAGARA), EU (*Imfinzi* ADRIATIC, *Enhertu* DESTINY-Breast06, *Imfinzi* AEGEAN)



Oncology – strong catalyst flow in Q1 2025

Positive key Phase III readouts reinforce leadership in breast and GI cancers

ASCO plenary

camizestrant SERENA-6

First positive oral SERD Phase III in 1L

AI + CDK4/6i switch to cami + CDK4/6i
1L *ESR1*m HR+/HER2- mBC

- Highly significant and clinically meaningful PFS improvement
- Trend in PFS2 improvement



~10k eligible patients across G7

ASCO plenary



MATTERHORN

Builds on AEGEAN and NIAGARA
perioperative success

+ FLOT
perioperative gastric / GEJ cancer

- Significant and clinically meaningful EFS improvement
- Strong trend to OS improvement



~43k eligible patients across G7



DESTINY-Breast09

Moves *Enhertu* earlier in HER2+ mBC

+ pertuzumab
1L HER2+ mBC

- Highly significant and clinically meaningful PFS improvement
- Early trend to OS improvement



~23k eligible patients across G7

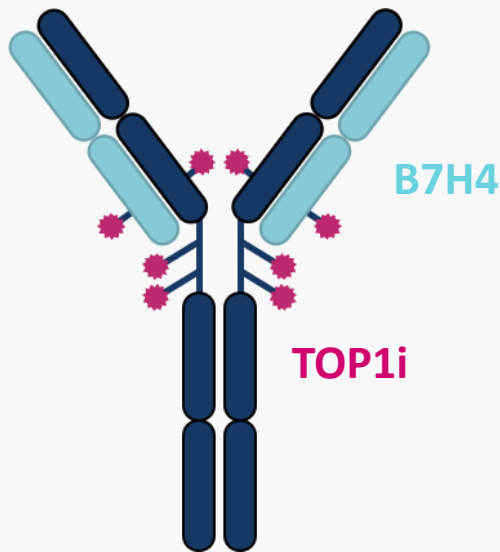
US Breakthrough Device Designation granted for TROP2-NMR companion diagnostic¹ in 2L *Datroway* TROPION-Lung17 trial



Oncology – advancing transformative technologies

Progress with in-house ADCs at Society of Gynecologic Oncology 2025

Encouraging puxitatug samrotescan Phase I/IIa data in endometrial cancer



- Promising ORR 34.6-38.5%
- Preliminary 7.0 months mPFS
- No TRAEs leading to discontinuation
- Low rates of haematologic toxicity

Initiating Phase III in endometrial cancer in 2025

AZD0901
Cldn18.2

Phase III

P-Sam
B7H4

Phase I/II

III

AZD5335
FR α

Phase I/II

AZD0305
GPC5D

Phase I/II

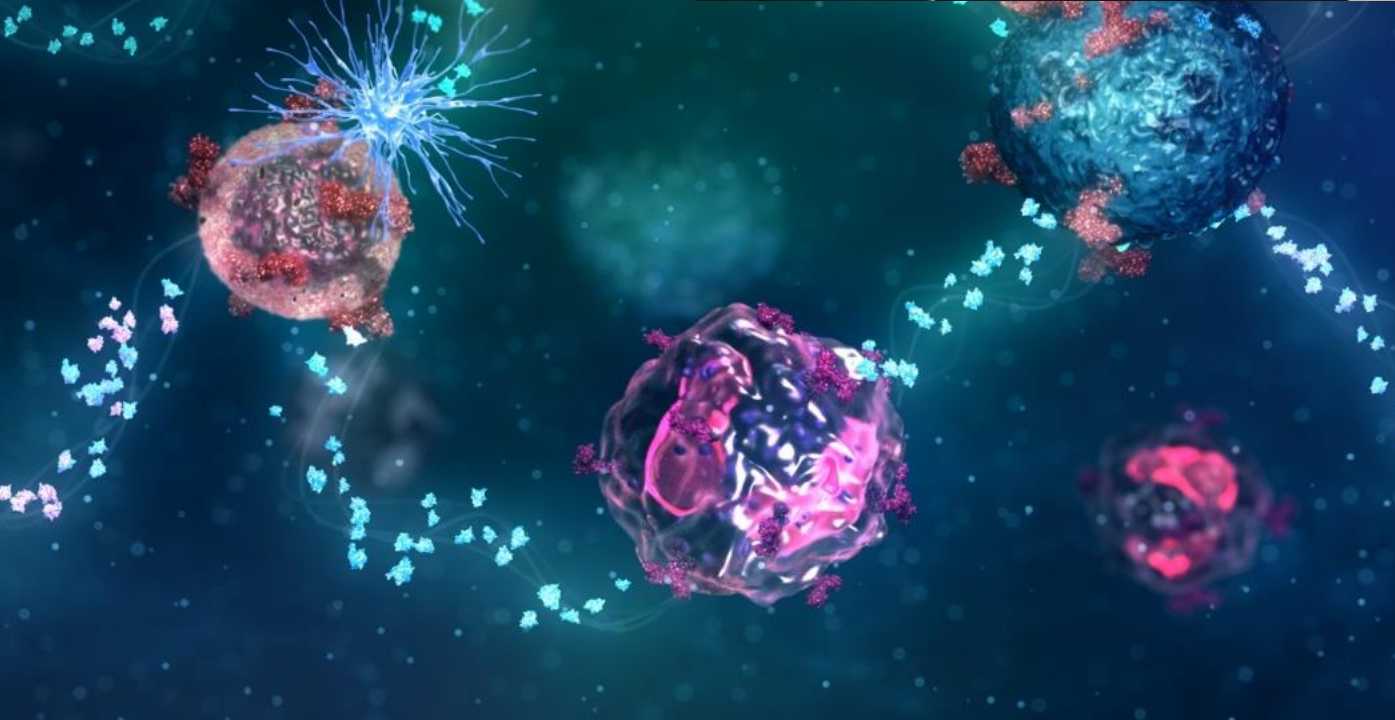
AZD9829
CD123

Phase I/II

AZD9592
EGFR/cMET

Phase I





BioPharmaceuticals

Ruud Dobber

BIOPHARMACEUTICALS BUSINESS

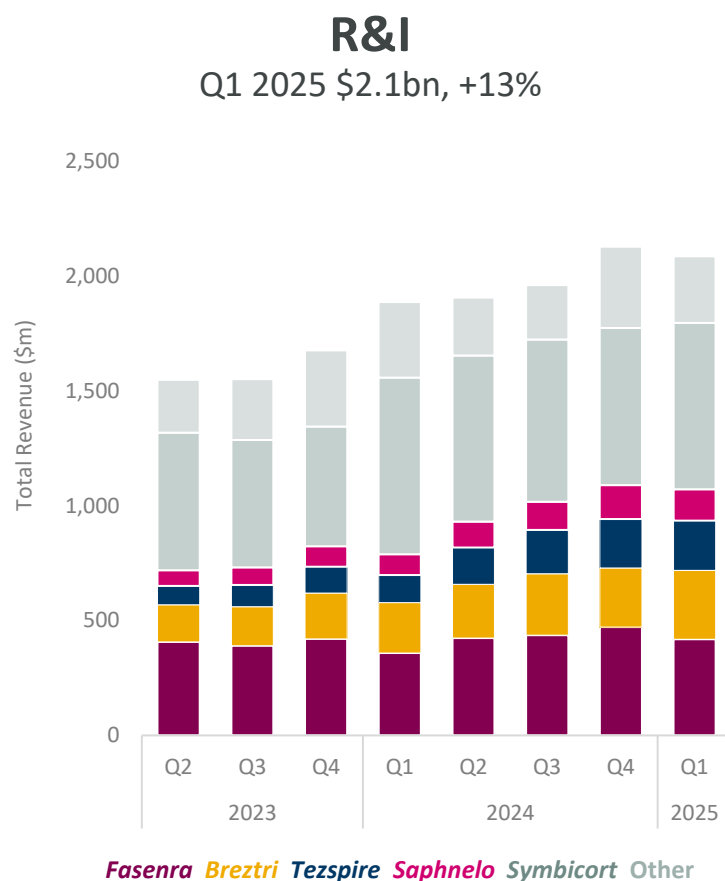
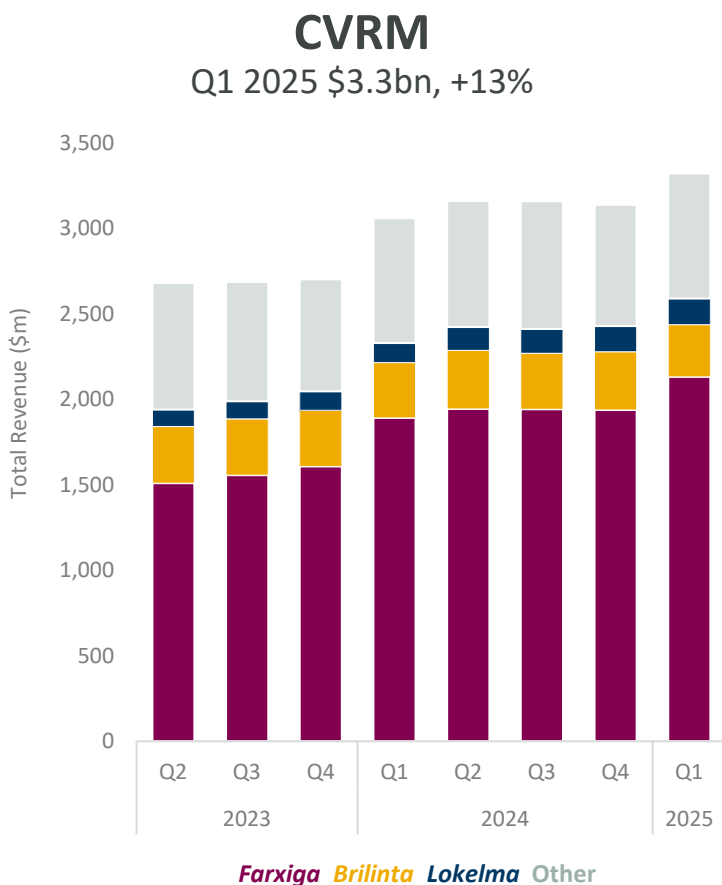
Sharon Barr

