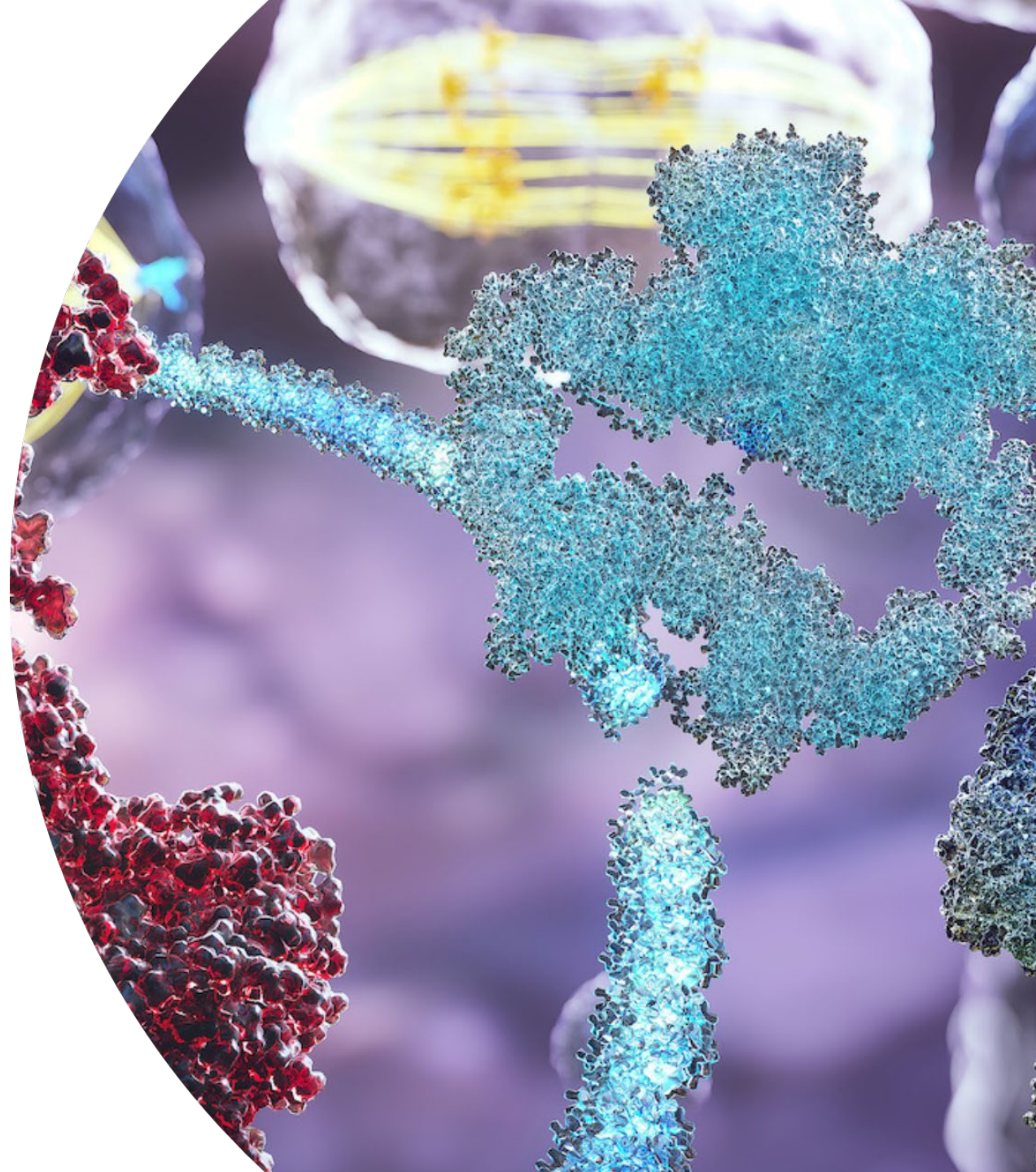




Q1 2024 Results

Conference call and webcast
for investors and analysts

25 April 2024



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 ability of the Group and Fusion to complete the transactions contemplated by the arrangement agreement with Fusion, including the parties' ability to satisfy the conditions set forth in the arrangement agreement with Fusion; the ability of the Group and Amolyt Pharma to complete the transactions contemplated by the acquisition agreement with Amolyt Pharma, including the parties' ability to satisfy the conditions set forth in the acquisition agreement with Amolyt Pharma; the Group's statements about the expected timetable for completing the acquisitions of Fusion and Amolyt Pharma; the Group's and Fusion's beliefs and expectations and statements about the benefits sought to be achieved in the Group's pending acquisition of Fusion; the Group's and Amolyt Pharma's beliefs and expectations and statements about the benefits sought to be achieved in the Group's proposed acquisition of Amolyt Pharma; the potential effects of the acquisition of Fusion on both the Group and Fusion and of the acquisition of Amolyt Pharma on both the Group and Amolyt Pharma; the possibility of any termination of the arrangement agreement with Fusion or of the acquisition agreement with Amolyt Pharma; the expected benefits and success of "FPI-2265" (Ac225-PSMA I&T) and any combination product or eneboparatide ("AZP-3601") and any combination product; the possibility that any milestone related to any contingent value right may take longer to achieve than expected or may never be achieved and the resulting contingent milestone payment may never be realized; the effects of disruption from the transactions contemplated by the acquisition agreement with Amolyt Pharma and the impact of the announcement and pendency of the transactions on Amolyt Pharma's business; the effects of disruption from the transactions contemplated by the arrangement agreement with Fusion and the impact of the announcement and pendency of the transactions on Fusion's business; the risk that shareholder litigation in connection with the offer or the acquisition may result in significant costs of defense, indemnification and liability; 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 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 the Group's 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 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. There can be no guarantees that the conditions to the closing of the proposed transaction with Fusion will be satisfied on the expected timetable or at all or that "FPI-2265" (Ac225-PSMA I&T) or any combination product will receive the necessary regulatory approvals or prove to be commercially successful if approved. There can be no guarantees that the conditions to the closing of the proposed transaction with Amolyt Pharma will be satisfied on the expected timetable or at all or that eneboparatide ("AZP-3601") will receive the necessary regulatory approvals or prove to be commercially successful if approved.



