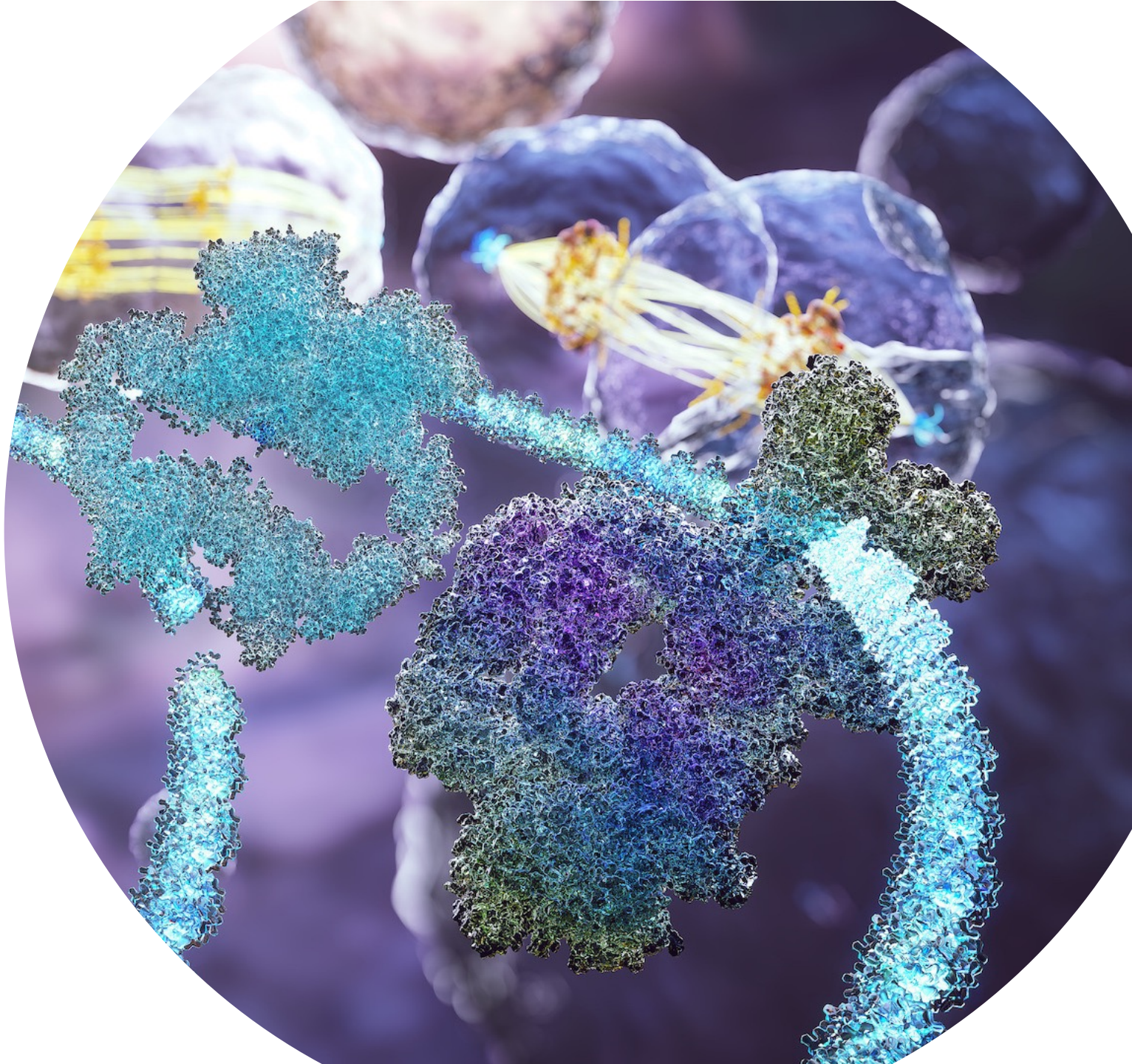




Q1 2023 Results

Conference call and webcast
for investors and analysts

27 April 2023



Forward-looking statements

In order, among other things, to utilise the 'safe harbour' provisions of the US Private Securities Litigation Reform Act of 1995, AstraZeneca (hereafter 'the Group') provides the following cautionary statement: 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 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 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 regulatory or ethical expectations on environmental impact, including climate change; 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-related risks to our products; the risk of failure to achieve strategic plans or meet targets or expectations; the risk of failure in financial control or the occurrence of fraud; the risk of unexpected deterioration in the Group's financial position; the impact that global and/or geopolitical events such as the COVID-19 pandemic and the Russia-Ukraine war 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 2023 results

Conference call agenda

CEO Opening Remarks

Pascal Soriot
Chief Executive Officer

Financial Results

Aradhana Sarin
Chief Financial Officer

Oncology

Dave Fredrickson
EVP, Oncology Business

Susan Galbraith
EVP, Oncology R&D

BioPharmaceuticals

Ruud Dobber
EVP, BioPharmaceuticals Business

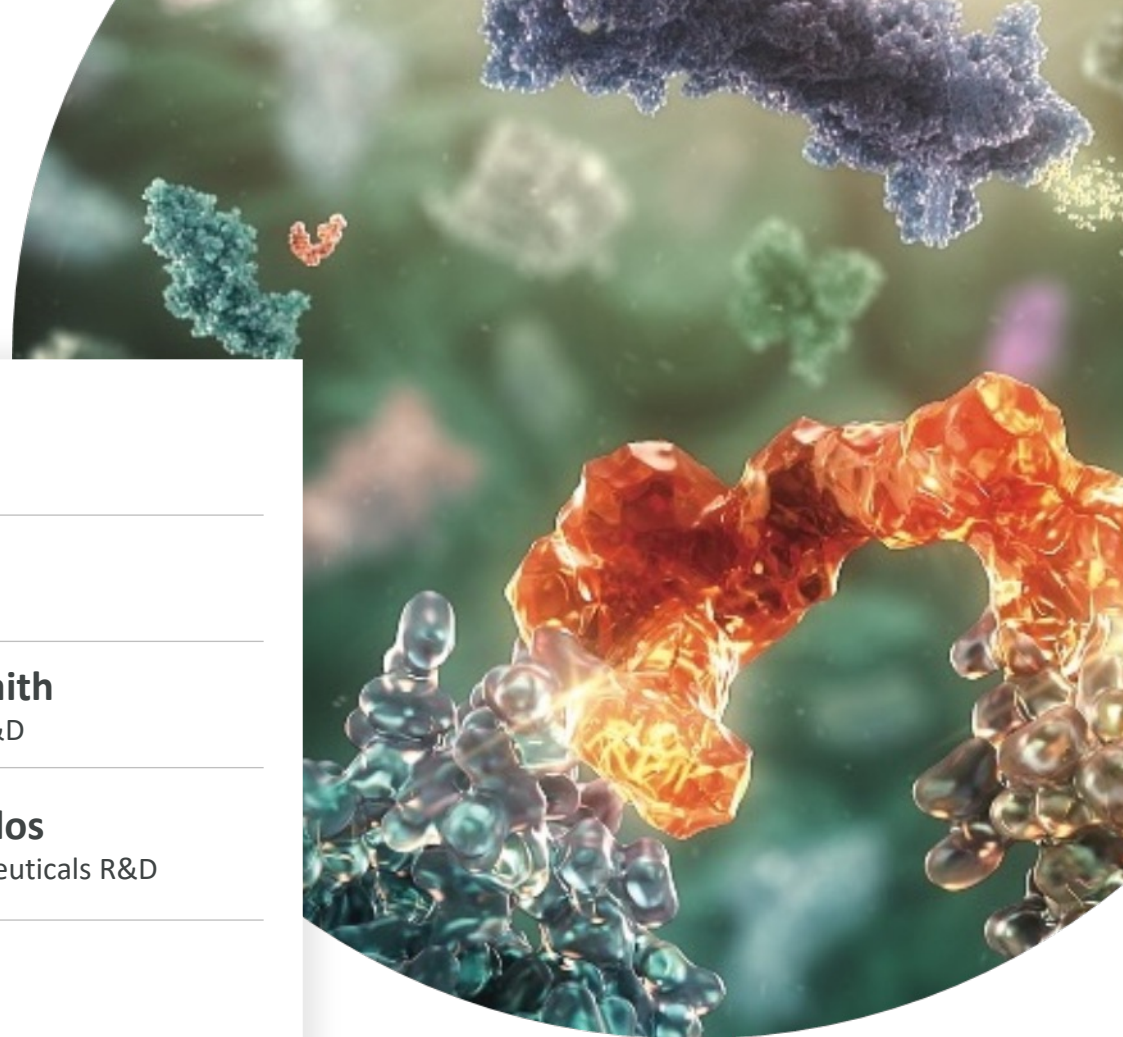
Mene Pangalos
EVP, BioPharmaceuticals R&D

Rare Disease

Marc Dunoyer
Chief Executive Officer, Alexion

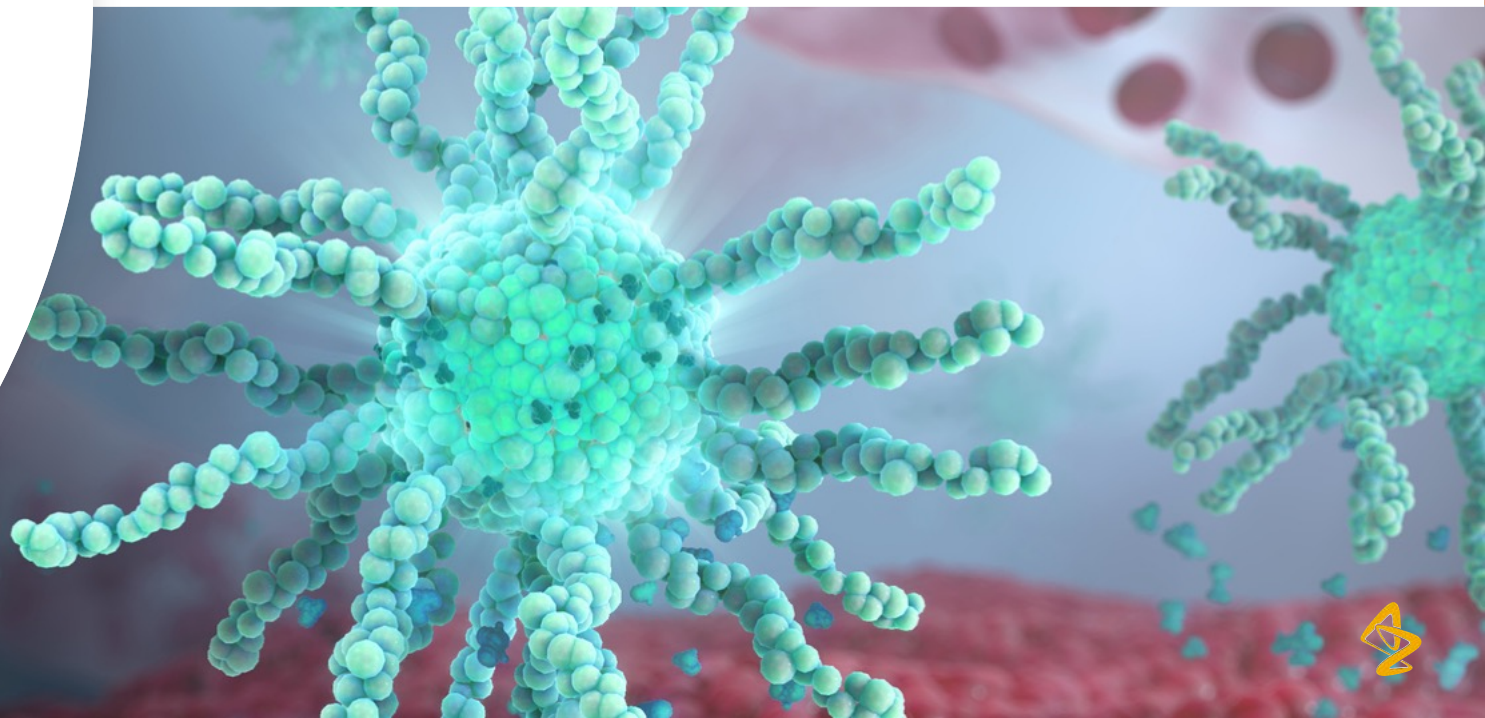
CEO Closing Remarks, Q&A

Pascal Soriot
Chief Executive Officer



CEO Opening Remarks

Pascal Soriot
CHIEF EXECUTIVE OFFICER

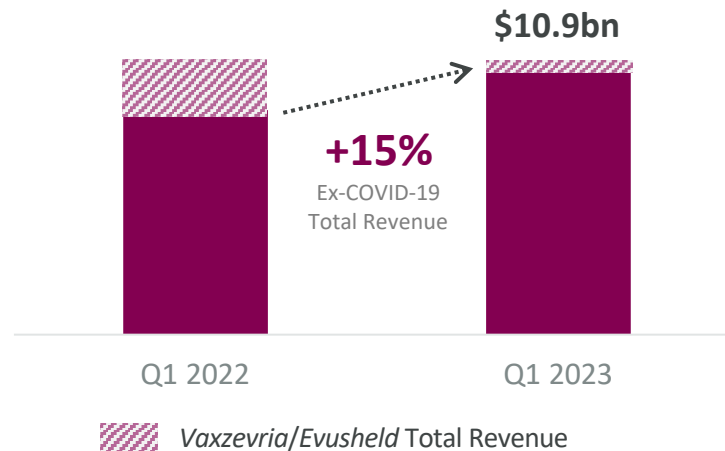


Q1 2023 – 15% growth from ex-COVID-19 medicines¹

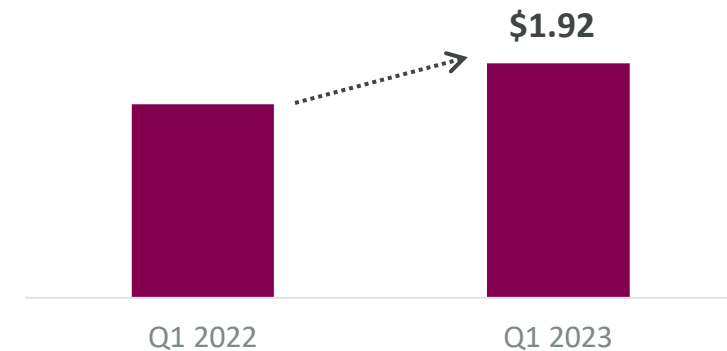
Stable Total Revenue, reiterating FY 2023 guidance

Growth in Oncology, CVRM, R&I and Rare Disease offset decline in COVID-19 medicines

Total Revenue | stable CER



Core EPS | +6% CER



Reiterating 2023 guidance: Core EPS to increase by a high single-digit to low double-digit %

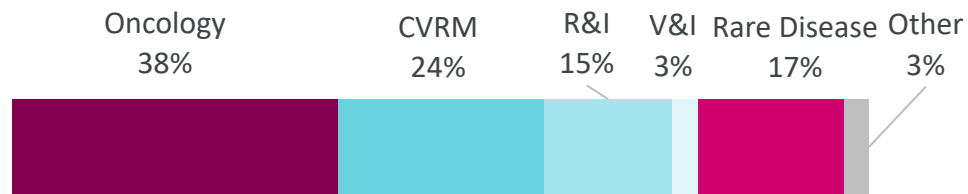
⁵ All growth rates at CER.

¹ Vaxzevria, Evusheld and AZD3152; CER = constant exchange rates; EPS = earnings per share.