BIOPHARMACEUTICALS R&D



BioPharmaceuticals – Q1 2025

Total Revenue \$5.6bn, +12%, strong demand growth in expanding classes of medicines



Q1 2025: key dynamics

- **Farxiga** +17%, global demand growth mainly driven by CKD and HF
- **Lokelma** +38%, market leader in growing class
- **Brilinta** (4%), generic competition ex-US
- **Fasenna** +19%, sustained IL-5 leadership and EGPA launch
- **Tezspire** +85%, continued launch momentum
- **Breztri** +39%, fastest growing medicine in expanding FDC triple class
- **Saphnelo** +51%, increasing penetration in i.v. segment
- **V&I** (1%), **Beyfortus** >2x Total Revenue

Key regulatory approvals:

EU (*Wainzua* NEURO-TTTransform)



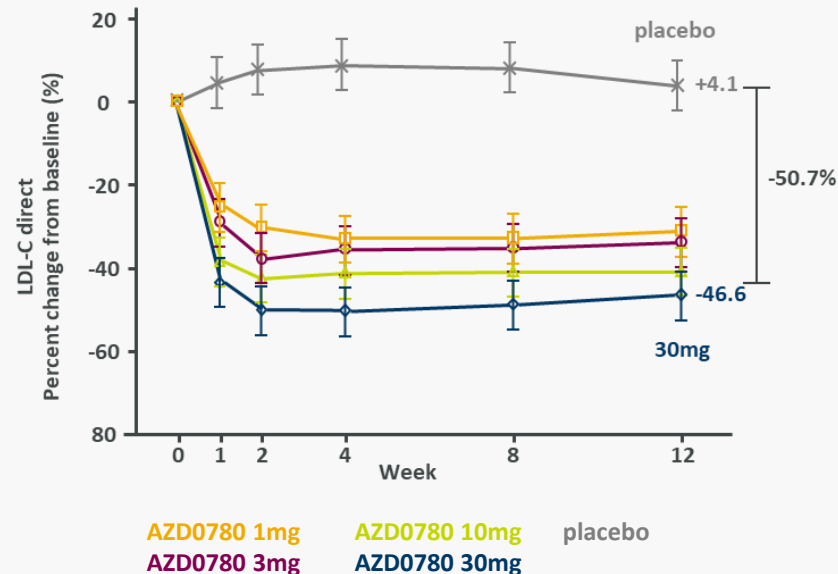
BioPharmaceuticals – potential best in class oral PCSK9 to address high unmet need in dyslipidaemia

AZD0780 (oPCSK9) Phase IIb PURSUIT met primary endpoint^{1,2}

≥50% LDL-C reduction
at 12 weeks³

84% of patients
met AHA/ACC guideline-
recommended LDL-C target

Oral small molecule
favourable tolerability, no food
effect or fasting requirement



Progressing into three Phase IIIs

AZURE-LDL

LDL-C reduction

AZURE-HeFH

Heterozygous familial
hypercholesterolemia
(HeFH)

AZURE-Outcomes

Cardiovascular
outcomes

\$5bn+ PYR⁵ with oral AZD0780 add-on and combination opportunities





Rare Disease

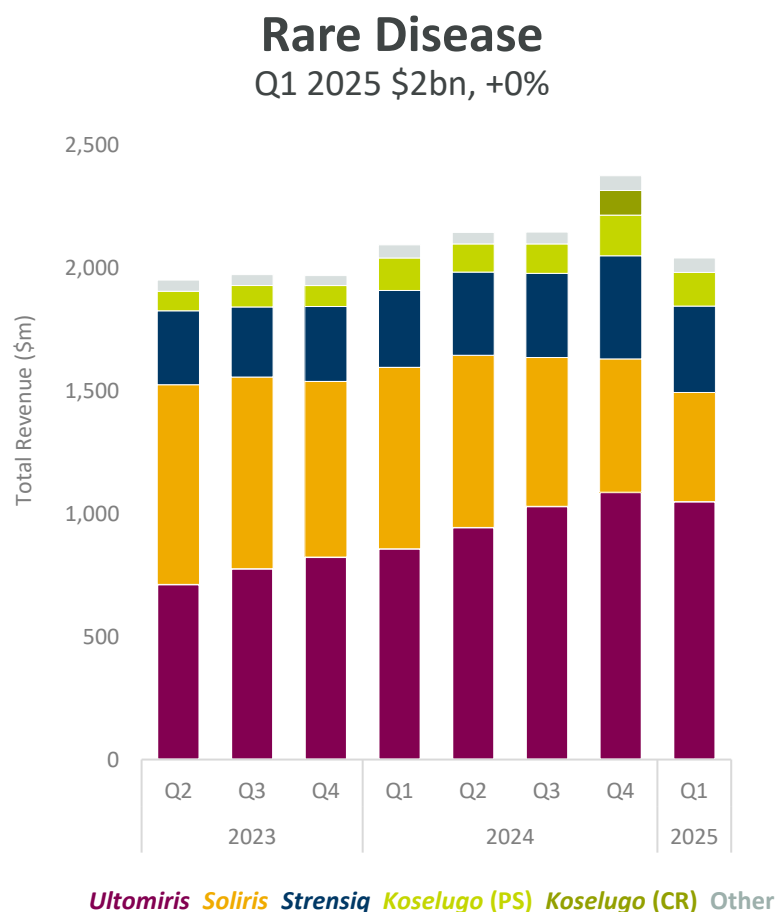
Marc Dunoyer

CHIEF EXECUTIVE OFFICER, ALEXION



Rare Disease – Q1 2025

Total Revenue \$2bn, confidence in future growth despite certain headwinds



Q1 2025: key dynamics

C5 Franchise

- **Ultomiris** +25%, demand growth across indications, offset by some competitive pressure in gMG and PNH and Part D redesign in US
- **Soliris** (38%), continued successful conversion to *Ultomiris*, additional impact from biosimilars in EU and unfavourable order timing in certain tender markets

Beyond Complement – market expansion and increased demand

- **Strensiq** +14%, continued global demand, partially offset by Part D redesign in US
- **Koselugo** +8%, continued global demand, partially offset by unfavourable order timing in certain tender markets
- **Key regulatory approvals:** JP (*Beyonttra*¹ ATTR-CM), CN (*Ultomiris* gMG)



Rare Disease – advancing *Ultomiris* in rare indications

Expanding into rare haematology and renal diseases

Potential blockbuster opportunity for *Ultomiris*
in HSCT-TMA across paededs and adults



TMA-314

Phase III demonstrated benefit
in paediatric patients

Clinically meaningful improvements

- Platelets
- LDH
- Urinary protein/creatinine ratio
- Overall survival

FDA BTD

TMA-313

Ongoing Phase III in
adult and adolescent patients

Data readout | H2 2025



~50k HSCT procedures per year¹

Advancing ambition
in rare renal diseases



aHUS
Launched

CSA-AKI
ARTEMIS | 2026

IgAN
ICAN | >2026

DGF
AWAKE | Initiating

pre-transplant

post-transplant

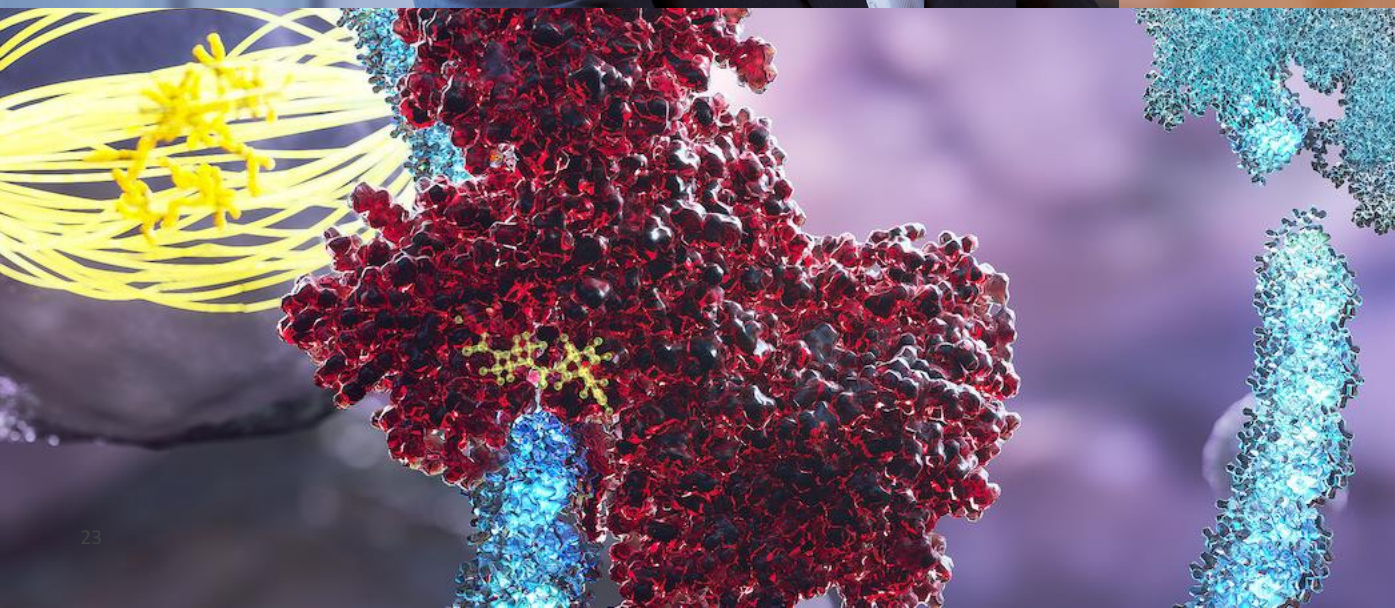
Ongoing or initiating Phase III *Ultomiris* trials