Q1 2024 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

Sharon Barr

EVP, BioPharmaceuticals R&D

Rare Disease

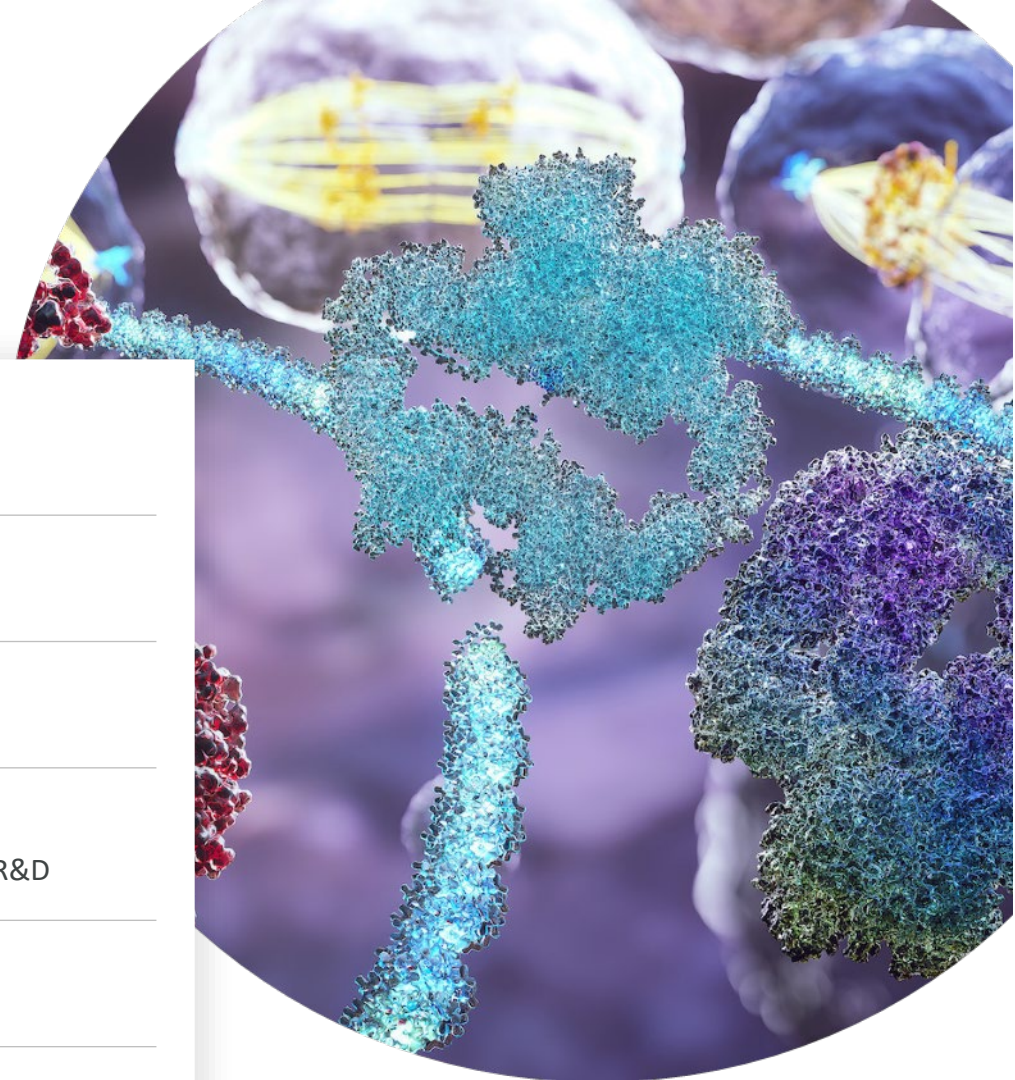
Marc Dunoyer

Chief Executive Officer, Alexion

CEO Closing Remarks, Q&A

Pascal Soriot

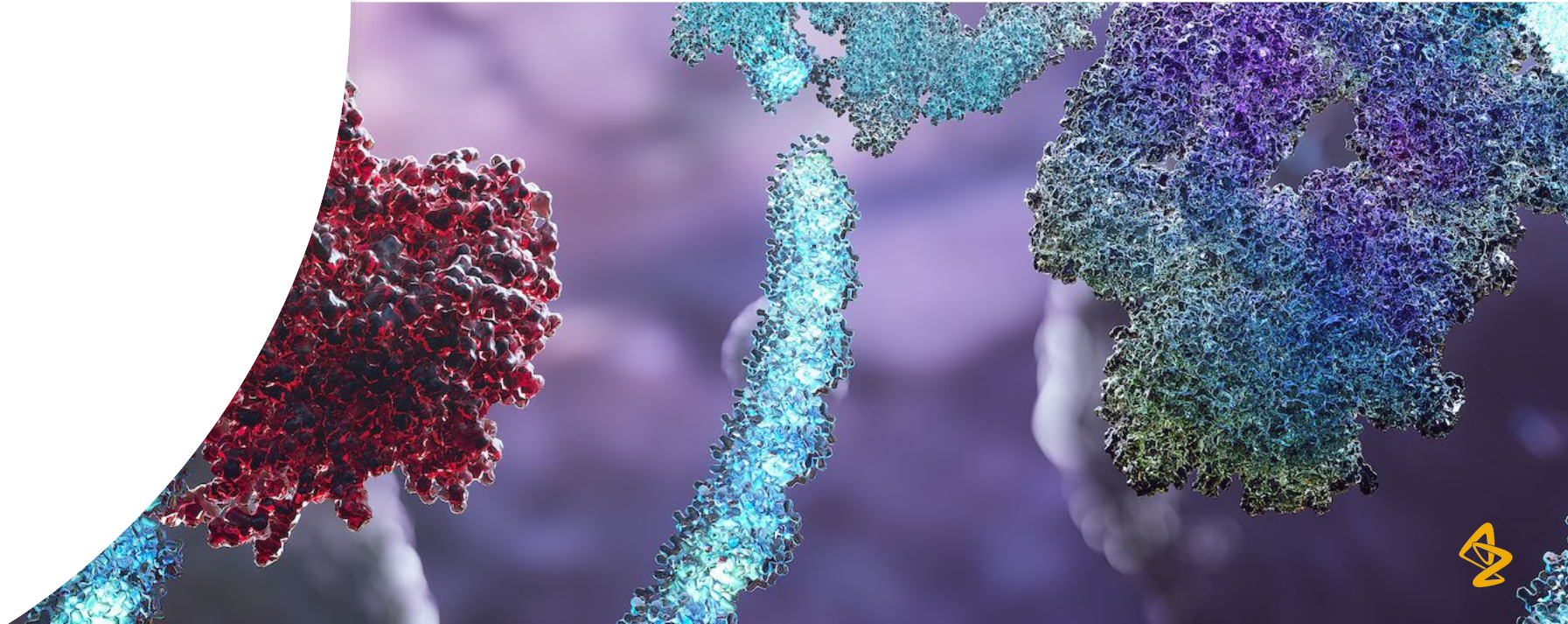
Chief Executive Officer



CEO Opening Remarks

Pascal Soriot