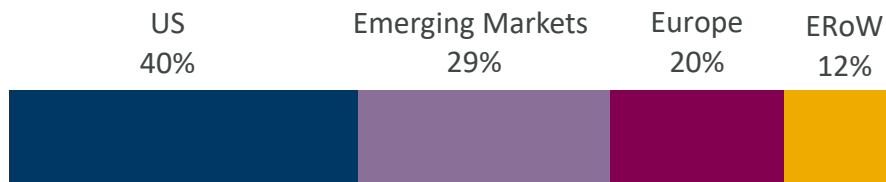


Q1 2023 – broad-based, diverse source of business

Q1 2023 | % Total Revenue by therapy area



Q1 2023 | % Total Revenue by geography



Strong growth across Oncology, CVRM, R&I and Rare Disease

Emerging Markets presence and strong growth

Value-enhancing business development strengthening growing pipeline

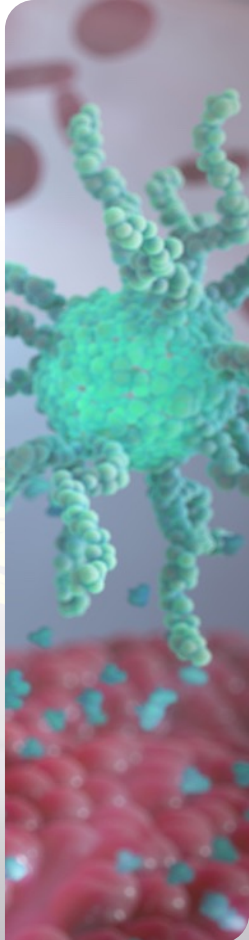
Increasing pipeline momentum

Industry-leading outlook underpinned by broad portfolio and geographic footprint



Accelerating our late-stage pipeline

Potential to initiate 30 Phase III trials, with six dosed to-date in 2023¹



Dato-DXd	Dato-DXd	Tezspire	AZD3152	camizestrant	Ultomiris
AVANZAR² 1L NSCLC	TROPION-Lung07² non-squamous 1L NSCLC	CROSSING EoE	SUPERNOVA COVID-19 prophylaxis	CAMBRIA-1 HR+/HER2- adjuvant BC	ARTEMIS CSA-AKI
Potential HLR: >2024	Potential HLR: >2024	Potential HLR: >2024	Potential HLR: H2 2023	Potential HLR: >2024	Potential HLR: >2024

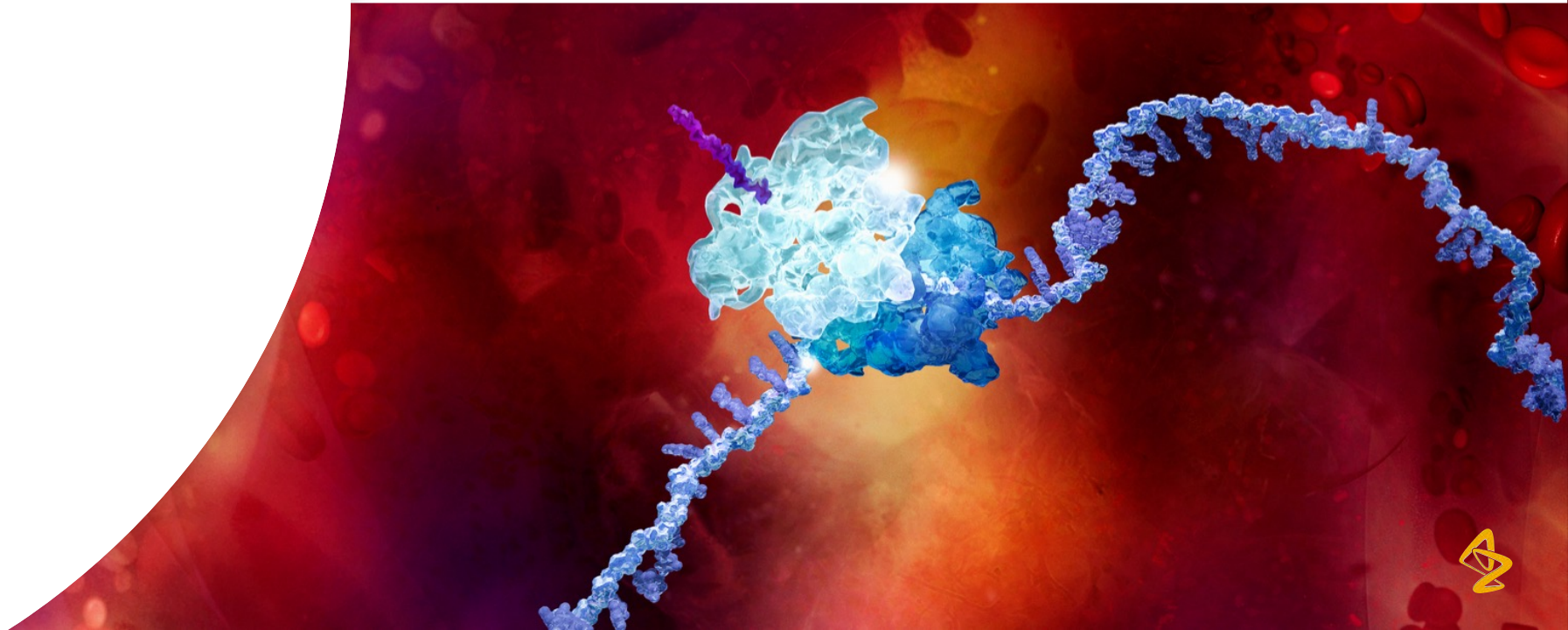
10 potential blockbuster opportunities from 30 Phase III trials planned in 2023

1. Phase III trial initiation defined as achievement of first patient dosed; 2. AVANZAR and TROPION-Lung07 Phase III first patient dosed ahead of Q4/FY 2022 results on 09 February 2023. NSCLC = non-small cell lung cancer; 1L = first-line (metastatic); EoE = eosinophilic esophagitis; HR+/HER2- = hormone receptor-positive/human epidermal growth factor receptor-negative; BC = breast cancer; CSA-AKI = cardiac surgery-associated acute kidney injury. Collaboration partners: Daiichi Sankyo (Dato-DXd), Amgen (Tezspire).

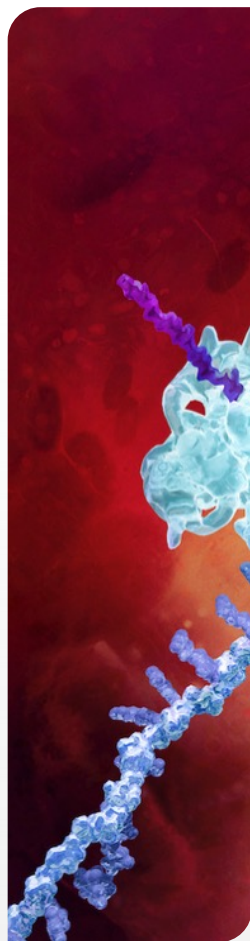


Financial Results

Aradhana Sarin
CHIEF FINANCIAL OFFICER



Q1 2023 – Reported profit and loss



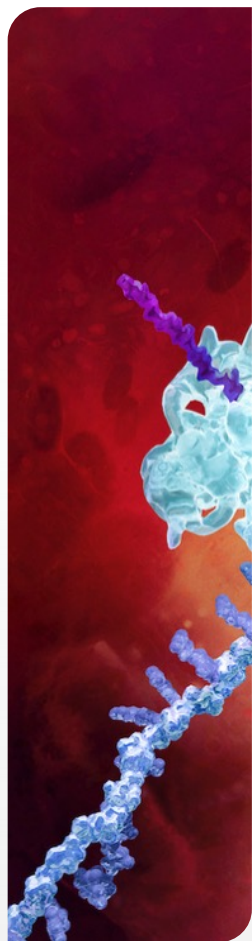
	Q1 2023 \$m	CER change %	% Total Revenue
Total Revenue	10,879	-	100
- <i>Product Sales</i>	10,566	1	97
- <i>Alliance Revenue</i>	286	90	3
- <i>Collaboration Revenue</i>	27	(89)	-
Gross margin	82.0%	+14 pp	
Total operating expense ¹	(6,804)	-	63
- R&D expense	(2,611)	28	24
- SG&A expense	(4,059)	(13)	37
Other operating income and expense	379	>3	3
Operating profit	2,549	>2x	23
Tax rate	20.2%		
Reported EPS	\$1.16	>4x	

⁹ Absolute values at actual exchange rates; changes at CER. Gross margin excludes the impact of Alliance and Collaboration Revenue and any associated costs, thereby reflecting the underlying performance of Product Sales.

¹ Total operating expenses include distribution, R&D and SG&A expenses. R&D = Research & Development; SG&A = Sales, General & Administrative; pp = percentage points; CER = constant exchange rates.



Q1 2023 – Core profit and loss



	Q1 2023 \$m	CER change %	% Total Revenue
Total Revenue	10,879	-	100
- Product Sales	10,566	1	97
- Alliance Revenue	286	90	3
- Collaboration Revenue	27	(89)	-
Gross margin	83.3%	+4 pp	
Total operating expense ¹	(5,488)	9	50
- R&D expense	(2,300)	10	21
- SG&A expense	(3,054)	8	28
Other operating income and expense	318	>3x	3
Operating profit	3,946	4	36
Tax rate	20%		
Core EPS	\$1.92	6	

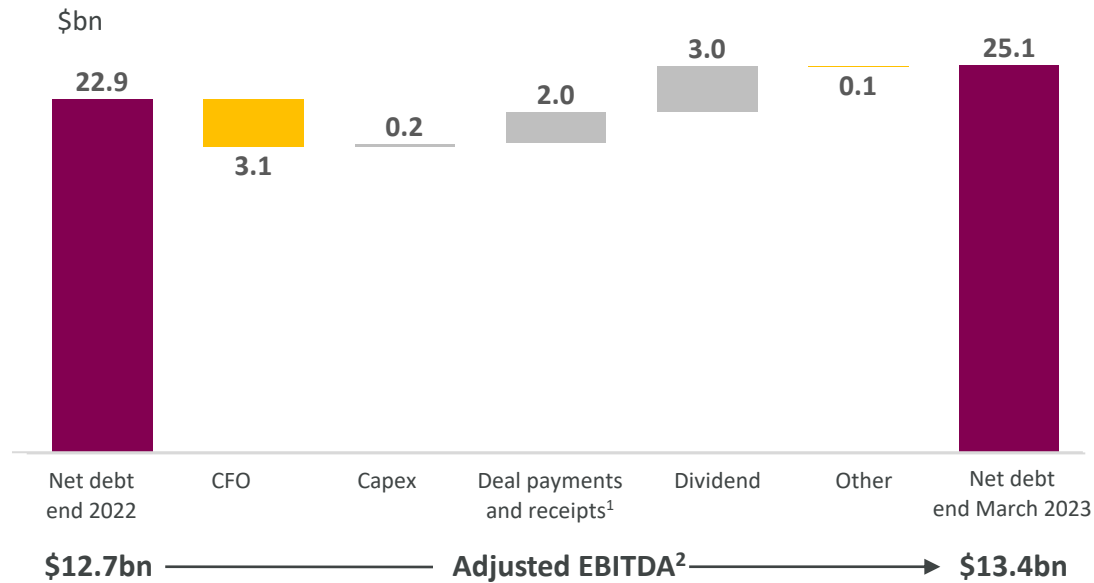
¹⁰ Absolute values at actual exchange rates; changes at CER. Gross margin excludes the impact of Alliance and Collaboration Revenue and any associated costs, thereby reflecting the underlying performance of Product Sales. 1. Total operating expenses include distribution, R&D and SG&A expenses. R&D = Research & Development; SG&A = Sales, General & Administrative; pp = percentage points; CER = constant exchange rates.



Cash Flow, Net Debt and 2023 Financial Guidance

Continued EBITDA improvement

Net Debt bridge



Net Debt/EBITDA: 2.3x

Net Debt/EBITDA adjusted for Alexion inventory fair value uplift: 1.9x

Reiterating 2023 Guidance

Total Revenue

- *Excluding COVID-19 medicines:* low double-digit % growth
- *Including COVID-19 medicines:* low-to-mid single-digit % growth

Core EPS

- High single-digit to low double-digit %

A low single-digit adverse FX-impact anticipated for both Total Revenue and Core EPS in 2023³



Artificial Intelligence at AstraZeneca

Leadership in AI is transforming the way we work and pace of innovation

AI is embedded across our organisation

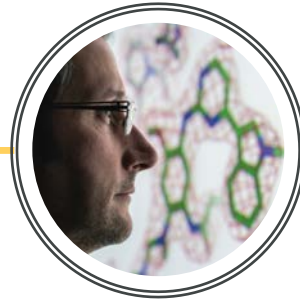
R&D • OPERATIONS • COMMERCIAL

400+

data scientists employed

100+

active AI projects within R&D alone



Drug discovery and development

- Clinical trials
- Supportive internal tools and platforms



Regulatory submission and monitoring

- Clinical forecasting and automation
- Pharmacovigilance



Manufacturing and supply

- Digital supply chain and manufacturing



Commercial launch and patient outcomes

- HCP experience
- Patient assistance
- Patient outcomes

Strategic investment in AI and digital tools has delivered demonstrable productivity gains and improvements in science-led innovation

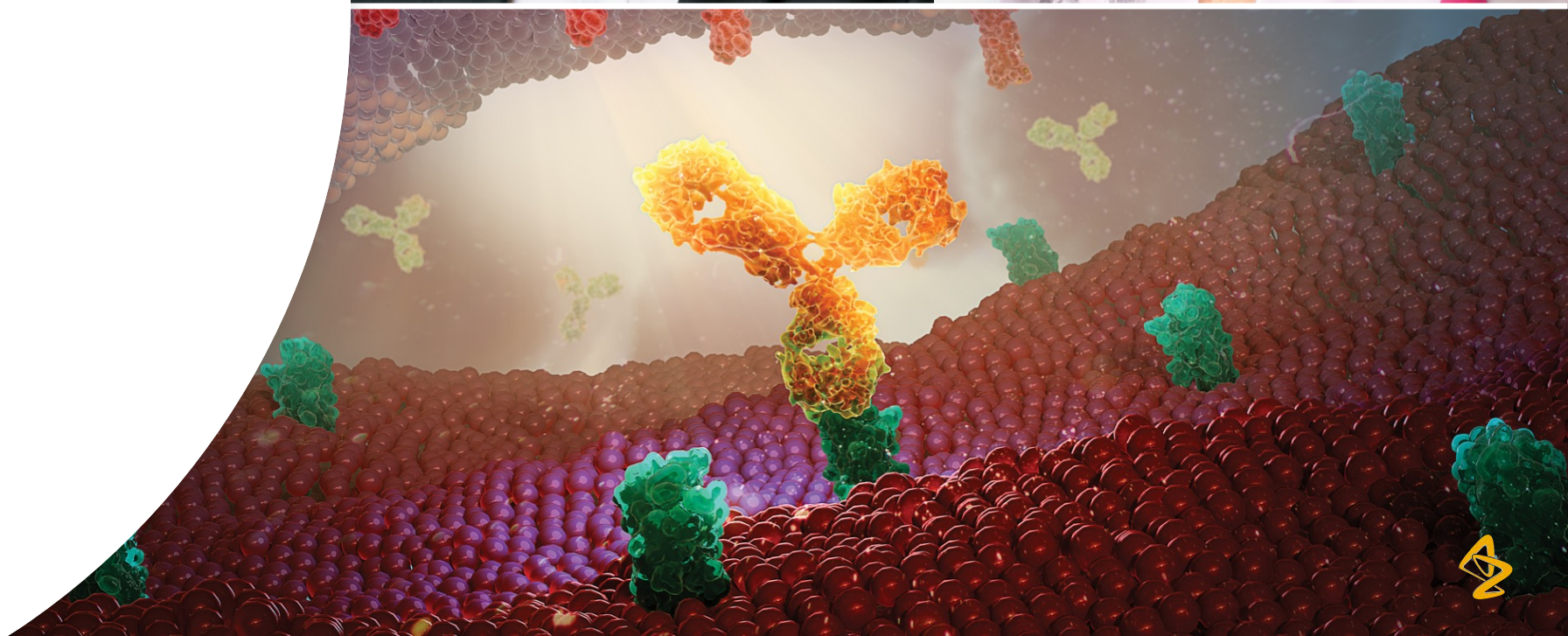


Oncology



Dave Fredrickson
ONCOLOGY BUSINESS

Susan Galbraith
ONCOLOGY R&D

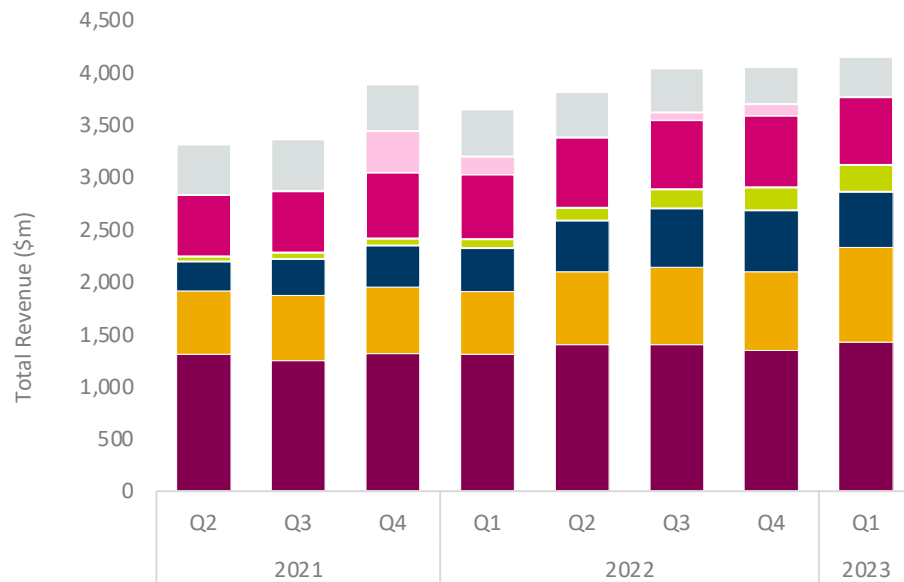


Oncology – Q1 2023

19% Total Revenue growth driven by differentiated portfolio and new indication launches