CEO Closing Remarks

Pascal Soriot
CHIEF EXECUTIVE OFFICER



2025 key Phase III readouts represent >\$10bn opportunity¹

H1 2025

H2 2025

First Phase III
data for 7 NMEs

- ✓ **SERENA-6**
camizestrant 1L *ESR1*m HR+/HER2-
breast cancer
- ✓ **CALYPSO**
eneboparatide hypoparathyroidism

- **CAEL-101/2**
anselamimab light chain amyloidosis
- **BaxHTN**
baxdrostat uncontrolled or
resistant hypertension
- **LATIFY**
ceralasertib Post-IO NSCLC
- **HICKORY/CHESTNUT**
efzimfotase alfa HPP
- **ALXN1720-MG-301**
gefurulimab gMG

Key indication
expansion
opportunities

- ✓ **DESTINY-Breast09**
Enhertu 1L HER2+
breast cancer
- **DESTINY-Breast11**
Enhertu early-stage
HER2+ breast
- **TROPION-Breast02**
Datroway 1L TNBC
- **KALOS/LOGOS**
Breztri uncontrolled
asthma
- **POTOMAC**
Imfinzi NMIBC

- **AVANZAR**
Datroway + Imfinzi 1L NSCLC
- **RESOLUTE**
Fasenra COPD
- **TMA-313**
Ultomiris adult HSCT-TMA
- **VOLGA**
Imfinzi cis-ineligible/refuse
MIBC
- **DESTINY-Breast05**
Enhertu early-stage
HER2+ breast



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



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 Fredrickson
EVP, ONCOLOGY
HAEMATOLOGY BUSINESS



Sharon Barr
EVP, BIOPHARMACEUTICALS R&D

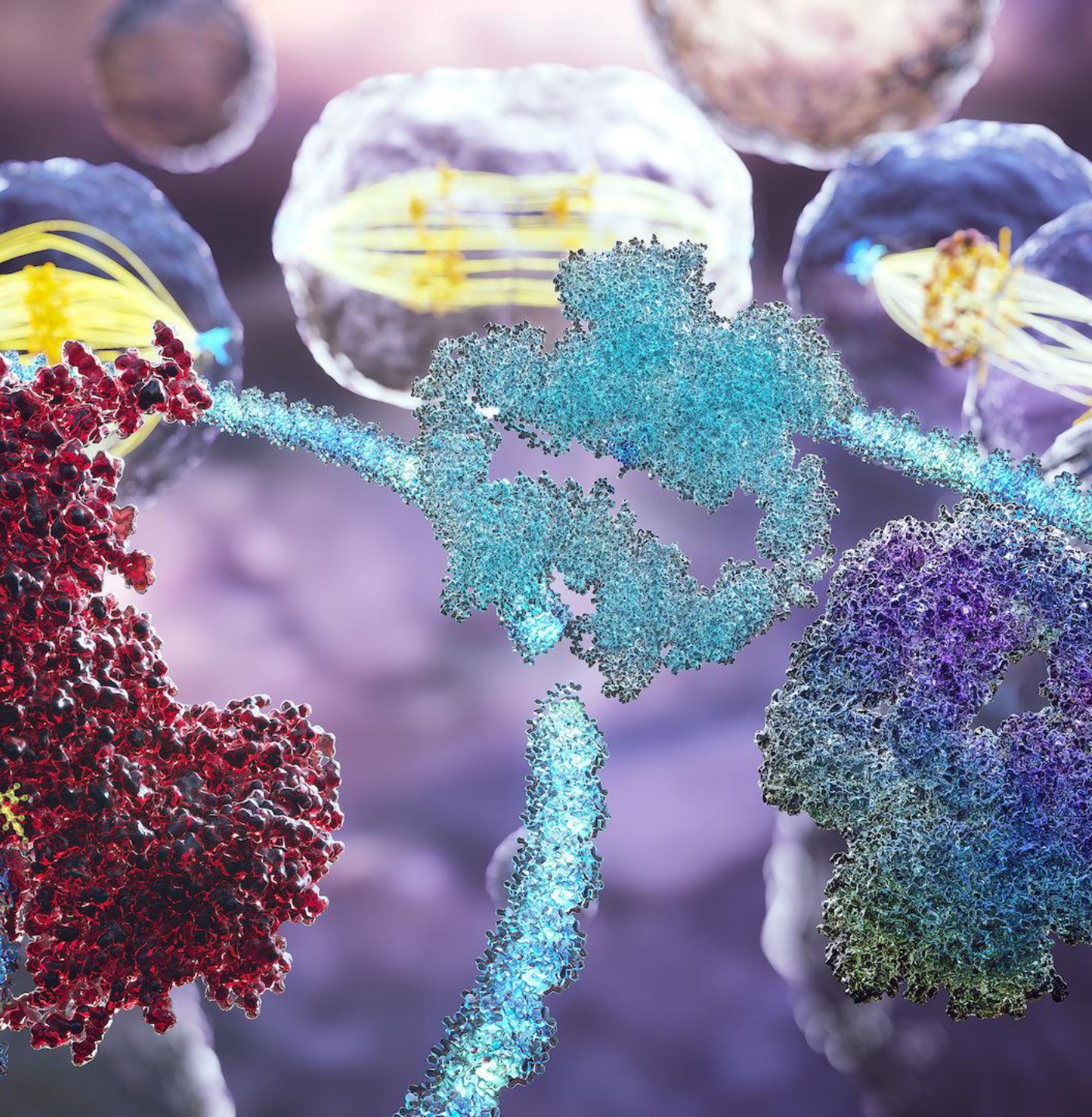


Ruud Dobber
EVP, BIOPHARMACEUTICALS
BUSINESS



Iskra Reic
EVP, INTERNATIONAL





Appendix



Appendix – AstraZeneca P&L reference table







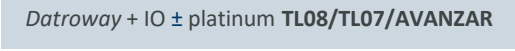



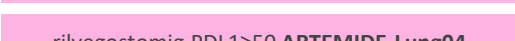






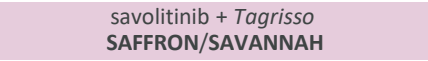


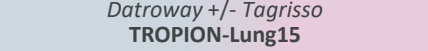
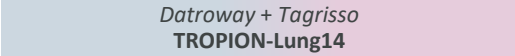


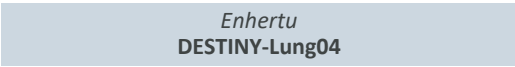

P&L line-item definitions

	P&L line-item definition
Product Sales	<ul style="list-style-type: none"> • Recognises sales from territories where Group has lead commercialisation • Recognises supply of <i>Beyfortus</i> to Sanofi
Alliance Revenue	<ul style="list-style-type: none"> • 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	<ul style="list-style-type: none"> • The sum of Product Sales and Alliance Revenue
Collaboration Revenue	<ul style="list-style-type: none"> • 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	<ul style="list-style-type: none"> • Sum of Product Sales, Alliance Revenue and Collaboration Revenue
Gross Margin	<ul style="list-style-type: none"> • Calculated by dividing Gross Profit by Total Revenue
Other operating income & expense	<ul style="list-style-type: none"> • 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 Operating margin	<ul style="list-style-type: none"> • Defined as Core Operating profit as a percentage of Total Revenue



AstraZeneca in Lung Cancer (NSCLC)

Overview of commercial portfolio and pivotal trials

		Resectable		Unresectable		Metastatic				
		Stg. I	Stg. II-III	Stg. I-II	Stg. III	1L		2L	3L+	
Est. epi (G7, 2025)		~150K	~70K	~50K	~100K	~310K		~190K	~70K	
IO sensitive ~70%		<i>Imfinzi</i> AEGEAN		 <i>SBRT → Imfinzi / Tagrisso</i> PACIFIC-4	 CRT → <i>Imfinzi</i> PACIFIC	 <i>Imfinzi</i> + <i>Imjudo</i> + CTx POSEIDON		<i>Imfinzi</i> + ceralasertib LATIFY		
	 rilvegostomig ± <i>Datroway</i> TROPION-Lung12				 <i>Imfinzi</i> combos PACIFIC-8, -9 improvements across PD-L1 spectrum	 <i>Datroway</i> + IO ± platinum TL08/TL07/AVANZAR				
						 rilvegostomig ± <i>Datroway</i> TROPION-Lung10				
						 rilvegostomig + CTx PDL1≥1 SQ ARTEMIDE-Lung02				
						 rilvegostomig + CTx PDL1≥1 NSQ ARTEMIDE-Lung03				
						 rilvegostomig PDL1≥50 ARTEMIDE-Lung04				
				 volrustomig + CTx eVOLVE-Lung02		 <i>Datroway</i> TROPION-Lung17 NSQ TROP2-NMR+				
EGFRm ~16-19%	 <i>Tagrisso</i> ADAURA2		 <i>Tagrisso</i> ADAURA	 CRT → <i>Tagrisso</i> LAURA	 <i>Tagrisso</i> FLAURA		 savolitinib + <i>Tagrisso</i> SAFFRON/SAVANNAH		 <i>Datroway</i> TROPION-Lung05	
					 <i>Tagrisso</i> + CTx FLAURA-2		 <i>Datroway</i> +/- <i>Tagrisso</i> TROPION-Lung15			
								 <i>Datroway</i> + <i>Tagrisso</i> TROPION-Lung14		 AZD9592 (EGFR/cMET ADC) EGRET
Other tumour drivers ~12%					 CRT → <i>Imfinzi</i> PACIFIC					
<i>HER2m</i> ~2%						 <i>Enhertu</i> DESTINY-Lung04		 <i>Enhertu</i> DESTINY-Lung02		



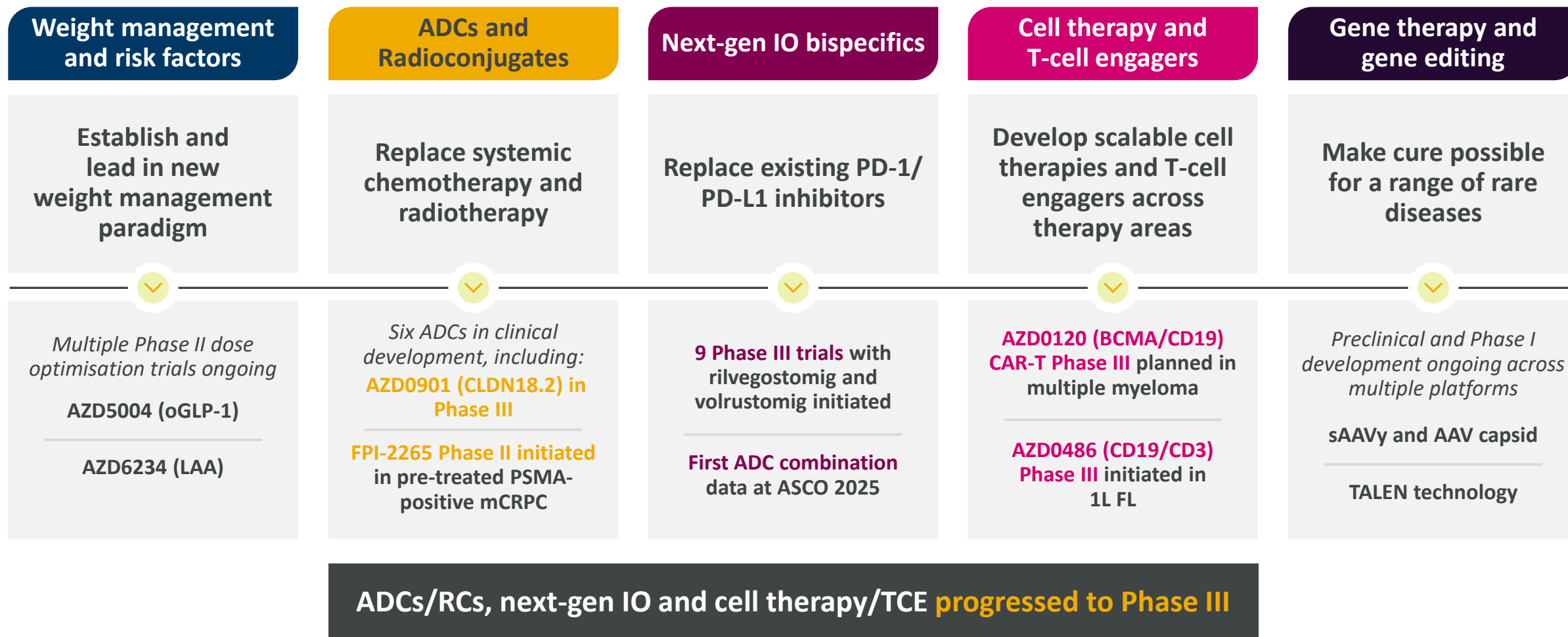
AstraZeneca in Breast Cancer

Overview of commercial portfolio and pivotal trials

	Early			Metastatic			
	Neoadjuvant	Adjuvant		1st line	2nd line	3rd line	4th line +
Est. epi (G7, 2025)	540k			135k	100k	75k	60k
HER2-positive 15-20%	<i>Enhertu</i> + THP DESTINY-Breast11	NST → residual disease → <i>Enhertu</i> DESTINY-Breast05		<i>Enhertu</i> ± pertuzumab DESTINY-Breast09	<i>Enhertu</i> DESTINY-Breast03	<i>Enhertu</i> DESTINY-Breast01/02	
HR-positive 65-75%		Good outcomes with current SoC for low-risk patients	RECURRENCE	camizestrant + palbociclib SERENA-4	<i>Truqap</i> + <i>Faslodex</i> CAPitello291 <i>PIK3CA, AKT1, PTEN alt.40%</i>	<i>Datroway</i> TROPION-Breast01	
		CTx → camizestrant ± abemaciclib CAMBRIA-2		AI + CDK4/6i → camizestrant + CDK4/6i SERENA-6 <i>ESR1m 35%</i>	<i>Enhertu</i> DESTINY-Breast06 HER2-low (1+, 2+) 60% HER2-ultralow (0-1+) 25%	<i>Enhertu</i> DESTINY-Breast04 HER2-low (1+, 2+) 60%	
		CTx → AI ± CDK4/6i 2-5 yrs → camizestrant CAMBRIA-1		<i>Truqap</i> + <i>Faslodex</i> + CDK4/6i CAPitello292			
TNBC 10-15%	<i>Datroway</i> + <i>Imfinzi</i> TROPION-Breast04	NST → residual disease → <i>Datroway</i> ± <i>Imfinzi</i> TROPION-Breast03		saruparib + camizestrant EvoPAR-Breast01 <i>tBRCAm, PALB2m 9%</i>	<i>Enhertu</i> DESTINY-Breast04 HER2-low (1+, 2+) 35%		HER2-low (1+, 2+) 35%
				<i>Datroway</i> + <i>Imfinzi</i> TROPION-Breast05 PD-L1+ 40%			
				<i>Datroway</i> TROPION-Breast02 PD-L1- 60%			
gBRCAm 5% of HR-positive 15% of TNBC		CTx → <i>Lynparza</i> OlympiA		<i>Lynparza</i> OlympiAD			



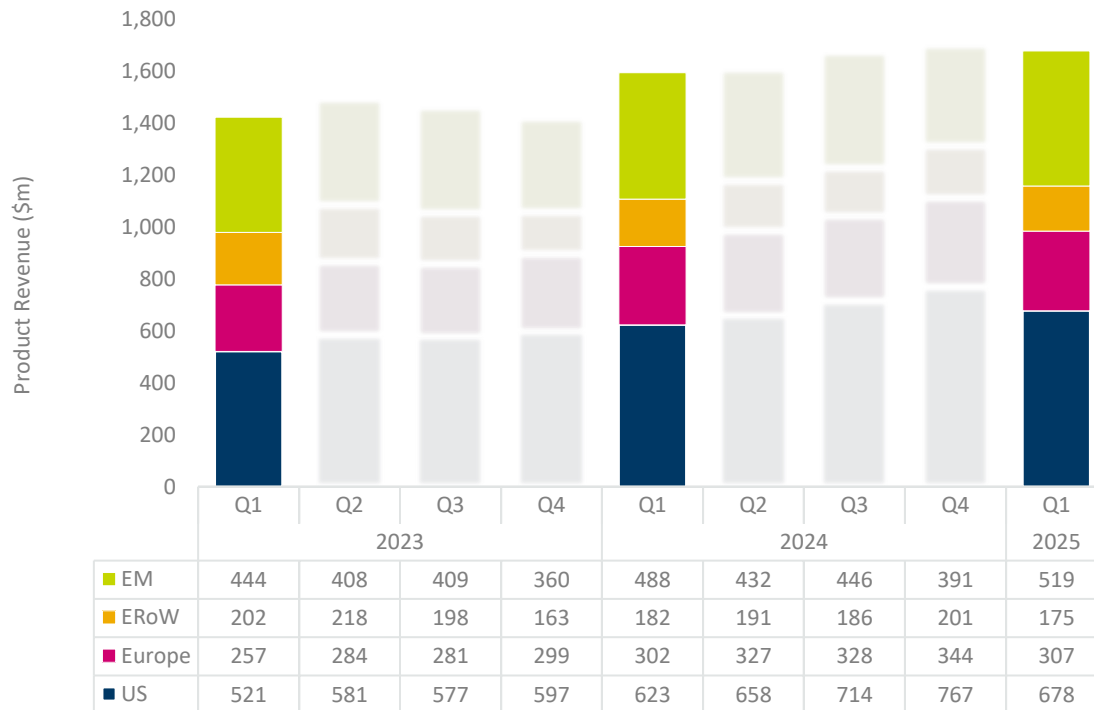
Appendix – Significant progress with transformative technologies to drive 2030+ growth