Oncology

Q1 2023 \$4.1bn, +19% at CER



Tagrisso Imfinzi/Imjudo Calquence Enhertu Lynparza (PS) Lynparza milestones Others

Q1 2023: key dynamics


- **Tagrisso** +15% CER, driven by global demand growth, CN recovery
 - **Lynparza PS** +10% CER, strong PROpel EU uptake offset by flattening testing rates and destocking in US
 - **Imfinzi/Imjudo** +56% CER, driven by global launch acceleration (TOPAZ-1, HIMALAYA, POSEIDON)
 - **Calquence** +31% CER, strong EU growth offset by destocking in US following maleate tablet approval (Q3 2022)
 - **Enhertu** >3x CER, strong global launch momentum
-
- New indications: EU (*Calquence* maleate tablet, *Imfinzi* HIMALAYA, POSEIDON), CN (*Calquence* for MCL, *Enhertu* DB03)



Oncology

Ambition for >50% of lung cancer patients to be eligible for AZN medicine by 2030

	resectable Stg. I-III	unresectable Stg. I-II	unresectable Stg. III	1L	metastatic 2L+	
Est. epi (G7)	~200K	~30K	~70K	~350K	~290K	
IO sensitive c.70%	<p><i>Imfinzi</i> AEGEAN</p> <p>volrustomig + CTx NEOCOAST-2</p>	<p><i>Imfinzi</i> w/ SBRT PACIFIC-4</p>	<p>CRT → <i>Imfinzi</i> PACIFIC</p> <p>CRT + <i>Imfinzi</i> PACIFIC-2</p> <p><i>Imfinzi</i> combos PACIFIC-8, -9 improvements across PD-L1 spectrum</p>	<p><i>Imfinzi</i> + <i>Imjudo</i> + CTx POSEIDON</p> <p>Dato-DXd + IO TROPION-Lung08 TROPION-Lung07 AVANZAR TROPION-Lung04/02</p> <p><i>Enhertu</i> + IO DESTINY-Lung03</p> <p>rilvegostomig (PD1/TIGIT) ARTEMIDE-1 volrustomig (PD1/CTLA4) sabestomig (PD1/TIM3)</p>	<p><i>Imfinzi</i> + ceralasertib LATIFY</p> <p>Dato-DXd TROPION-Lung01</p> <p>AZD9592 (EGFR/cMET ADC) EGRET</p>	
EGFRm c.16%	<p><i>Tagrisso</i> ADAURA</p> <p><i>Tagrisso</i> neo-ADAURA</p>		<p>CRT → <i>Tagrisso</i> LAURA</p>	<p><i>Tagrisso</i> FLAURA</p> <p><i>Tagrisso</i> + CTx FLAURA2</p>	<p>savolitinib + <i>Tagrisso</i> SAFFRON/SAVANNAH</p> <p>AZD9592 (EGFR/cMET ADC) EGRET</p>	
Other tumor drivers c.12%				<p>CRT → <i>Imfinzi</i> PACIFIC</p>		<p>Dato-DXd TROPION-Lung01 TROPION-Lung05</p>
HER2 c.2%					<p><i>Enhertu</i> DESTINY-Lung04</p>	<p><i>Enhertu</i> DESTINY-Lung02</p>

 established SoC

Leading the future of lung cancer treatment

- *Tagrisso* established TKI backbone in EGFRm
- *Imfinzi* leading IO in unresectable
- Advancing best-in-class ADCs to replace systemic chemotherapy
- Delivering next-wave bispecifics to improve on PD1/PD-L1
- Developing novel combinations, including IO + ADC
- Investing behind new technologies and platforms, including cell therapy, testing/screening

Est epi (G7) = estimated epidemiology across G7 (US, EU5, JP); Stg. = stage; CTx = chemotherapy; SBRT = stereotactic body radiation therapy; CRT = chemoradiotherapy; pembro = pembrolizumab; IO = immunotherapy; ADC = antibody-drug conjugate; PD1 = programmed cell death protein 1; EGFR = epidermal growth factor receptor; c-MET = mesenchymal-epithelial transition factor; TIGIT = T-cell immunoreceptor with immunoglobulin and ITIM domains; CTLA4 = cytotoxic T-lymphocyte associated protein 4;

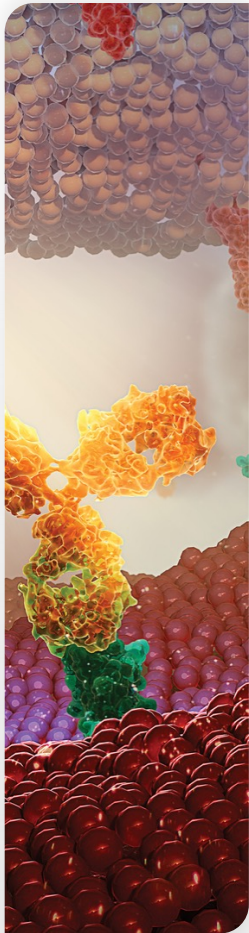
TIM3 = T-cell immunoglobulin and mucin domain-containing protein 3; SoC = standard of care; TKI = tyrosine kinase inhibitor.

Collaboration partners: Daiichi Sankyo (*Enhertu*, Dato-DXd), Compugen (rilvegostomig).



Oncology – R&D highlights

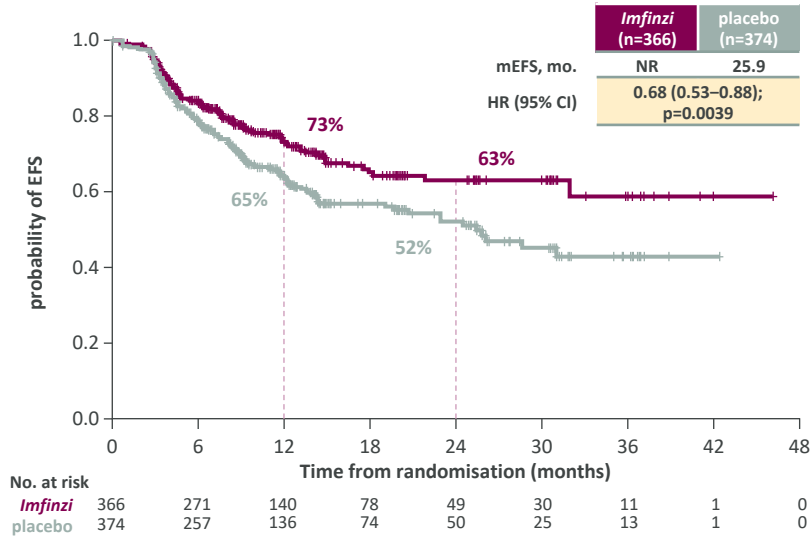
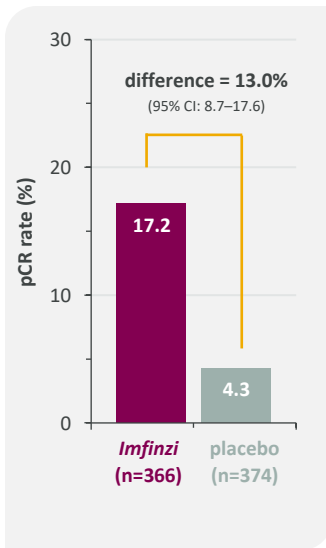
Moving into earlier lines of lung cancer with highest potential for cure



AEGEAN: updated pCR and interim EFS



Imfinzi-based treatment before and after surgery in early-stage NSCLC



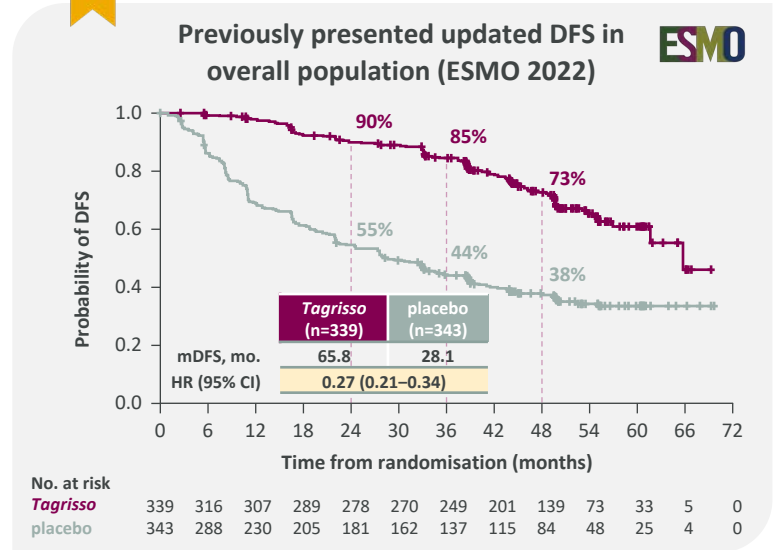
No. at risk		366	271	140	78	49	30	11	1	0
Imfinzi	placebo	374	257	136	74	50	25	13	1	0

Potential to become a backbone combination approach that may alter the course of a patient's cancer

ADAURA: final OS

adjuvant *Tagrisso* in early stage EGFRm NSCLC

ASCO plenary
First Phase III trial to demonstrate survival benefit in this setting

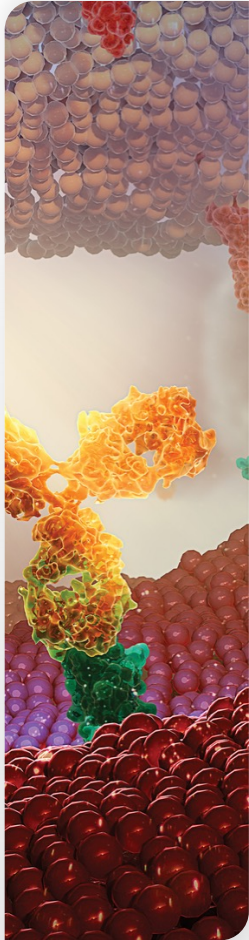


No. at risk		339	316	307	289	278	270	249	201	139	73	33	5	0
Tagrisso	placebo	343	288	230	205	181	162	137	115	84	48	25	4	0

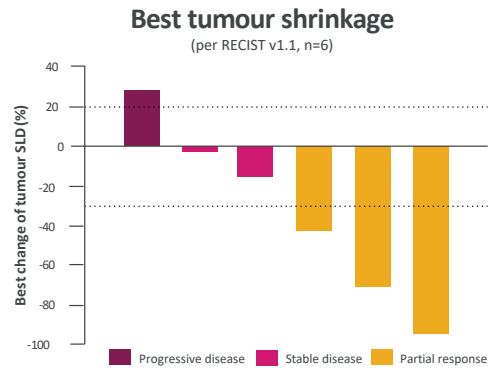


Oncology – R&D highlights

AACR demonstrates harnessing of in-house capabilities to build early-stage pipeline



First clinical data for CAR-T therapy



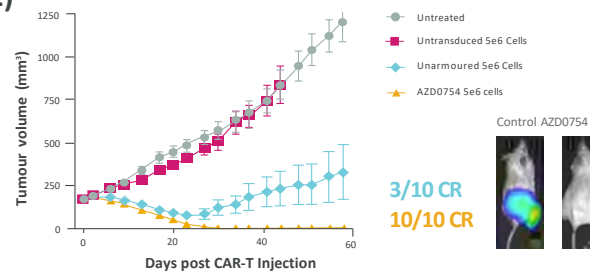
C-CAR031 (armoured GPC3)¹

- First-in-human trial in patients with hepatocellular carcinoma
- Well-tolerated with promising anti-tumour activity including objective responses in several patients

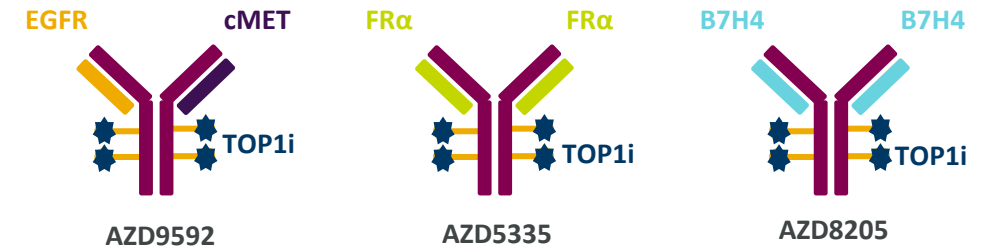
AZD0754 (armoured STEAP2)

- Favourable *in vitro* properties
- Robust dose dependent *in vivo* efficacy in STEAP2-expressing CDX and PDX models
- Encouraging preclinical safety

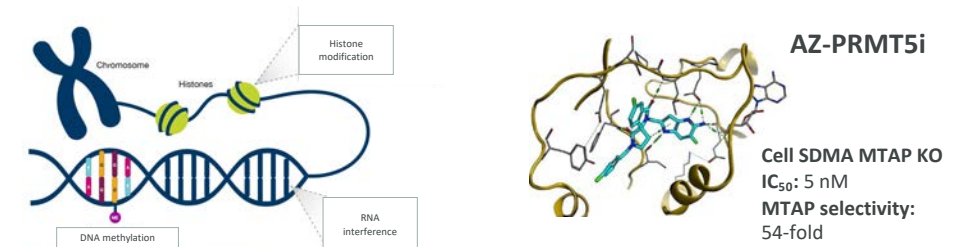
Tumour volume in mice after CAR-T injection



Preclinical data for three AstraZeneca in-house developed ADCs



First disclosure and preclinical data for epigenetics molecule targeting PRMT5, AZ-PRMT5i



1. Designed by AstraZeneca and manufactured and developed in China by Cellular Biomedicine group.

AACR = American Association for Cancer Research annual meeting; RECIST = Response Evaluation Criteria In Solid Tumours; CAR-T = chimeric antigen receptor T-cell; SLD = sum of longest diameters; GPC3 = Glypican 3; CDX = cell line-derived xenograft; PDX = patient-derived xenograft; ADC = antibody-drug conjugate; EGFR = epidermal growth factor receptor; c-MET = mesenchymal-epithelial transition factor; TOP1i = topoisomerase 1 inhibitor; FR α = folate receptor alpha; MTAP = methylthioadenosine phosphorylase; SDMA = symmetric dimethylarginine; KO = knockout; IC₅₀ = half-maximal inhibitory concentration; nM = nanomolar.