Oncology

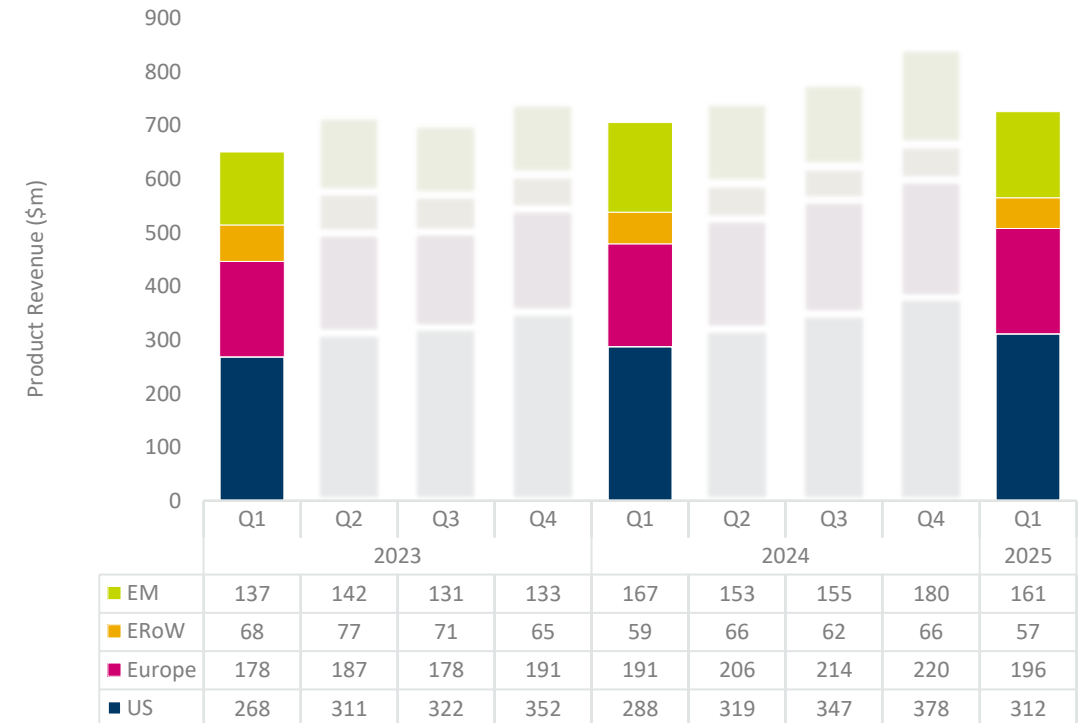
Tagrisso

8% growth at CER to \$1,679m in Q1 2025



Lynparza

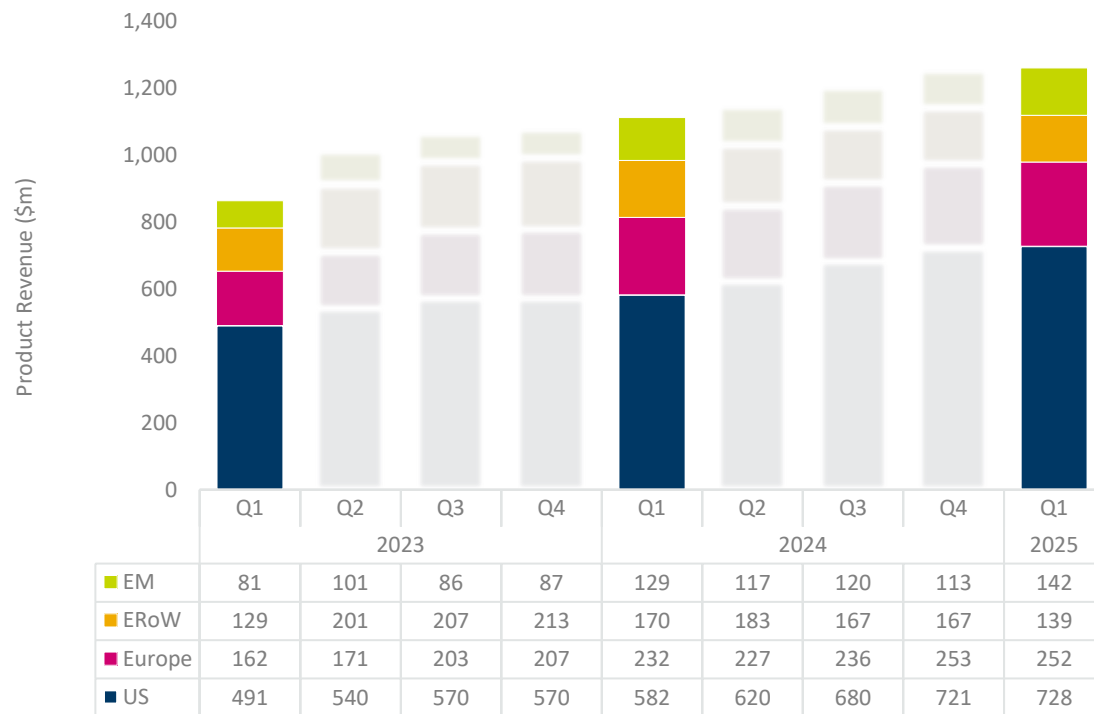
5% growth at CER to \$726m in Q1 2025



Oncology

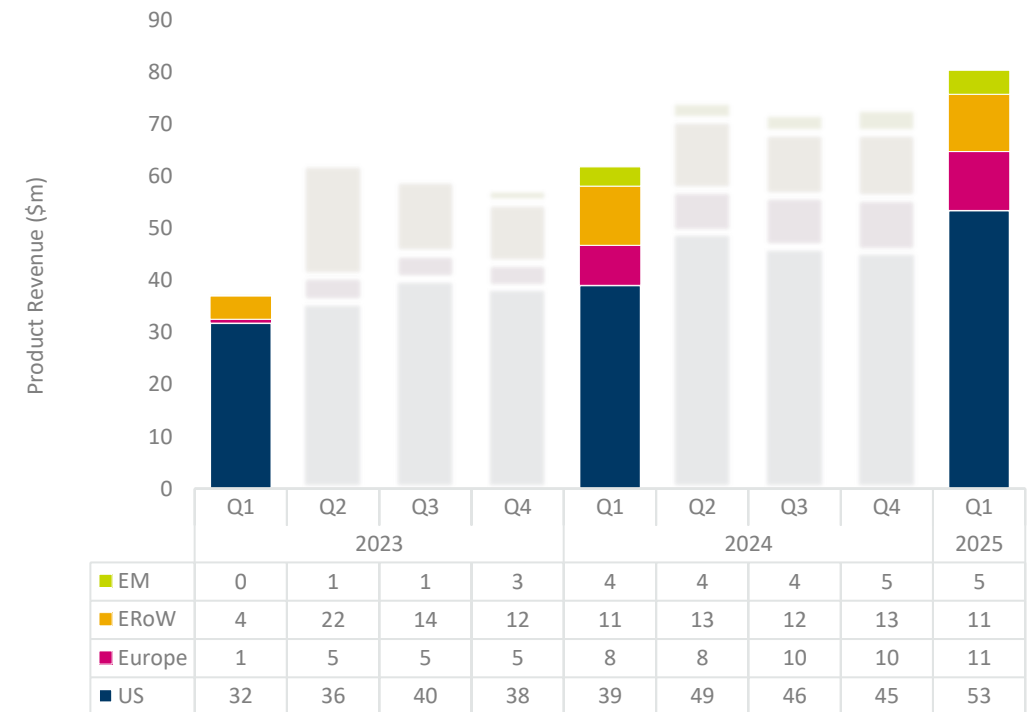
Imfinzi

16% growth at CER to \$1,261m in Q1 2025



Imjudo

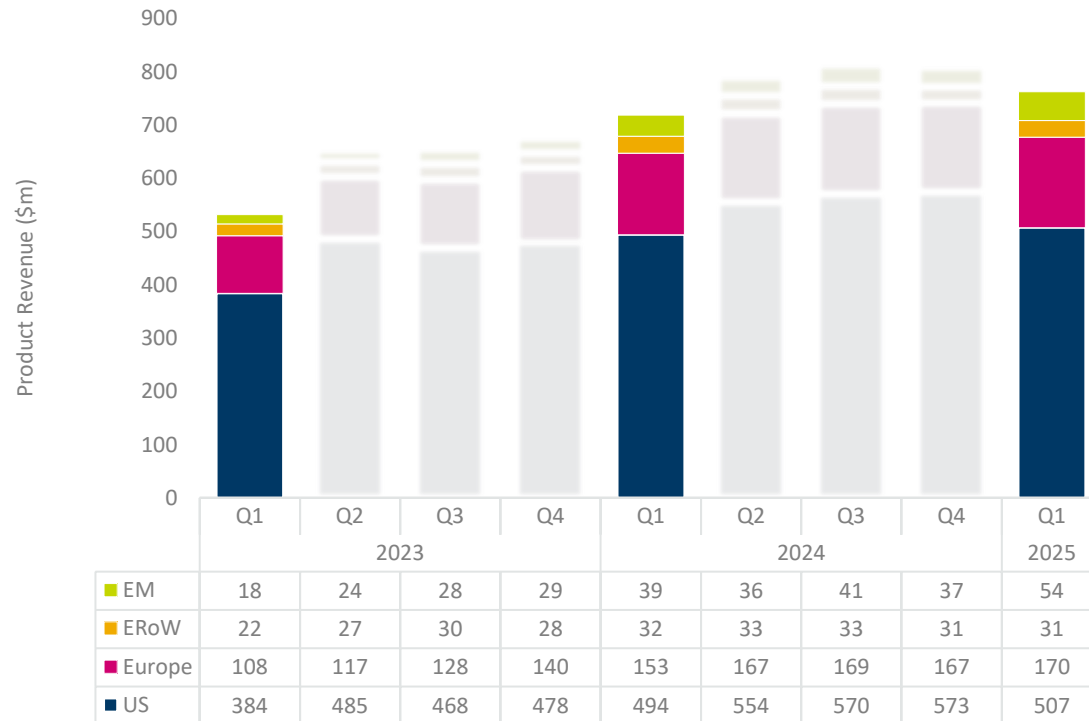
33% growth at CER to \$80m in Q1 2025



Oncology

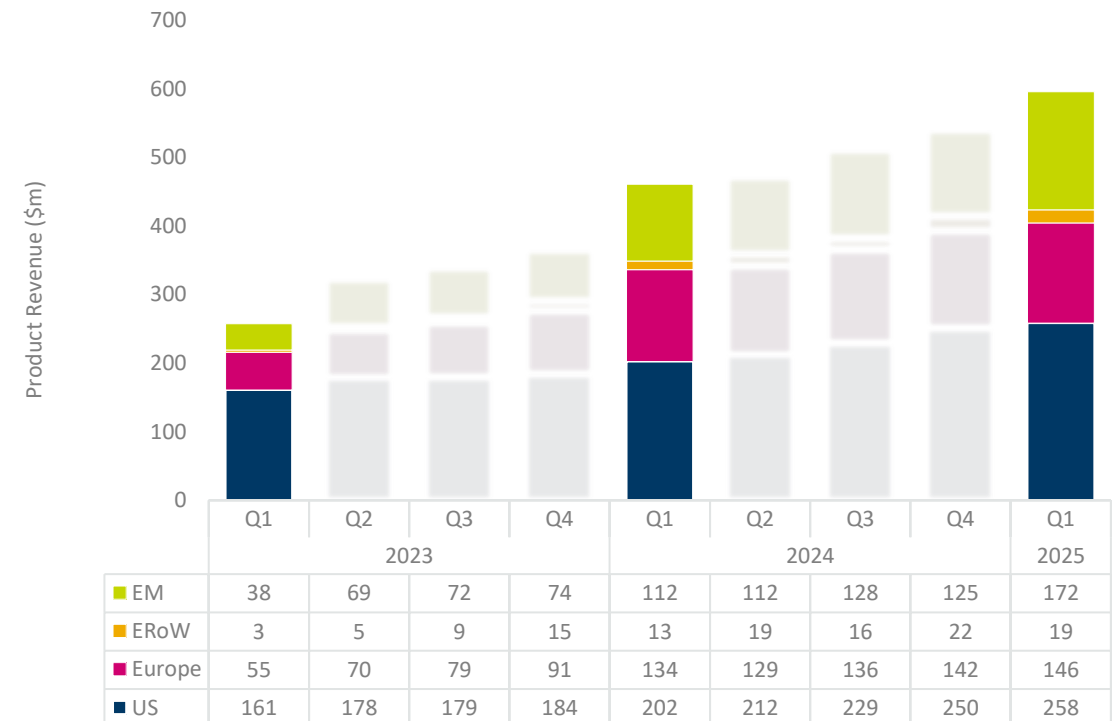
Calquence

8% growth at CER to \$762m in Q1 2025



Enhertu

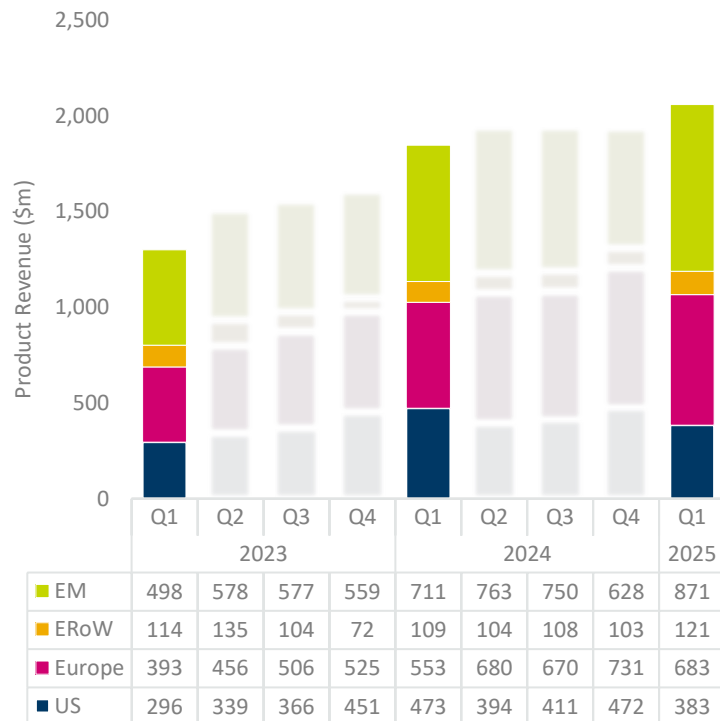
34% growth at CER to \$596m in Q1 2025



BioPharmaceuticals: Cardiovascular, Renal & Metabolism

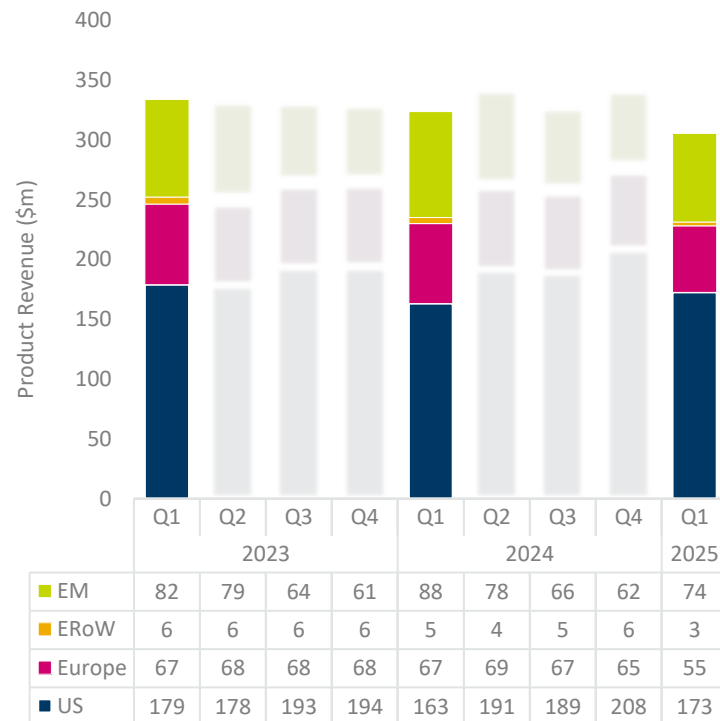
Farxiga

16% growth at CER to \$2,058m in Q1 2025



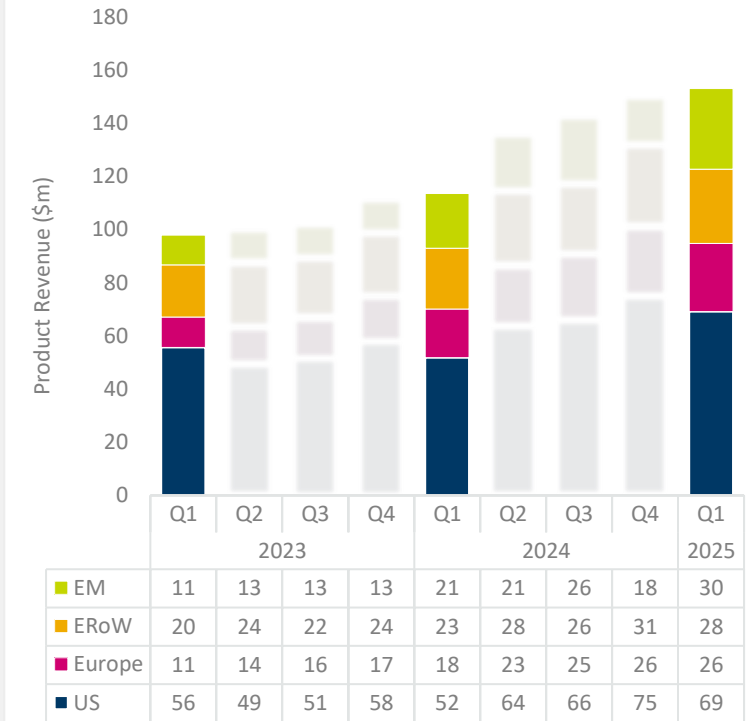
Brilinta

4% decrease at CER to \$305m in Q1 2025



Lokelma

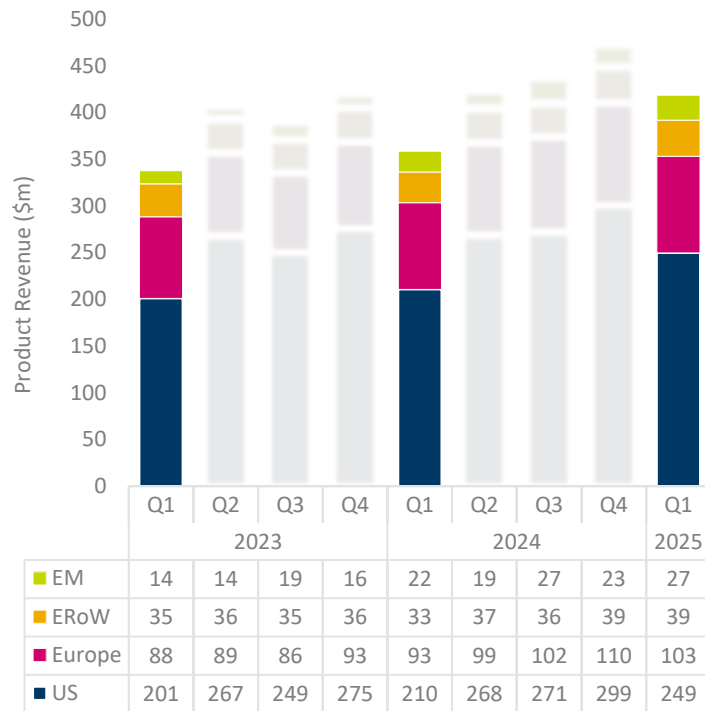
38% growth at CER to \$153m in Q1 2025



BioPharmaceuticals: Respiratory & Immunology

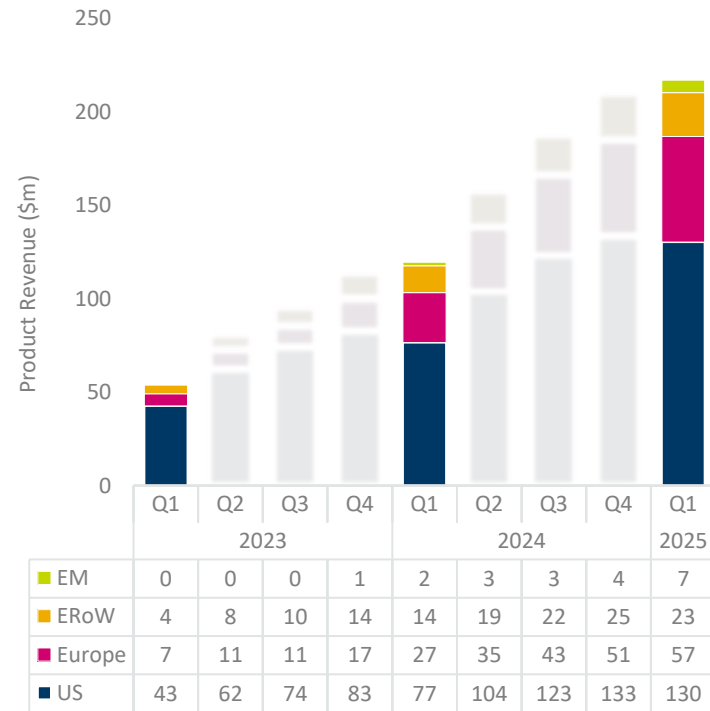
Fasenra

19% growth at CER to \$418m in Q1 2025



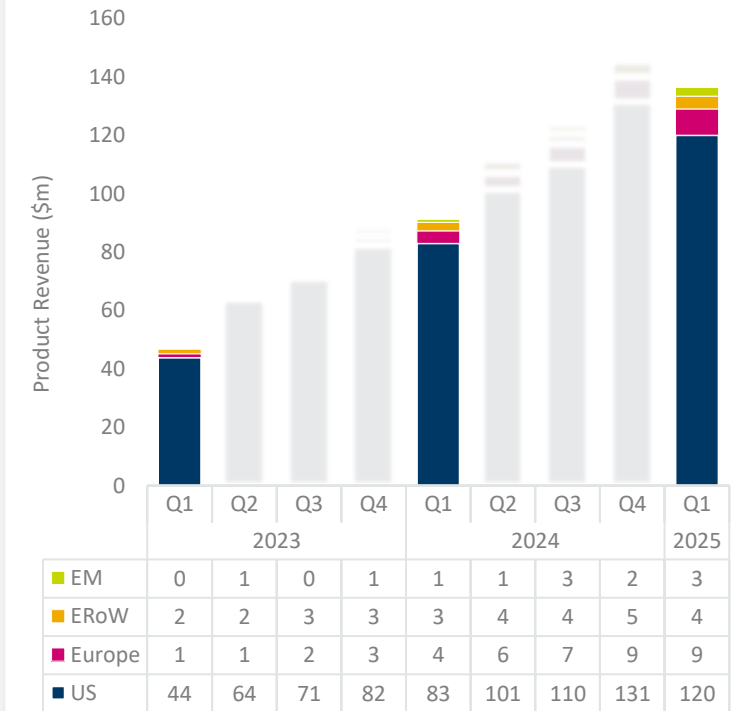
Tezspire

85% growth at CER to \$217m in Q1 2025



Saphnelo

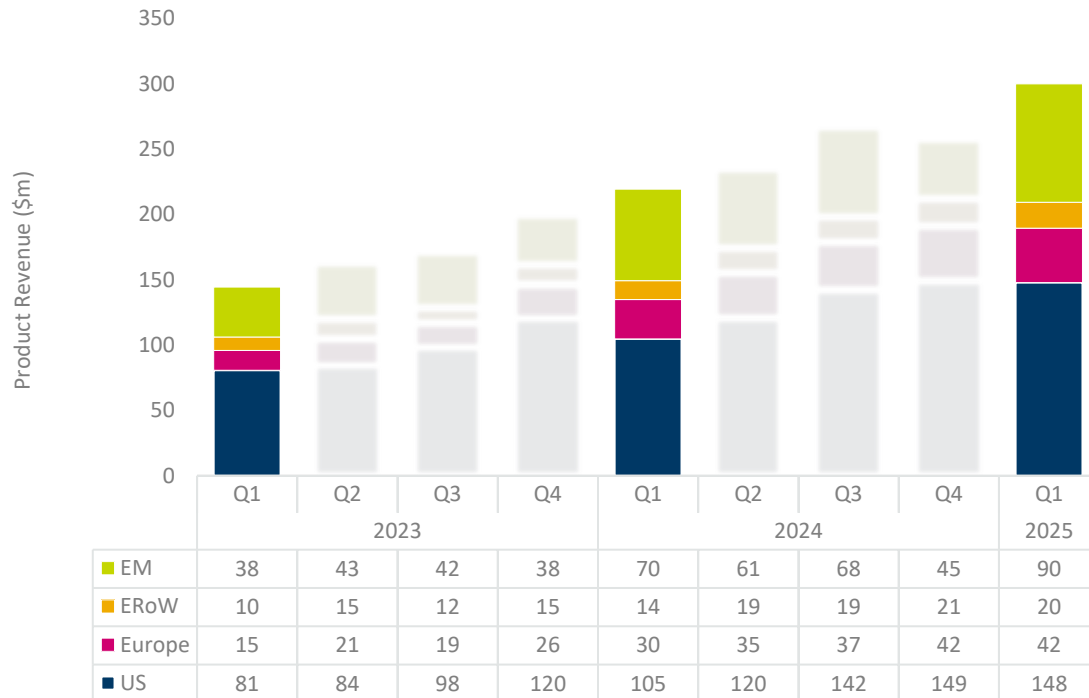
51% growth at CER to \$136m in Q1 2025



BioPharmaceuticals: Respiratory & Immunology

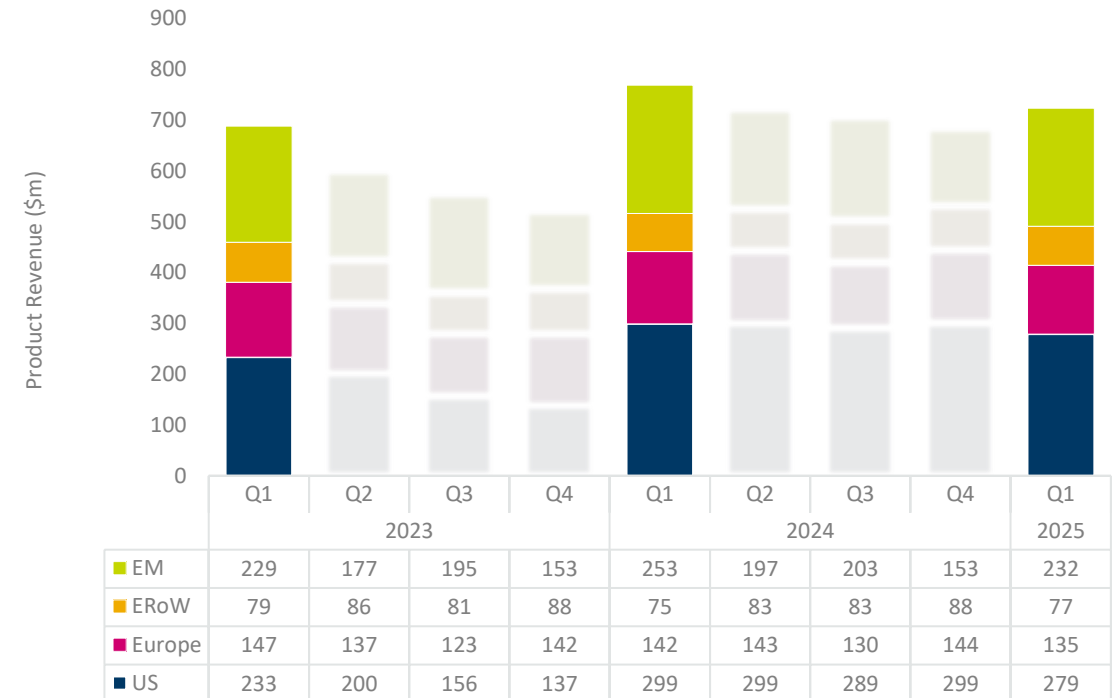
Breztri

39% growth at CER to \$300m in Q1 2025



Symbicort

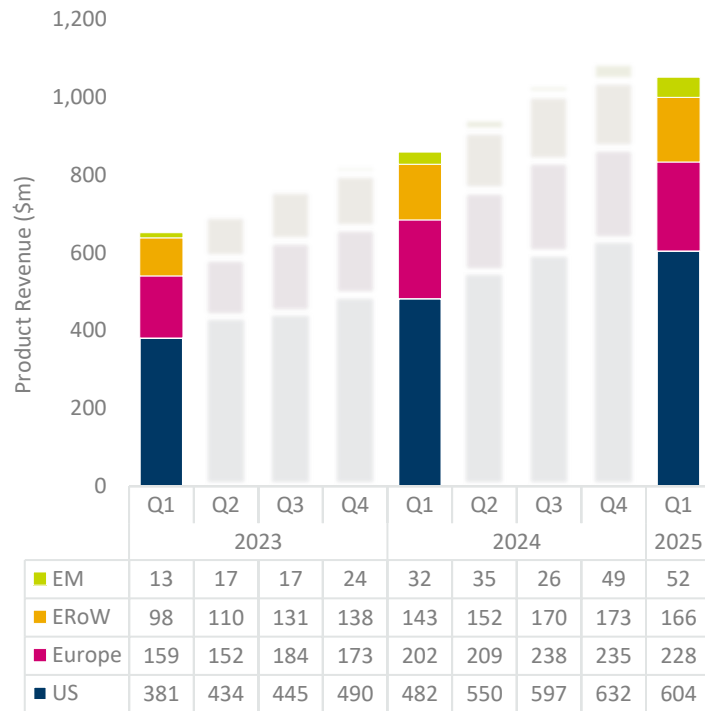