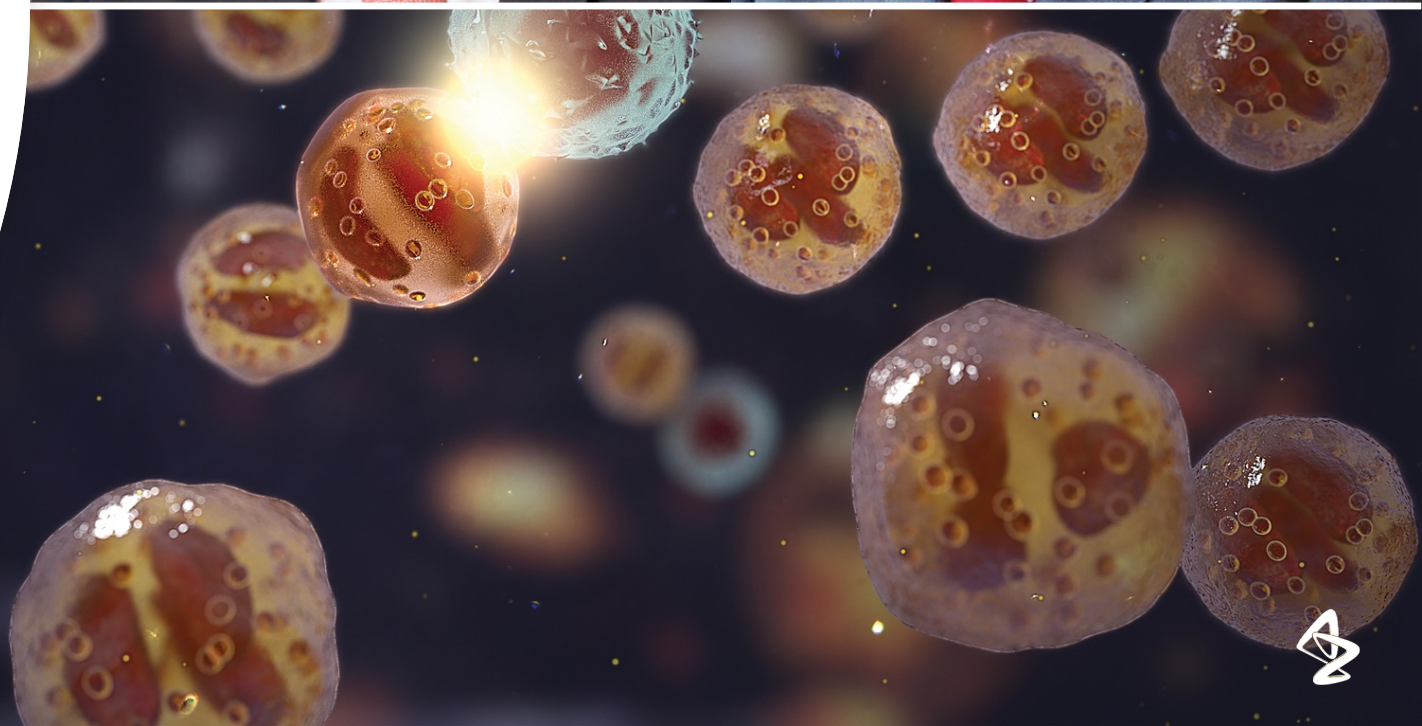
BioPharmaceuticals

Ruud Dobber

BIOPHARMACEUTICALS BUSINESS

Mene Pangalos

BIOPHARMACEUTICALS R&D

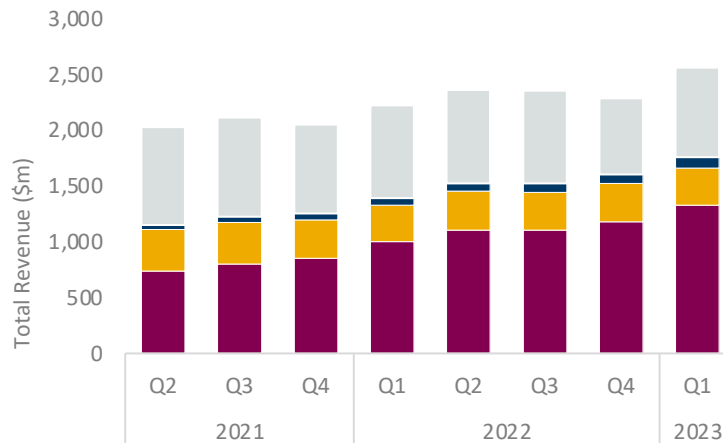


BioPharmaceuticals – Q1 2023

Increasing momentum across CVRM and R&I

CVRM

Q1 2023 \$2.6bn, +22% at CER

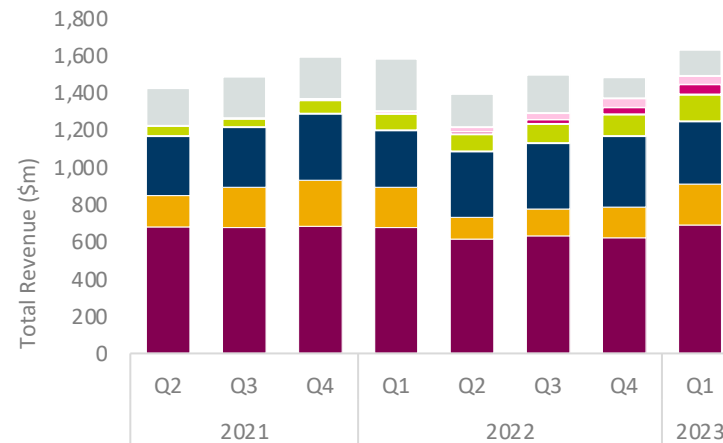


Farxiga Brilinta Lokelma Others

- **Farxiga** +39% to \$1.3bn
- Strong double-digit growth from **Lokelma** and roxadustat

R&I

Q1 2023 \$1.6bn, +8% at CER

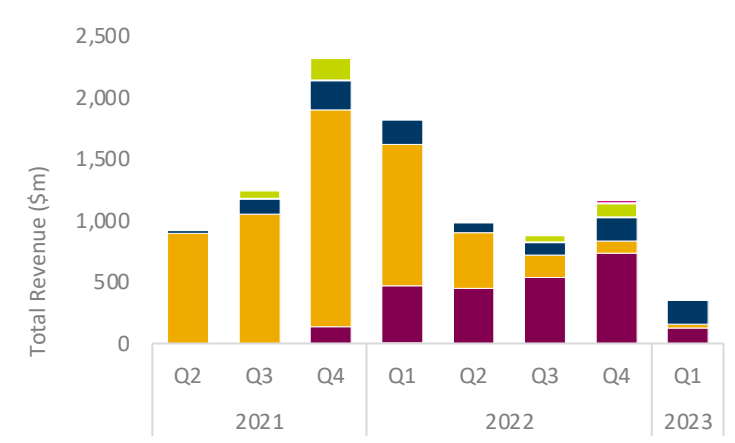


Symbicort Pulmicort Fasenra Breztri Tezspire Saphnelo Others

- **Fasenra** +13% to \$338m
- **Breztri** +73% to \$144m
- **Tezspire** +32% QoQ growth to \$54m

V&I

Q1 2023 \$355m, -79% at CER



COVID-19 mAbs¹ Vaxzevria Synagis Flumist Beyfortus

- COVID-19 medicines declined by \$1.5bn
- **Synagis** +5%

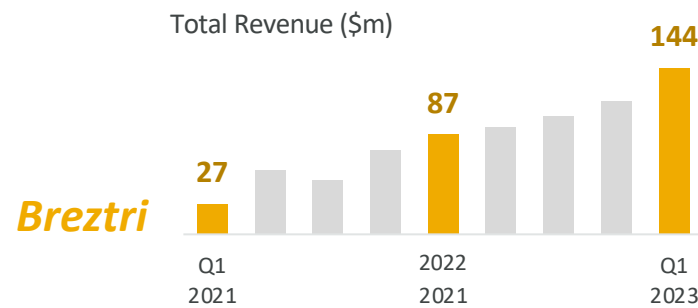
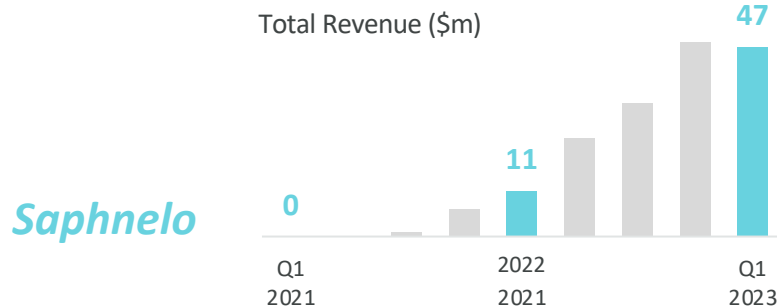
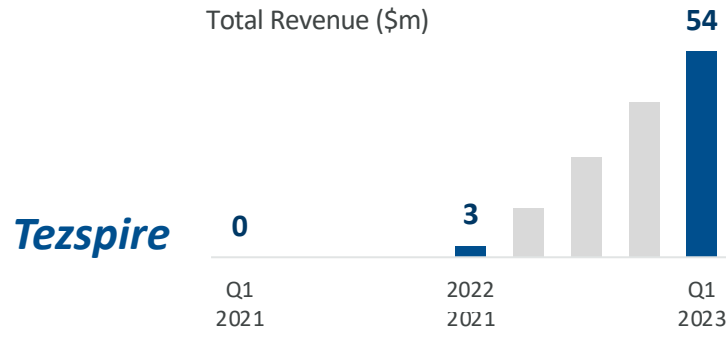
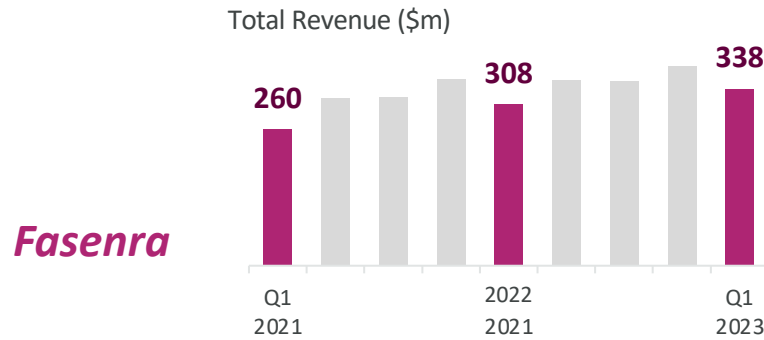
1. COVID-19 mAbs = Evusheld and AZD3152, the antibody currently in development



BioPharmaceuticals – Q1 2023

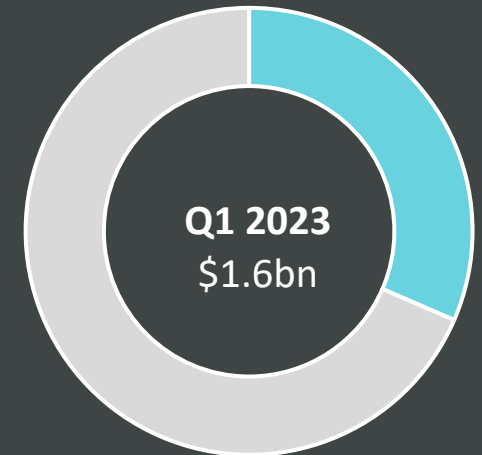
Key medicines driving R&I growth

Building launch momentum in R&I



+46% growth in Q1 from key R&I medicines

Fasenra • Tezspire • Saphnelo • Breztri

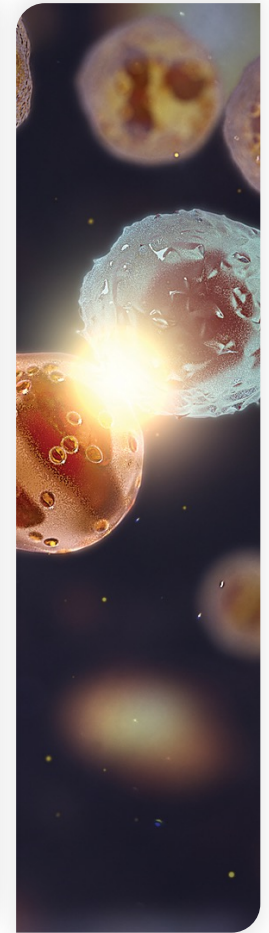
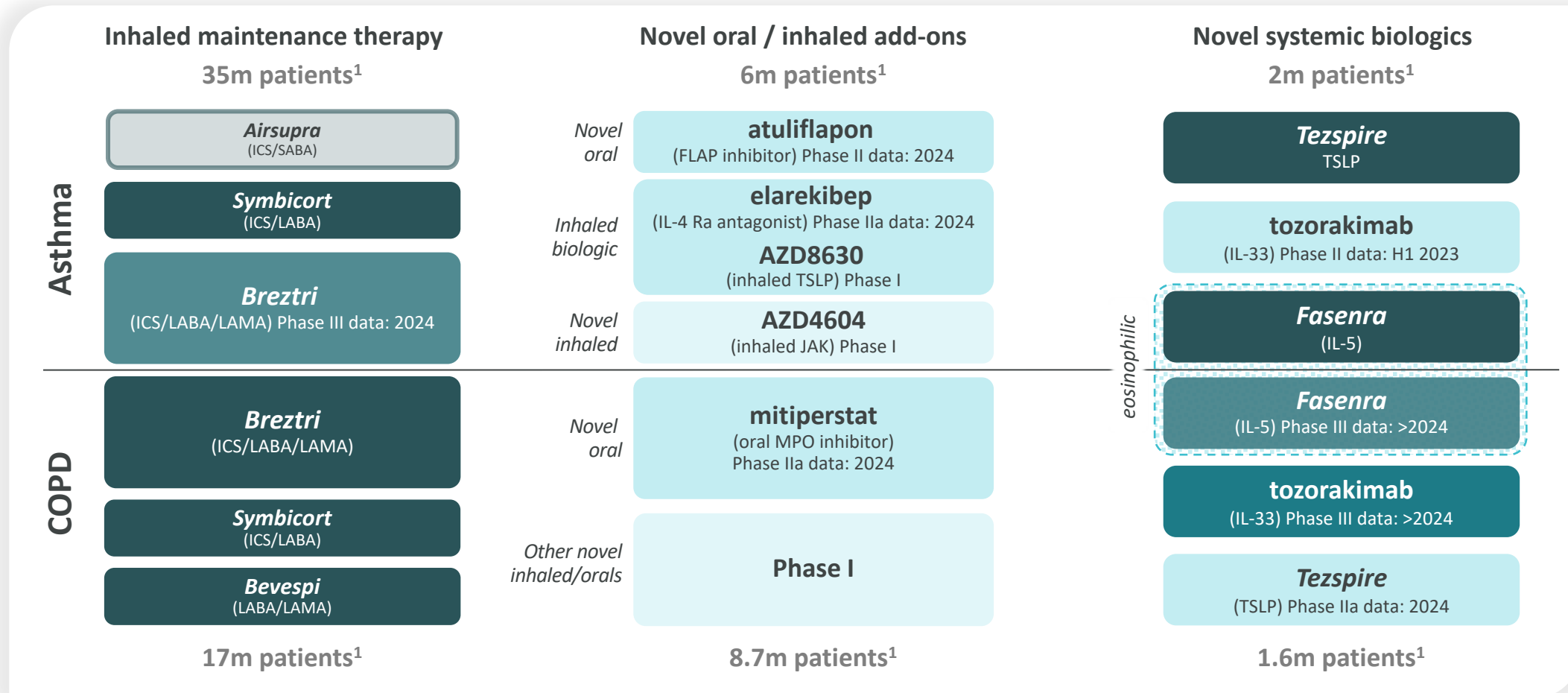


R&I Q1 2023 Total Revenue



BioPharmaceuticals – R&D highlights

Leading respiratory portfolio across asthma and COPD



launched
 approved/launch prep
 Phase III
 Phase II
 Phase I

1. Populations relate to T7 (US, EU5 and CN) and reflect AstraZeneca projections from IQVIA prescription and units sold data

ICS = inhaled corticosteroid; SABA = short-acting beta-agonist; LABA = long-acting beta-agonist; LAMA = long-acting muscarinic antagonist; MPO = myeloperoxidase; TSLP = thymic stromal lymphopoietin; IL-33 = interleukin-33; IL-5 = interleukin-5; JAK = janus kinase;

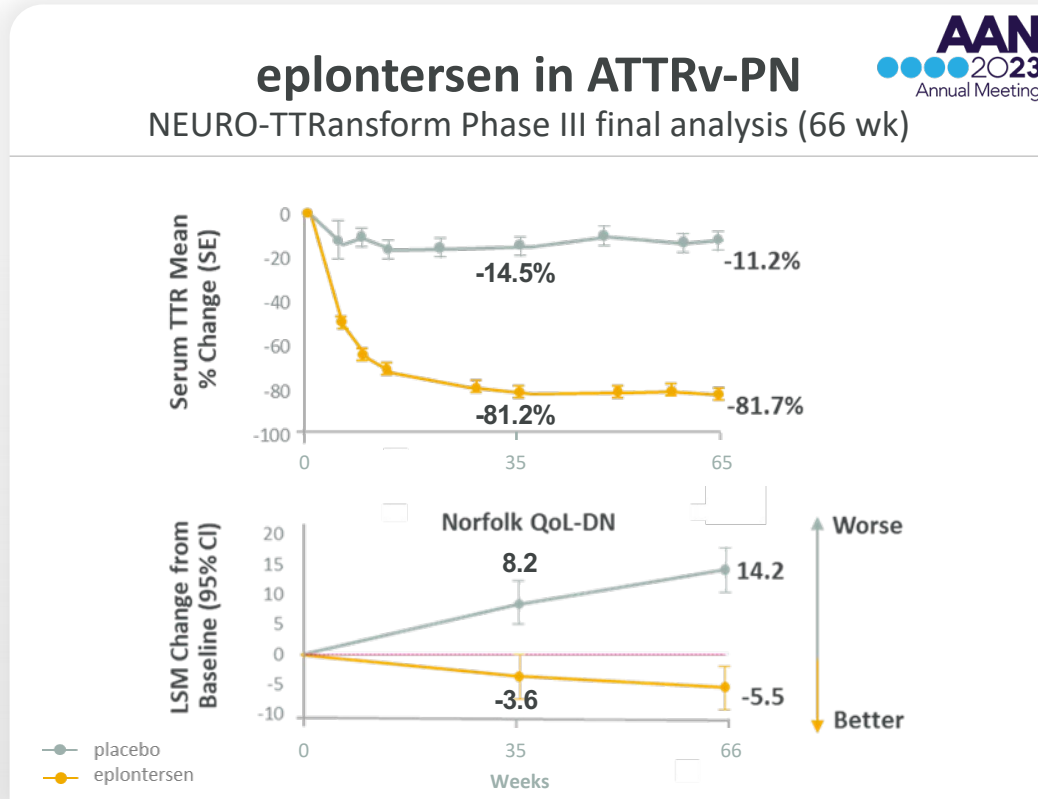
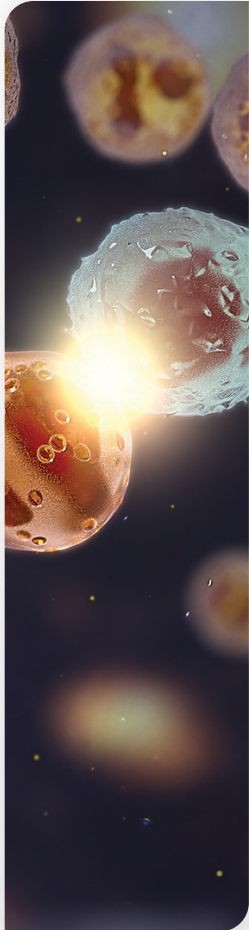
IL-4 Ra = interleukin-4 receptor alpha; FLAP = 5-lipoxygenase-activating protein.

Collaboration partners: Amgen (*Tezspire*).

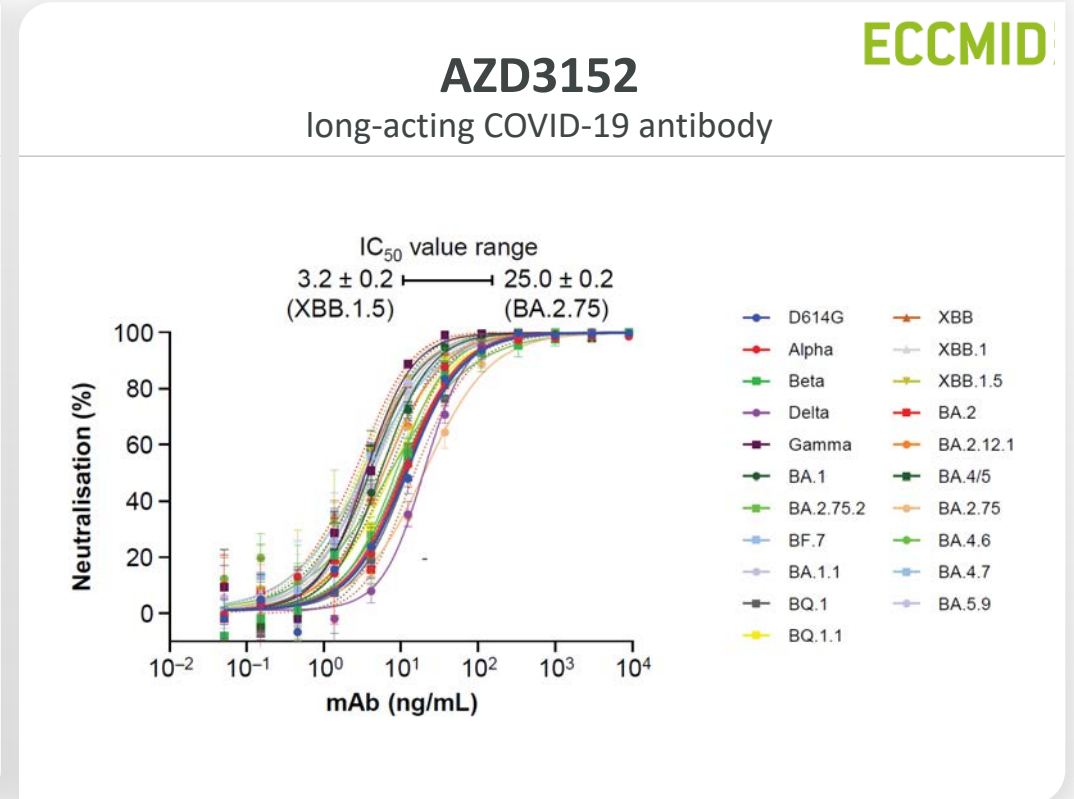


BioPharmaceuticals – R&D highlights

Congress highlights at AAN and ECCMID



Met co-primary endpoints, with favourable safety and tolerability profile



***In vitro* studies show neutralisation across all known variants, including Arcturus**



AI in R&D

Accelerating the time to deliver clinical leads



DRUG DISCOVERY & DEVELOPMENT | *reinventing the traditional drug discovery process*

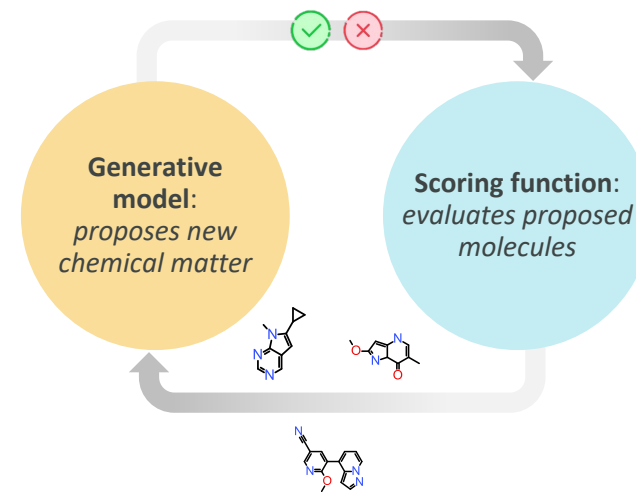
Knowledge graphs | *empowering data insights*



Visual representation of relationships built from data, external literature, etc.

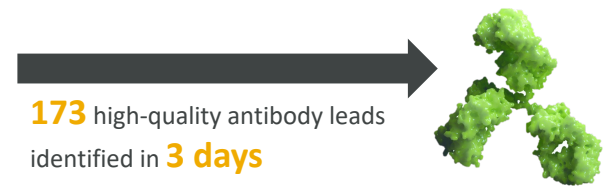
- Which genes are upregulated in Disease A?
- What genes are linked to Disease A?
- Which targets are druggable?
- What are the pathway relationships between these targets?

Accelerating discovery speed | *small and large molecules*



AI-enabled process accelerates generation of high-quality small molecules by >2x

Traditional process generated
0 leads over **3 months**



173 high-quality antibody leads
identified in **3 days**

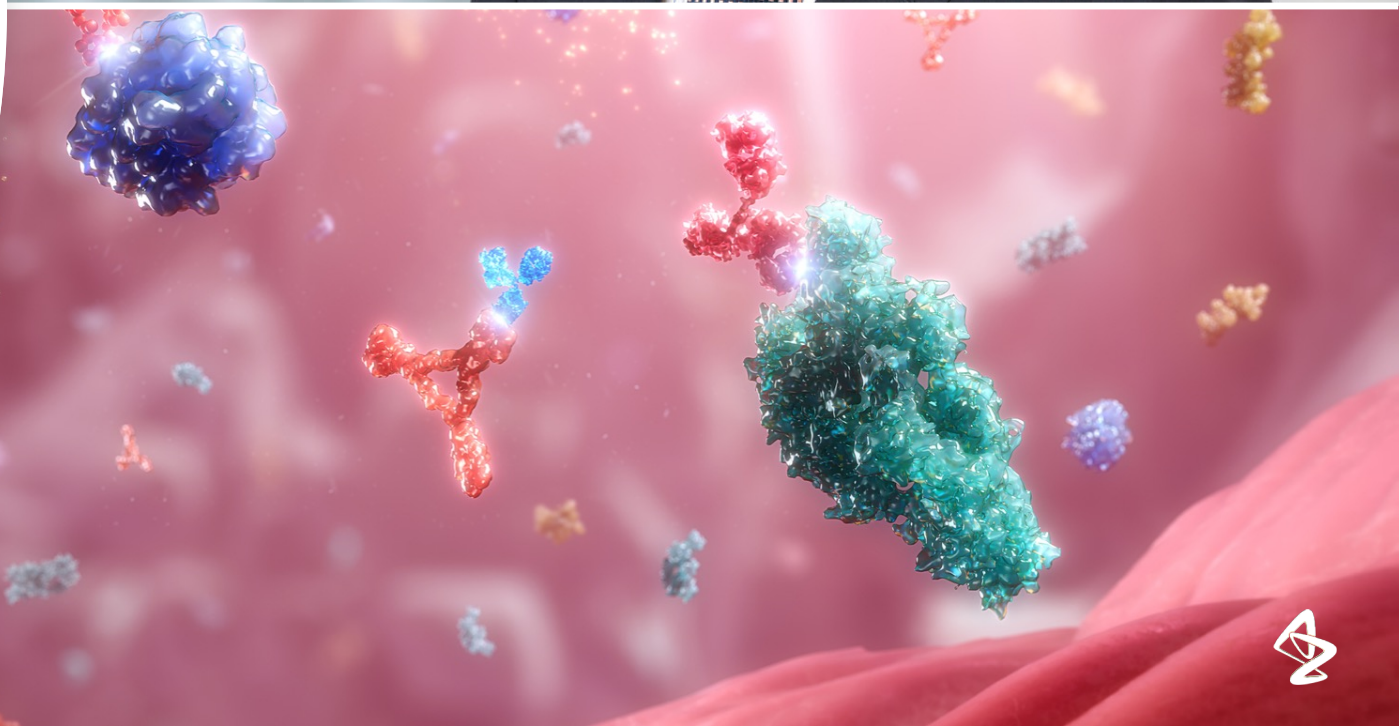
AI-enabled process cut time to identify target antibody leads to 3 days



Rare Disease

Marc Dunoyer

CHIEF EXECUTIVE OFFICER,
ALEXION

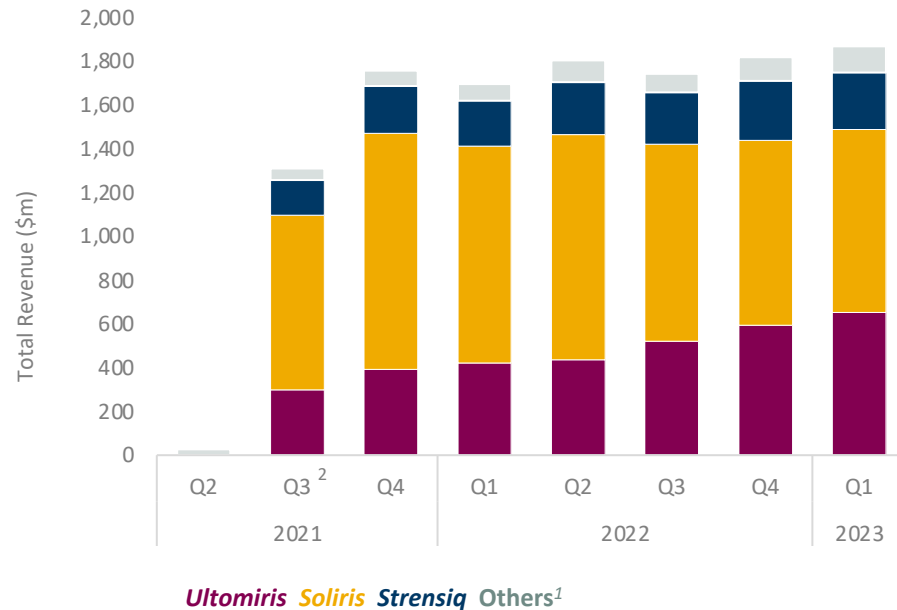


Rare Disease – Q1 2023

Accelerated conversion in C5, continued strength beyond complement

Rare Disease

Q1 2023 \$1.9bn, +14% at CER



Q1 2023: key dynamics

Durable C5 Franchise growth

- **Ultomiris** +61% driven by successful conversion from **Soliris**, new patients and market expansion
- **Soliris** (13%) decline reflecting successful conversion, partially offset by NMOsD growth

Strensiq +28% and Koselugo >2x

- Reflecting strength of patient demand and geographic expansion

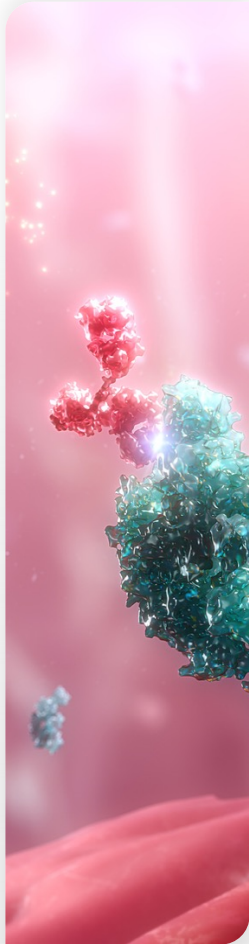
Strong commercial execution across indications and geographies

1. Includes *Kanuma* and *Koselugo*. 2. Q3 2021 Total Revenues reported only comprise of those booked by AstraZeneca following completion of the acquisition of Alexion on 21 July 2021. C5 = C5 inhibitors *Ultomiris* and *Soliris*;

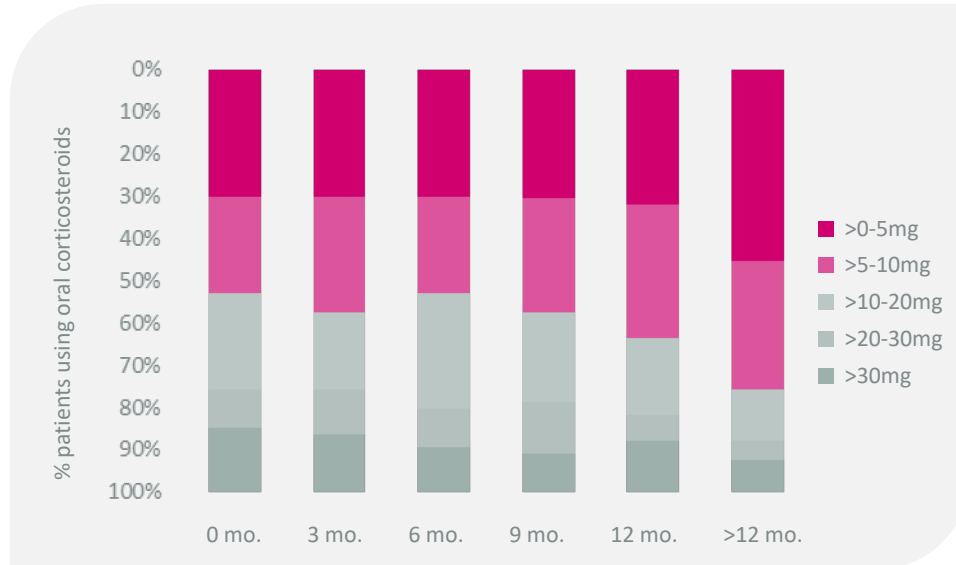


Rare Disease – R&D highlights

Pioneering in rare neurology to improve patient outcomes



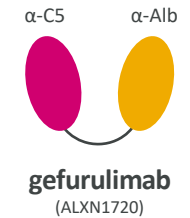
Soliris gMG registry¹ oral corticosteroid usage decreased over time



>75% patients on low-dose OCS after 1 year

gefurulimab (ALXN1720) pipeline expansion to further extend C5 leadership

C5 bi-specific, heavy chain (V_HH)
antibody demonstrated
**complete terminal
complement inhibition²**
in Phase I



Phase III PREVAIL in gMG:

R
1:1
n=254

gefurulimab QW s.c.

placebo

Primary endpoint:
change from baseline in
MG-ADL total scores
at Week 26

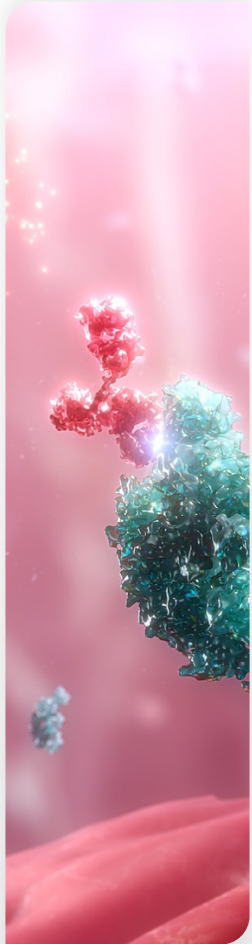
Potential best-in-class weekly, self-admin s.c.

1. Pulley et al. American Academy of Neurology Annual Meeting 2023 based on *Soliris* data; concomitant therapies include: use of azathioprine (AZA), mycophenolate mofetil (MMF), intravenous immunoglobulin (IVIg)/plasma exchange (PLEX) and oral corticosteroids at initiation of, and during *Soliris*. 2. Ortiz et al. American Academy of Neurology Annual Meeting 2023. Complete terminal complement inhibition is defined as serum-free C5 concentrations less than 0.5 micrograms per millilitre; gMG = generalised myasthenia gravis; mo. = month; mg = milligram; OCS = oral corticosteroids; V_HH = single domain; Alb = albumin; r = randomised; QW = once weekly; s.c. = subcutaneous; MG-ADL = Myasthenia Gravis-Activities of Daily Living.