3% decrease at CER to \$723m in Q1 2025



Rare Disease

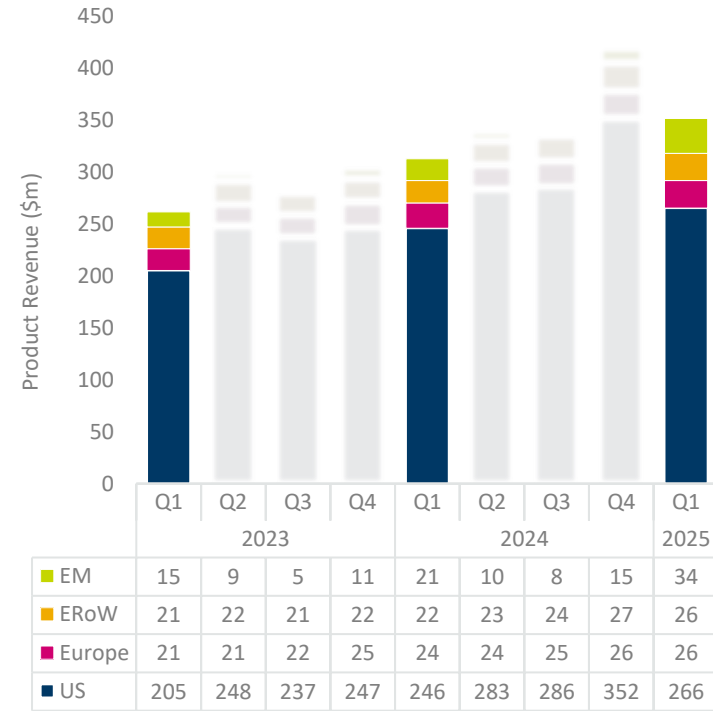
Ultomiris

25% growth at CER to \$1,050m in Q1 2025



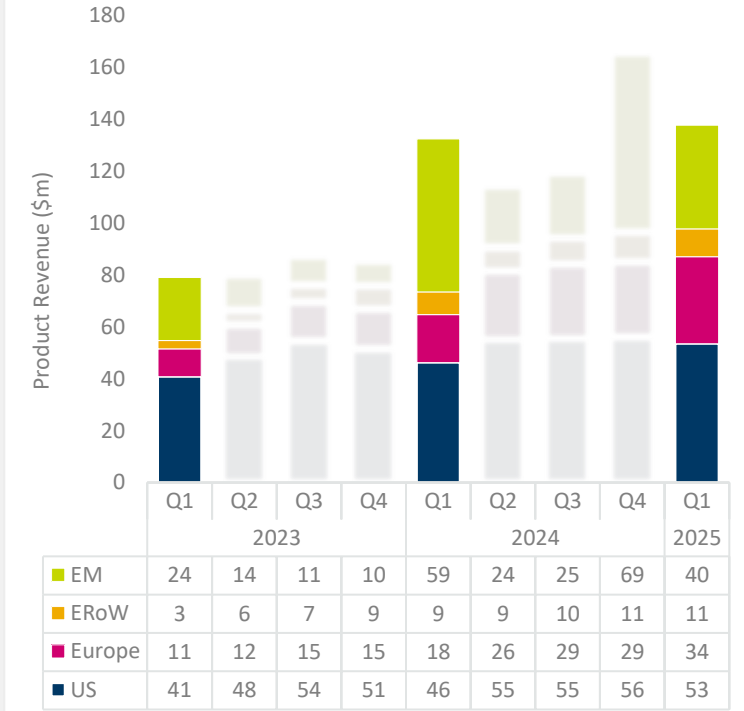
Strensiq

14% growth at CER to \$352m in Q1 2025



Koselugo

8% growth at CER to \$138m in Q1 2025



Glossary

1L, 2L, 3L	first-, second-, third-line	FLOT	fluorouracil, leucovorin, oxaliplatin and docetaxel	oPCSK9	oral protein convertase subtilisin/kexin type 9
AAV	adeno-associated virus	FRα	folate receptor alpha	ORR	overall response rate
ACC	American College of Cardiology	FX	foreign exchange	OS	overall survival
ADC	antibody-drug conjugate	G7	US, Japan, EU5	P&L	Profit & Loss
AHA	American Heart Association	gBRCAm	germline BRCA-mutated breast cancer	PALB2m	partner and localizer of BRCA2
aHUS	atypical haemolytic uraemic syndrome	GC	gastric cancer	PARPi	poly-ADP ribose polymerase inhibitor
AI	aromatase inhibitor	GEJ	gastroesophageal junction	PCSK9	protein convertase subtilisin/kexin type 9
AKT1	AKT serine/threonine kinase 1	GEJC	gastroesophageal junction cancer	PD-1	programmed cell death protein-1
B7H4	B7 homolog 4	GI	gastrointestinal	PDL1	programmed death-ligand 1
BCMA	B-cell maturation antigen	gMG	generalised myasthenia gravis	PD-L1	programmed cell death ligand 1
BTKi	Bruton's tyrosine kinase	GPRC5D	G protein-coupled receptor class C group 5 member D	PFS	progression free survival
CAR-T	chimeric antigen receptor T-cells	HeFH	heterozygous familial hypercholesterolemia	PIK3CA	phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit
CD123	interleukin-3 receptor alpha chain	HER2-	human epidermal growth factor receptor 2	PNH	paroxysmal nocturnal haemoglobinuria
CD19	cluster of differentiation 19	HER2+	human epidermal growth factor receptor 2-positive	PS	Product Sales
CD3	cluster of differentiation 3	HER2-low	human epidermal growth factor receptor 2-low	PSMA	prostate-specific membrane antigen
CDK4/6i	cyclin-dependent kinase 4/6 inhibitor	HER2-ultralow	human epidermal growth factor receptor 2-ultralow	PTEN	phosphatase and TENsin homolog deleted on chromosome 10
CER	constant exchange rates	HF	heart failure	PYR	Peak-Year Revenue
CFO	net cash inflow from operating activities	HPP	hypophosphatasia	R&D	Research & Development
cis	cisplatin	HR+	hormone receptor positive	R&I	Respiratory & Immunology
CKD	chronic kidney disease	HSCT	hematopoietic stem cell transplantation	RC	radioconjugate
CLDN18.2	Claudin-18.2	HSCT-TMA	hematopoietic stem cell transplantation-associated thrombotic microangiopathy	SERD	selective estrogen receptor degrader
CLL	chronic lymphocytic leukaemia	i.v.	intravenous	SG&A	Selling, General & Administrative
cMET	c-mesenchymal epithelial transition factor	IgAN	IgA nephropathy	SoC	standard-of-care
CN	China	IL-5	interleukin-5	SQ	squamous
COPD	chronic obstructive pulmonary disease	IO	immuno-oncology	Stg.	stage