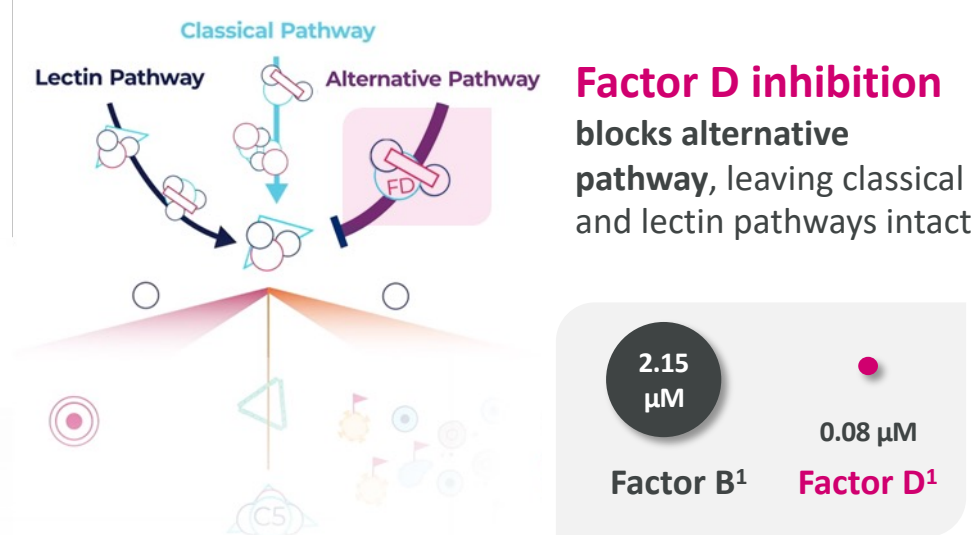
Rare Disease – R&D

Industry-leading Factor D portfolio



Factor D inhibition

more tractable target, lower circulating concentration in plasma



Factor D more likely to maintain consistent control than Factor B inhibitors

Factor D portfolio

novel small molecule assets with high affinity for Factor D

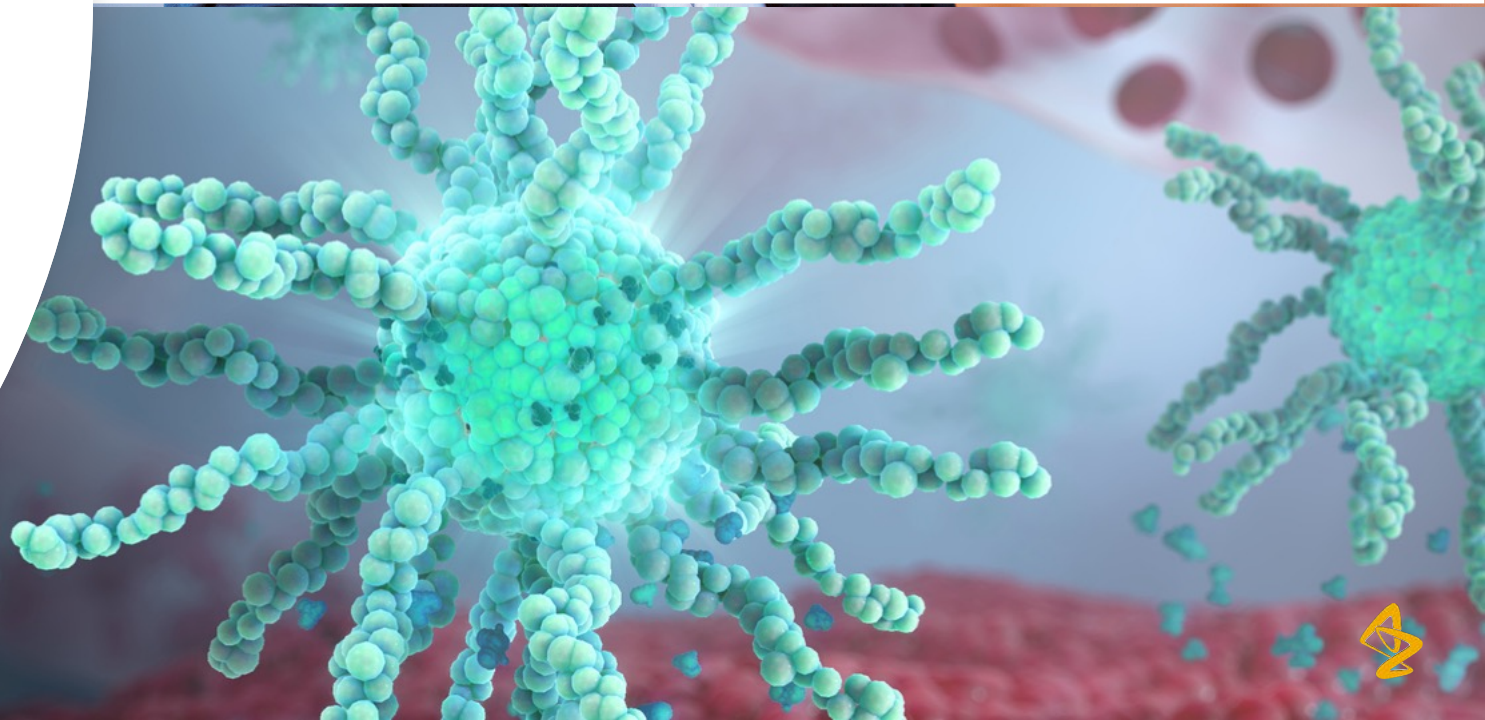
	dosing	development status
danicopan (ALXN2040)	TID	PNH-EVH submitted, Phase II GA ongoing as potential first oral treatment
vemircopan (ALXN2050)	BID	Phase II ongoing PNH monotherapy, gMG, Renal (LN, IgAN)
ALXN2080	QD	entered Phase I in Q3 2022 with potential application in non-rare indications

Robust Factor D portfolio with broad application



CEO Closing Remarks

Pascal Soriot
CHIEF EXECUTIVE OFFICER



AstraZeneca – Q1 2023

Accelerating pipeline momentum, disciplined investment fuels industry-leading growth

Pipeline advances in 2023

with 18 Phase III read-outs anticipated, including:

H1 2023

Dato-DXd – TROPION-Lung01 – 2nd-line/3rd-line NSCLC

Tagrisso – FLAURA2 – 1st-line NSCLC

H2 2023

Enhertu – DESTINY-Breast06 – 2L+ HR-positive/HER2-low BC

Tagrisso – LAURA – Stage III unresectable EGFRm NSCLC

Fasenra – MANDARA – EGPA

Dato-DXd – TROPION-Breast01 – HR-positive/HER2-negative mBC



Total Revenue ambition¹:
low double-digit % 2021-2025
Industry-leading growth 2025+



Remain focused on operating
margin expansion



At least 15 NMEs
approved by 2030



Emissions reduction:
98% by end 2025 – Scope 1 & 2
50% by 2030 – Scope 3

1. Indicates Company ambition to achieve Total Revenue low double-digit CAGR through 2025 (2021 base year, Alexion pro-forma) and industry-leading Total Revenue beyond 2025; this is not formal guidance. Dato-DXd = datopotamab deruxtecan; NSCLC = non-small cell lung cancer; HER2 = human epidermal growth factor receptor 2; HR = hormone receptor; BC = breast cancer; EGFRm = epidermal growth factor receptor mutant; EGPA = eosinophilic granulomatosis with polyangiitis; ESG = environmental, sustainability and governance; CAGR = compound annual growth rate; NMEs = new molecular entities. Collaboration partners: Daiichi Sankyo (Enhertu, Dato-DXd).



Question & Answer Session



Pascal Soriot
EXECUTIVE DIRECTOR &
CHIEF EXECUTIVE OFFICER



Aradhana Sarin
EXECUTIVE DIRECTOR &
CHIEF FINANCIAL OFFICER



Marc Dunoyer
CHIEF EXECUTIVE OFFICER,
ALEXION



Susan Galbraith
EXECUTIVE VICE PRESIDENT,
ONCOLOGY R&D



Dave Fredrickson
EXECUTIVE VICE PRESIDENT,
ONCOLOGY BUSINESS



Mene Pangalos
EXECUTIVE VICE PRESIDENT,
BIOPHARMACEUTICALS R&D



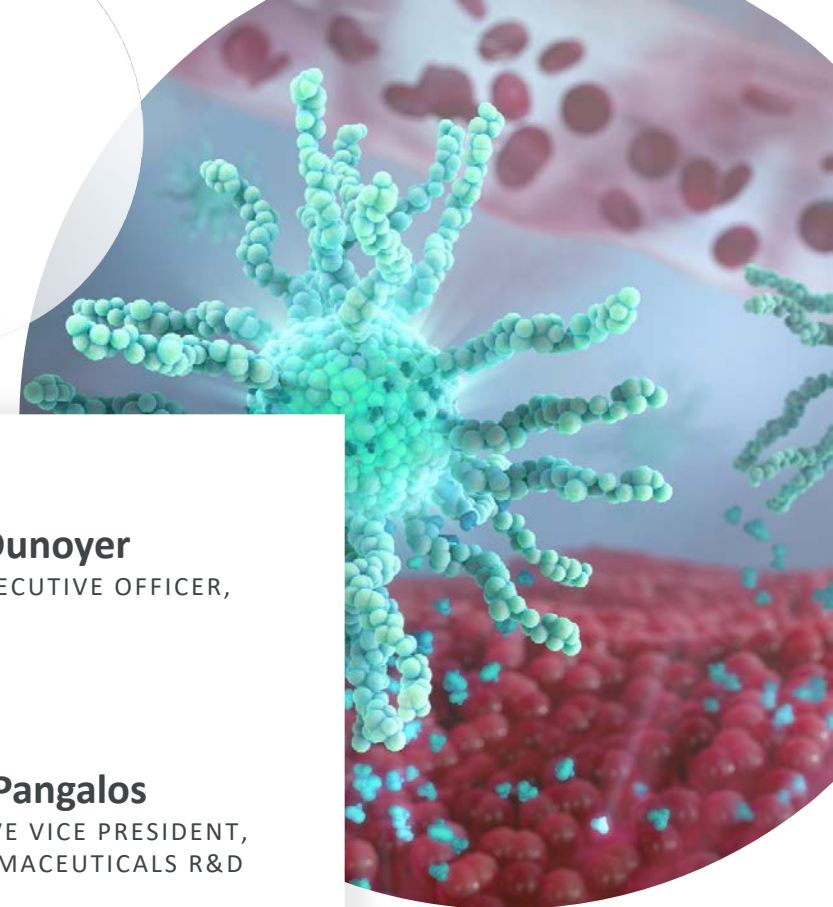
Ruud Dobber
EXECUTIVE VICE PRESIDENT,
BIOPHARMACEUTICALS
BUSINESS



Iskra Reic
EXECUTIVE VICE PRESIDENT,
VACCINES AND IMMUNE
THERAPIES



Leon Wang
EXECUTIVE VICE PRESIDENT,
INTERNATIONAL



Appendix

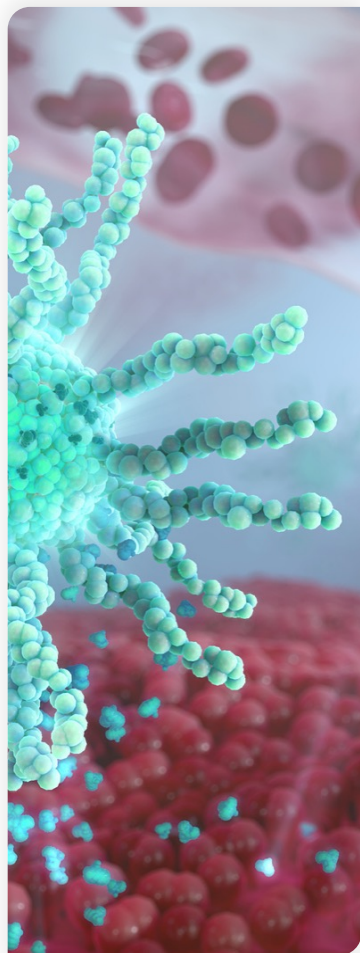
- Pipeline Highlights since Q4/FY2022 results
- ESG & Corporate Sustainability
- Oncology landscapes: breast and lung
- Emerging Markets
- Key Performance by Therapy Area



Delivering on science-led innovation

Selected key pipeline highlights since Q4/FY 2022 results

Oncology BioPharmaceuticals Rare Disease



5 regulatory approvals in major markets, including:

Imfinzi +/- Imjudo (EU)

non-small cell lung cancer (1st-line) (POSEIDON)

Imfinzi + Imjudo (EU)

hepatocellular carcinoma (1st-line) (HIMALAYA)

Enhertu (CN)

HER2+ breast cancer (2nd-line) (DESTINY-Breast03)

Calquence (EU)

maleate tablet formulation

Calquence (CN)

mantle cell lymphoma

1 positive CHMP opinion:

Ultomiris (EU)

neuromyelitis optica spectrum disorder
(CHAMPION-NMO)



2022 Sustainability highlights

Progress on our overall strategy includes:

14

public and private sector organisations convened by AstraZeneca CEO through the SMI to accelerate transition to net-zero health systems

87%

of employee survey respondents say that they understand their contributions to our sustainability priorities

25/27

of sustainability targets in Sustainability Data Summary are 'on plan'

Access to Healthcare

126,684

healthcare workers and others trained¹ (cumulative)

By 2025: 170,000

>44.63m

people reached through Access to Healthcare programmes (cumulative)

By 2025: 50M

>12.83m

people reached through our patient assistance programmes (cumulative)