CR	Collaboration Revenue	JP	Japan	tBRCAm	tumor BRCA mutation
CRT	chemoradiotherapy	LAA	long-acting amylin	TCE	T-cell engager
CSA-AKI	cardiac surgery-associated acute kidney injury	LDH	lactate dehydrogenase	THP	docetaxel, trastuzumab and pertuzumab
CTx	chemotherapy	LDL-C	low-density lipoprotein cholesterol	TL07	TROPION-Lung07
CVRM	Cardiovascular, Renal and Metabolism	mBC	metastatic breast cancer	TL08	TROPION-Lung08
Dato-DXd	Datroway	mCRPC	metastatic castration-resistant prostate cancer	TNBC	triple negative breast cancer
DGF	delayed graft function	MIIBC	muscle-invasive bladder cancer	TRAE	treatment-related adverse event
EBITDA	earnings before interest, tax, depreciation and amortisation	mPFS	median progression-free survival	TROP2	trophoblast cell surface antigen 2
EGFR	epidermal growth factor receptor	NME	new molecular entity	UK	United Kingdom
EGFRm	epidermal growth factor receptor-mutant	NMIBC	non-muscle invasive bladder cancer	US	United States
EPS	earnings per share	NMR	normalised membrane ratio	V&I	Vaccines & Immune Therapies
ERoW	Established Rest of World	NRDL	national reimbursement drug list		
ESR1m	estrogen receptor alpha-mutated	NSCLC	non-small cell lung cancer		
EU	Europe	NSQ	non-squamous		
FDC	fixed-dose combination	NST	neoadjuvant systemic treatment		
FL	follicular lymphoma	oGLP-1	oral glucagon-like peptide-1		



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