Environmental protection

59.3%

reduction in Scope 1 and Scope 2 greenhouse gas emissions

By 2026: 98% from 2015 base year

18.7%

reduction in our water use

By 2025: 20% below 2015 baseline

18.6%

reduction in our waste

By 2025: 10% below 2015 baseline

Ethics and transparency

49.5%

senior middle management roles held by women

By 2025: reach gender equality in management positions

8 countries

with supplier diversity programmes

By 2025: 10 new countries outside of the US

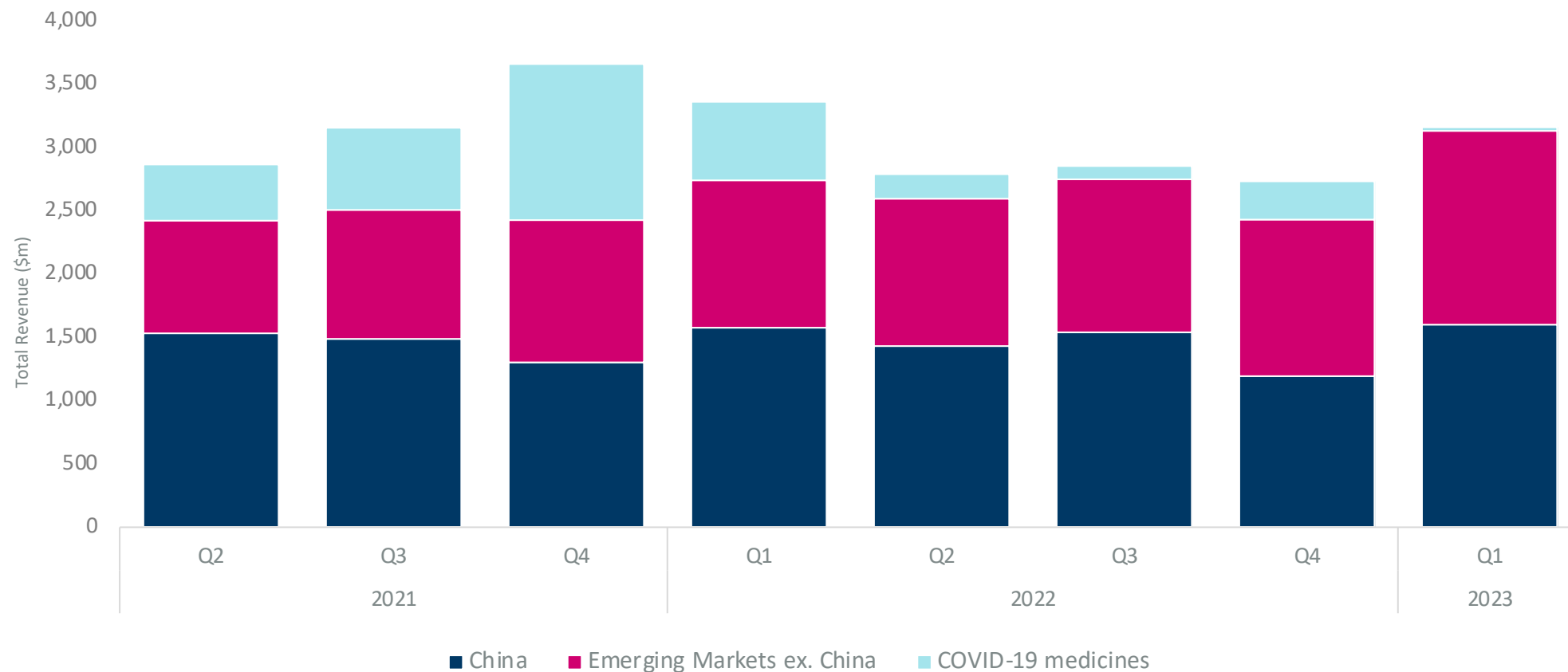
83%

of employee survey respondents feel we have a 'speak up' culture



Emerging Markets – Q1 2023

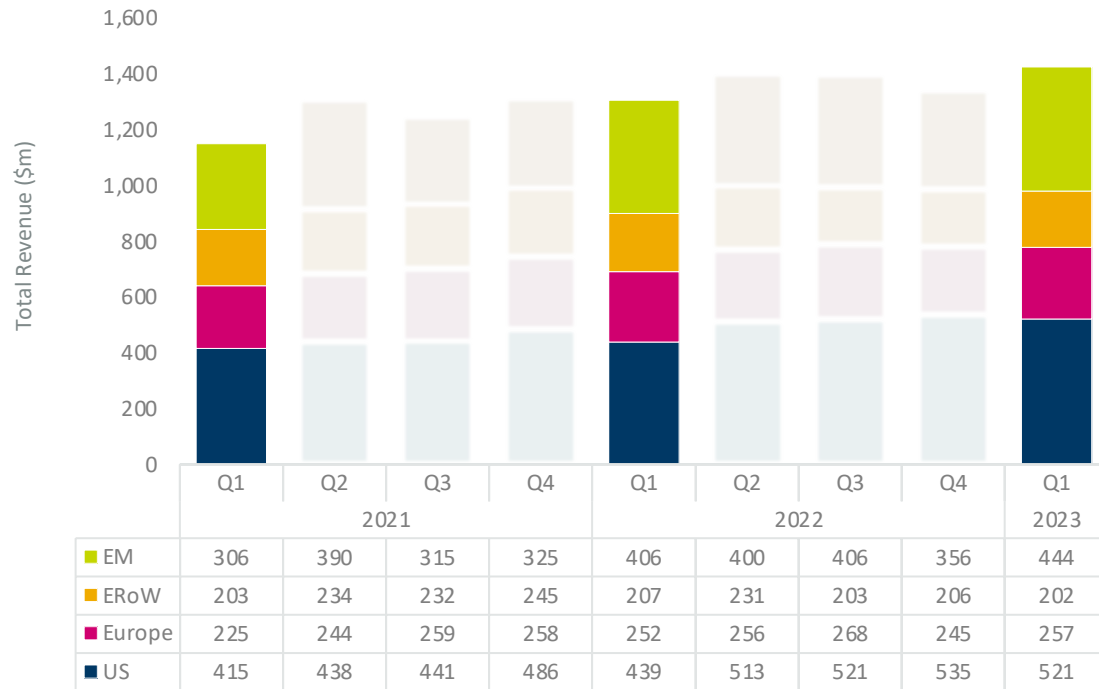
Total Revenue +1% at CER to \$3.2bn. +22% at CER excluding COVID-19 medicines



Oncology

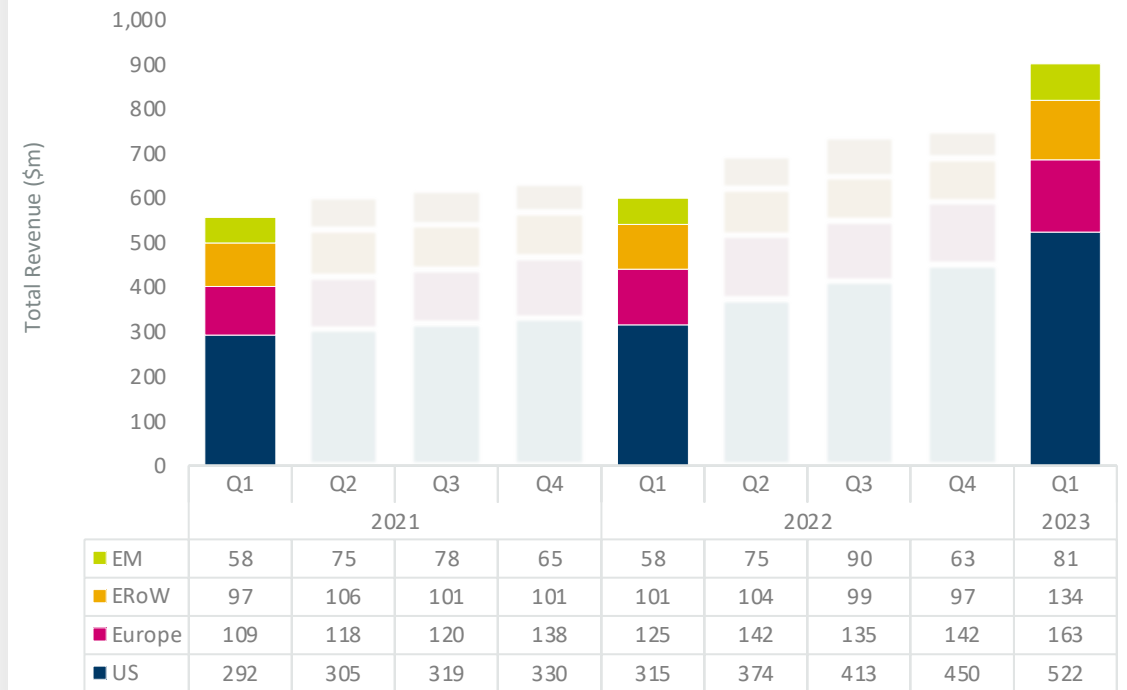
Tagrisso

15% growth at CER to \$1,424m



Imfinzi

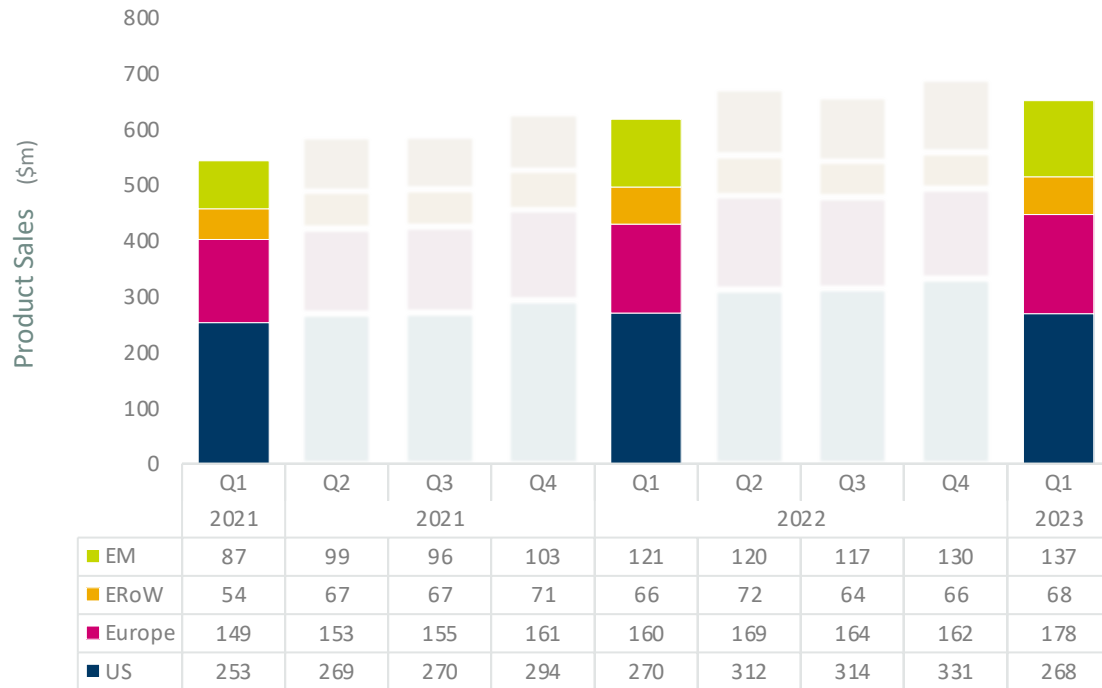
56% growth at CER to \$900m



Oncology

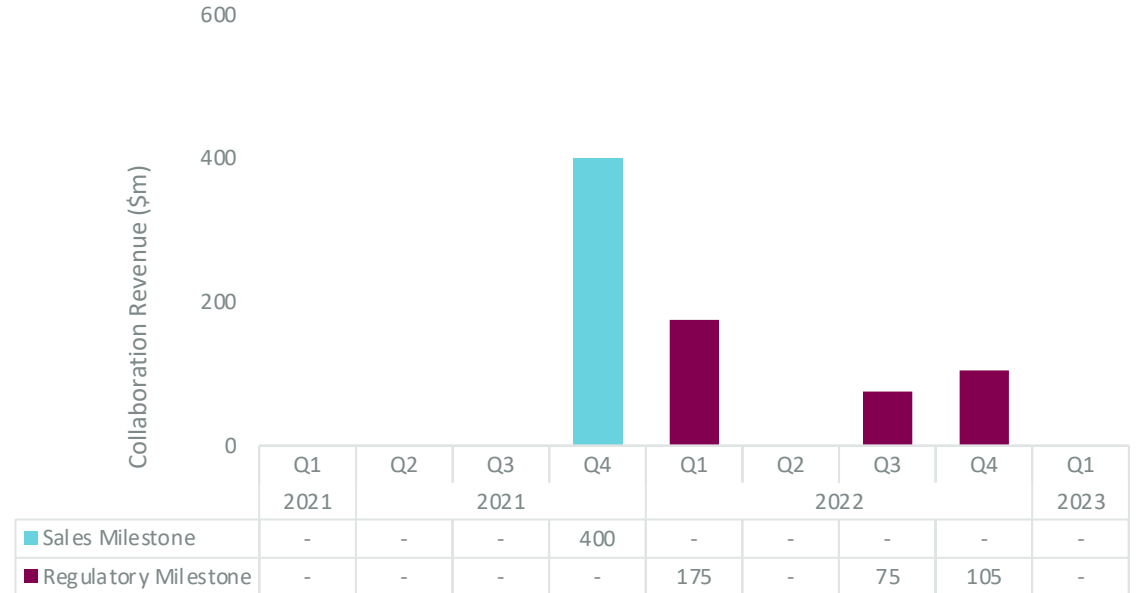
Lynparza

14% decrease at CER to \$651m



Lynparza

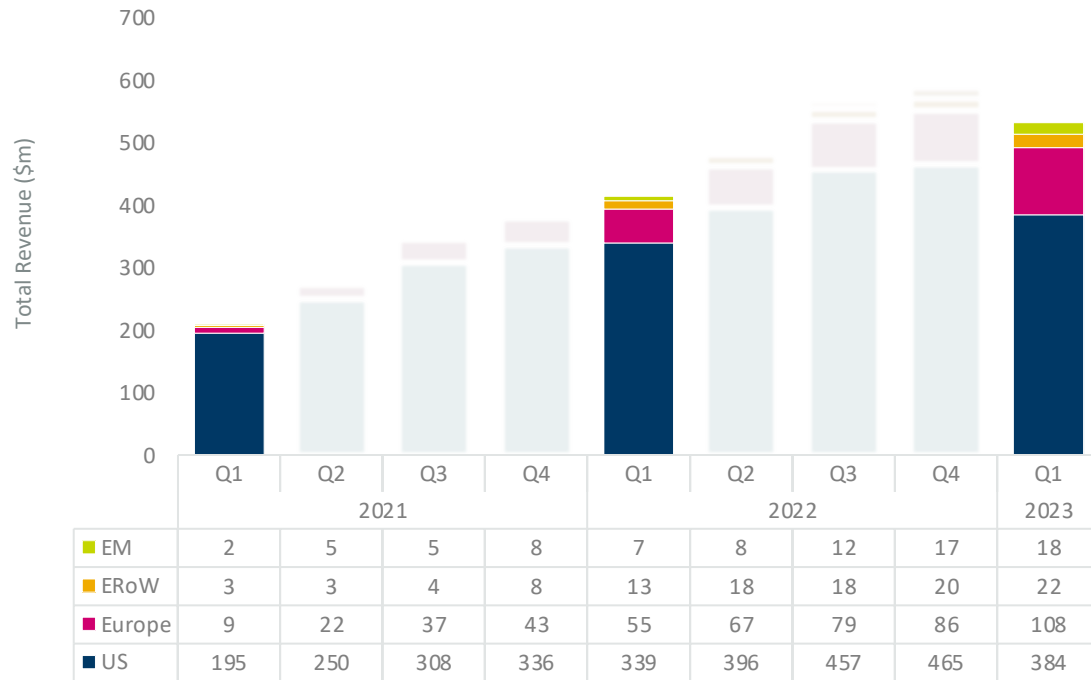
Collaboration Revenue: \$3.8bn recorded cumulative, \$3.9bn future potential



Oncology

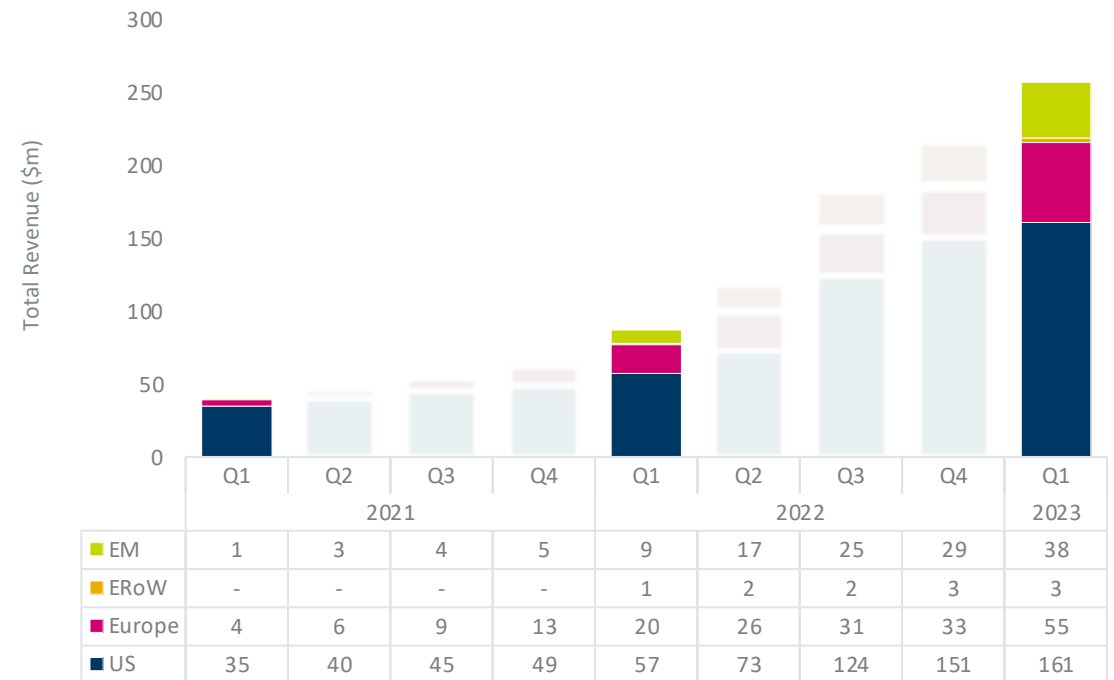
Calquence

31% growth at CER to \$532m



Enhertu

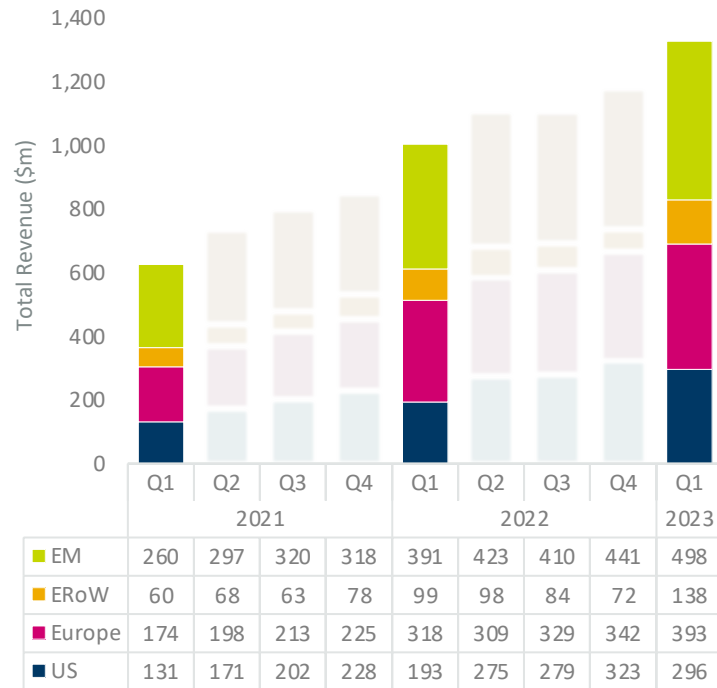
>3x growth at CER to \$257m



BioPharmaceuticals: Cardiovascular, Renal & Metabolism

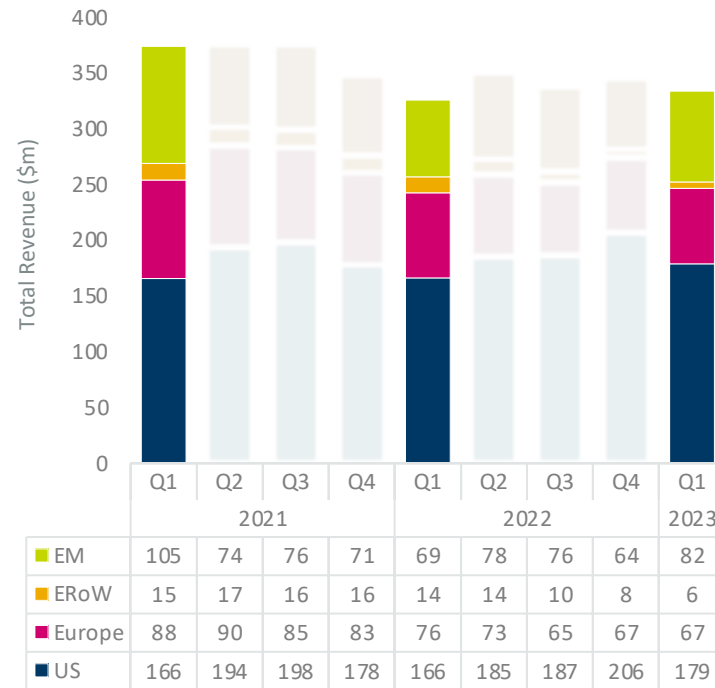
Farxiga

39% growth at CER to \$1,324m



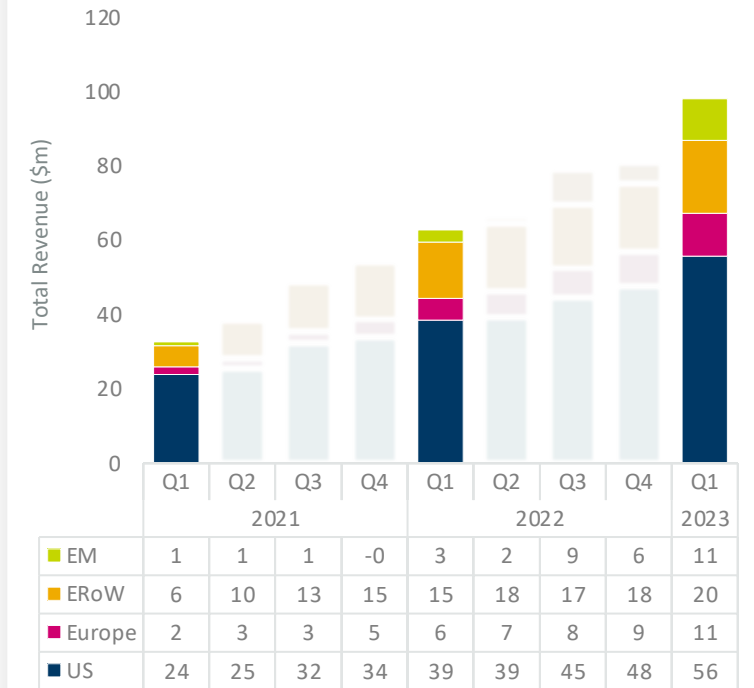
Brilinta

5% growth at CER to \$334m



Lokelma

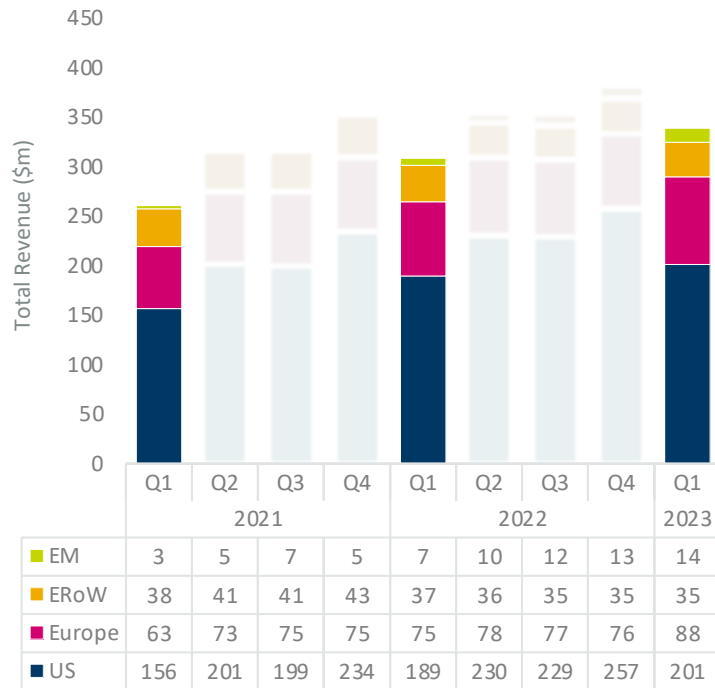
64% growth at CER to \$98m



BioPharmaceuticals: Respiratory & Immunology

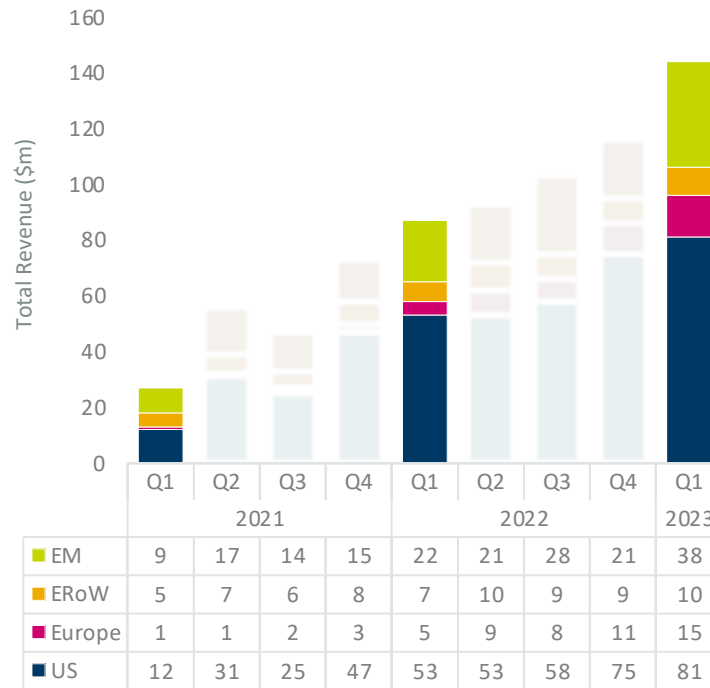
Fasenra

13% growth at CER to \$338m



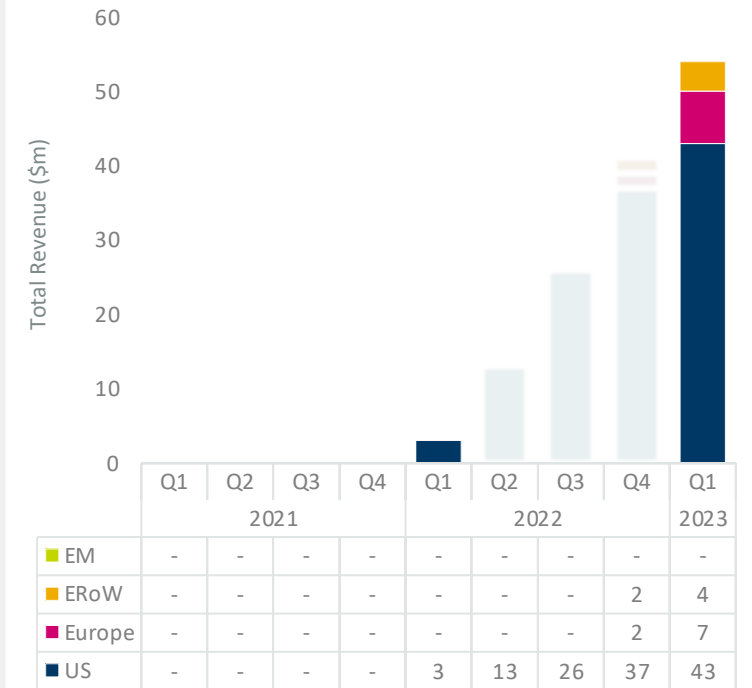
Breztri

73% growth at CER to \$144m



Tezspire

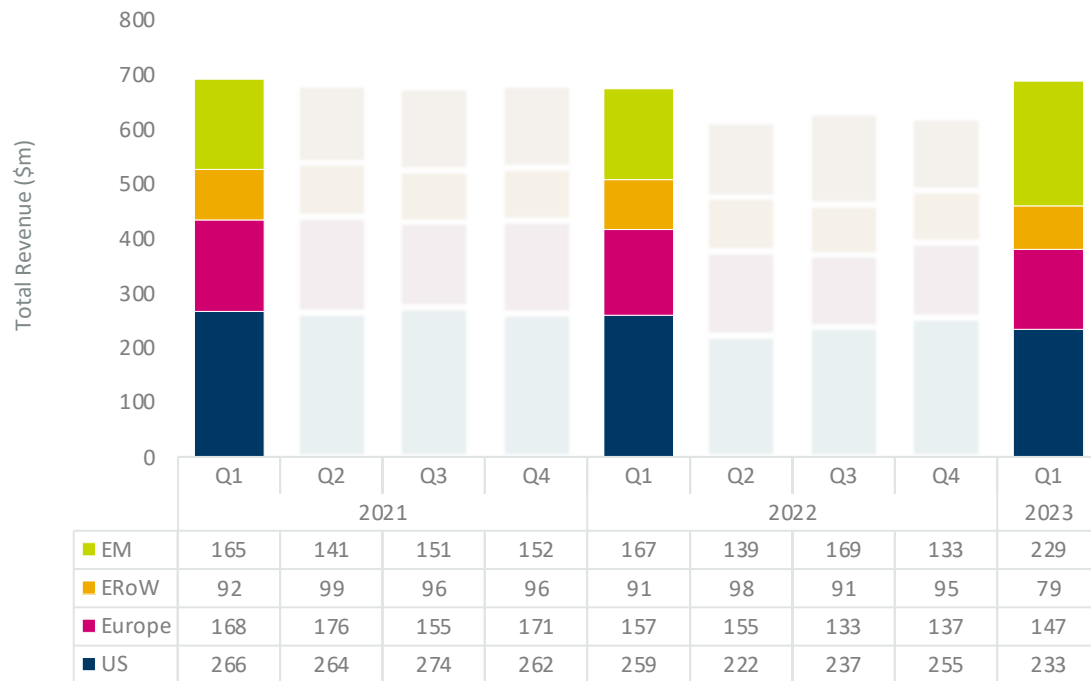
32% sequential growth at CER to \$54m



BioPharmaceuticals: Respiratory & Immunology

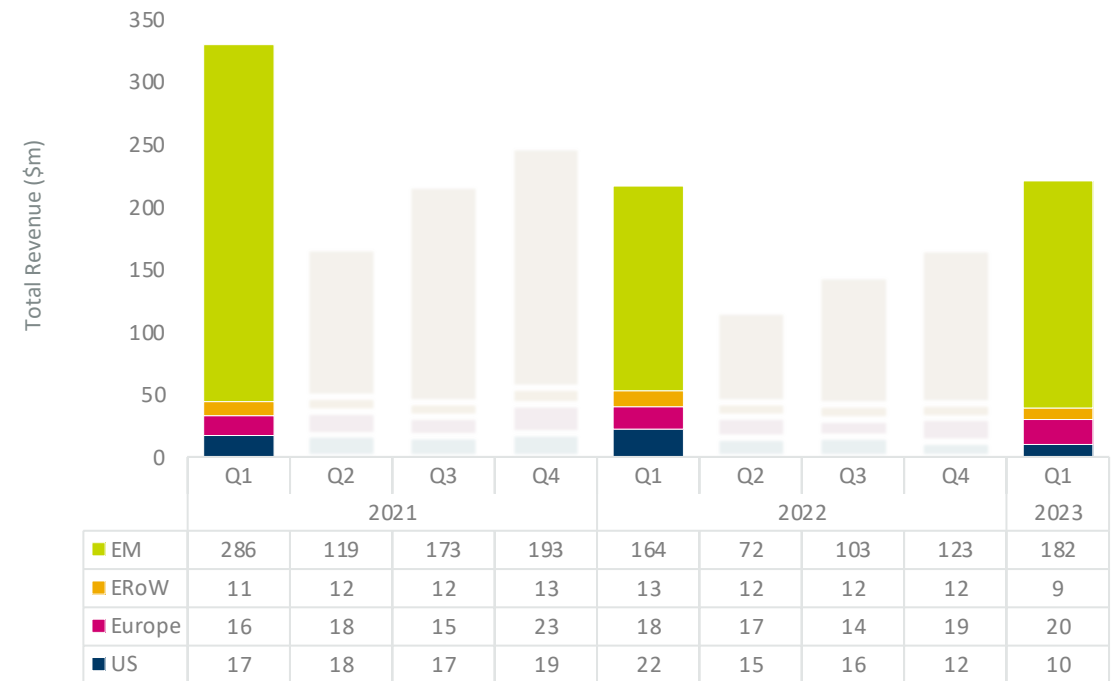
Symbicort

7% growth at CER to \$688m



Pulmicort

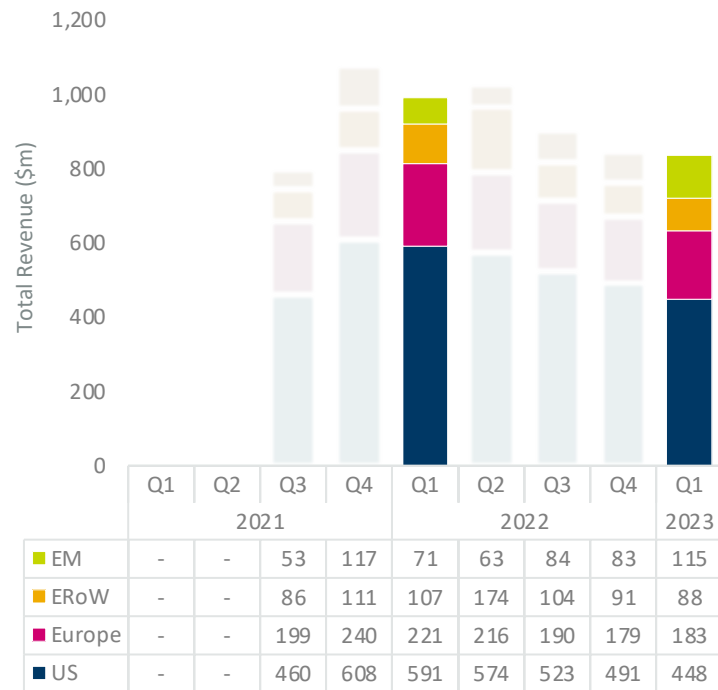
9% growth at CER to \$221m



Rare Disease

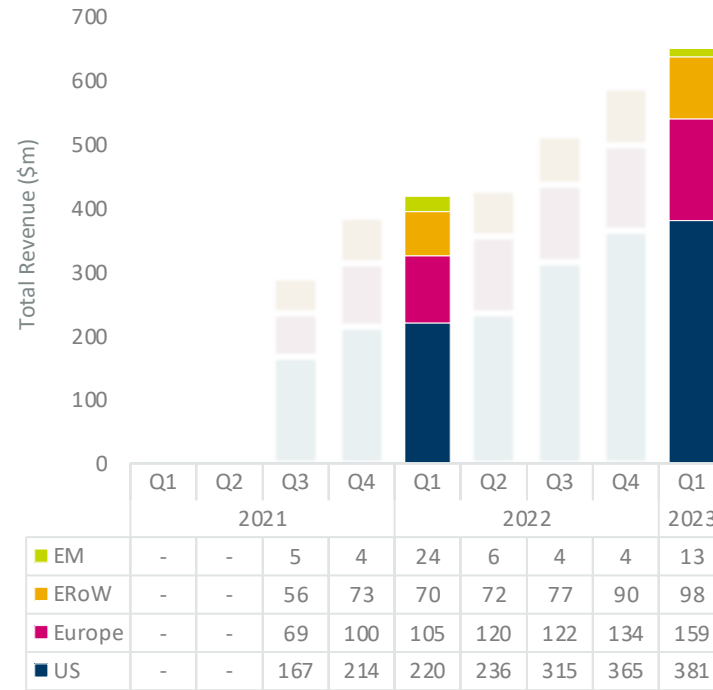
Soliris

13% decrease at CER to \$834m



Ultomiris

61% growth at CER to \$651m



Strensiq

28% growth at CER to \$262m