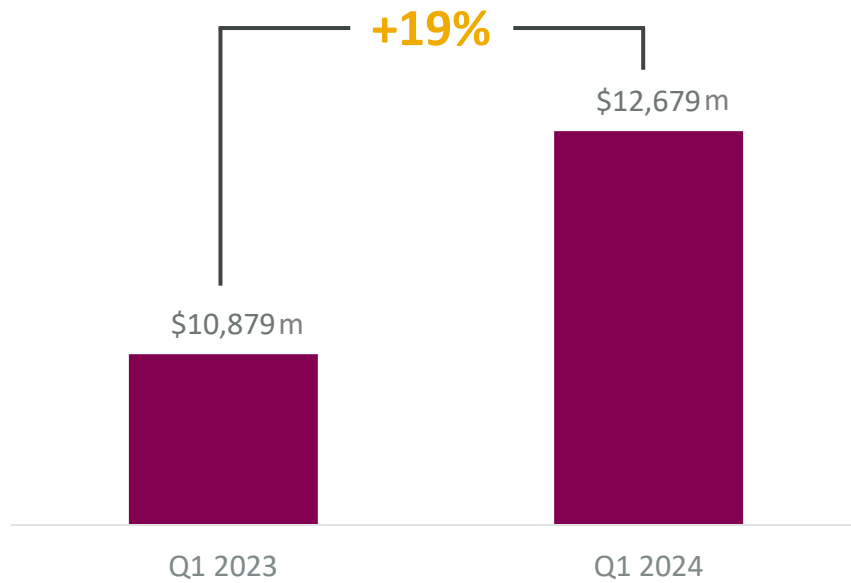
CHIEF EXECUTIVE OFFICER



Strong delivery in Q1 2024

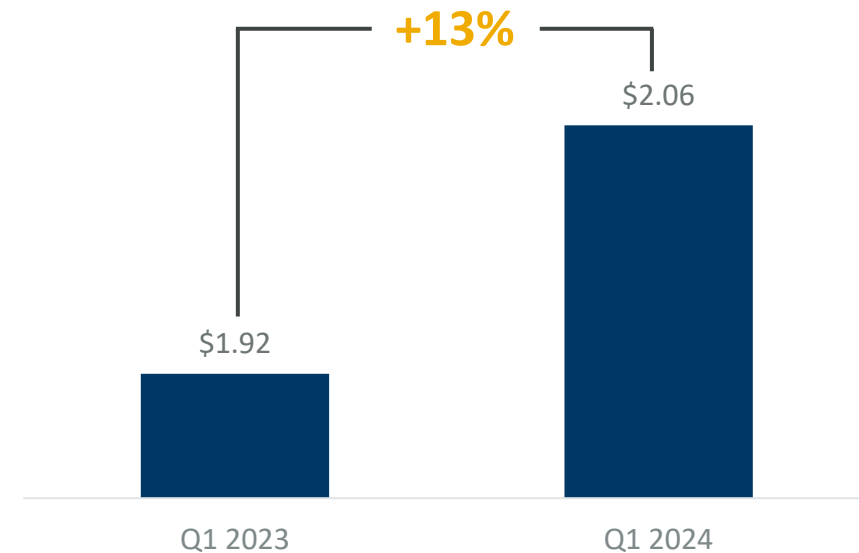
Total Revenue growth

fueled by global Product Sales



Core EPS growth

despite substantial decrease in Other Operating Income¹

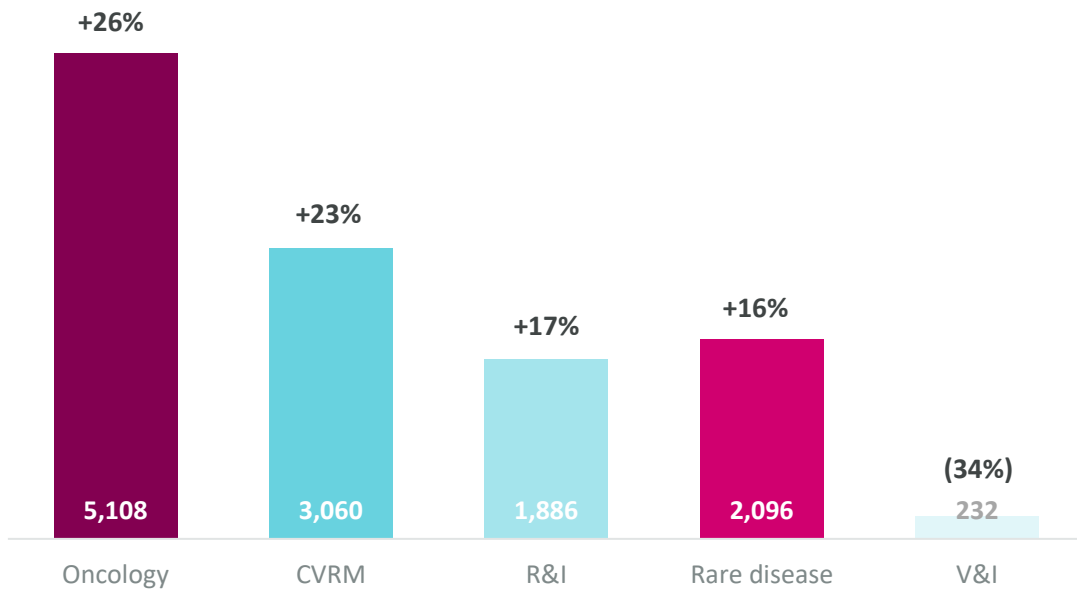


2024 dividend: 7% increase to \$3.10

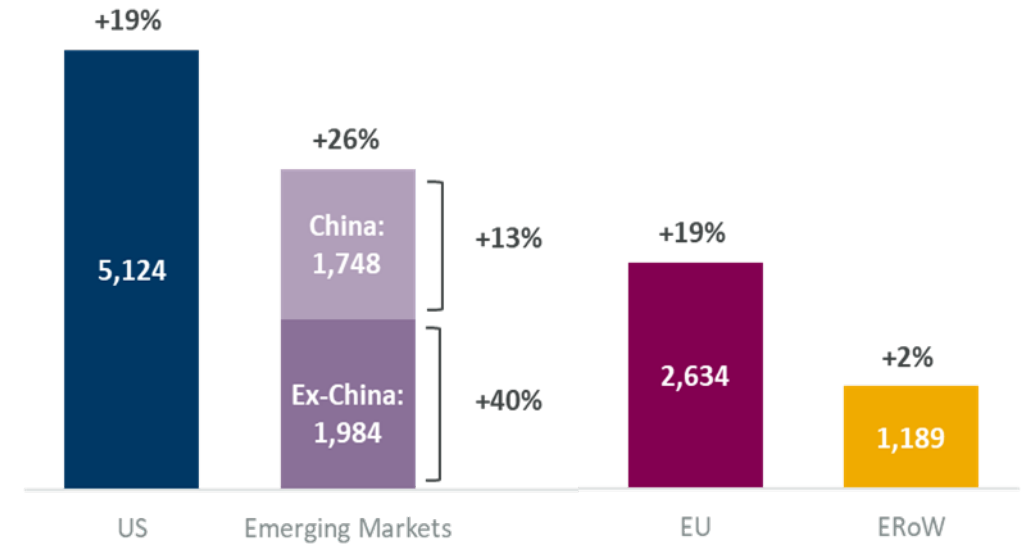


Growth across therapy areas and geographies

Total Revenue, \$m



Total Revenue, \$m



Double-digit growth across therapy areas

Strong Emerging Markets performance, +26%



Q1 2024 pipeline events unlock significant growth potential

Unprecedented Phase III results

Phase III ADRIATIC trial

- *Imfinzi* potentially first IO therapy in LS-SCLC

Phase III LAURA trial

- Reinforcing *Tagrisso* as backbone TKI, expanding in early-stage *EGFRm* NSCLC

2024 **ASCO**
ANNUAL MEETING

Phase III ADRIATIC and LAURA trials selected for plenary

Transformative new approvals

Tagrisso + CTx
1L *EGFRm* NSCLC

Enhertu
Tumour agnostic

Ultomiris
NMOSD

Continued investment in recent launches

 **WAINUA**

 **Truqap**

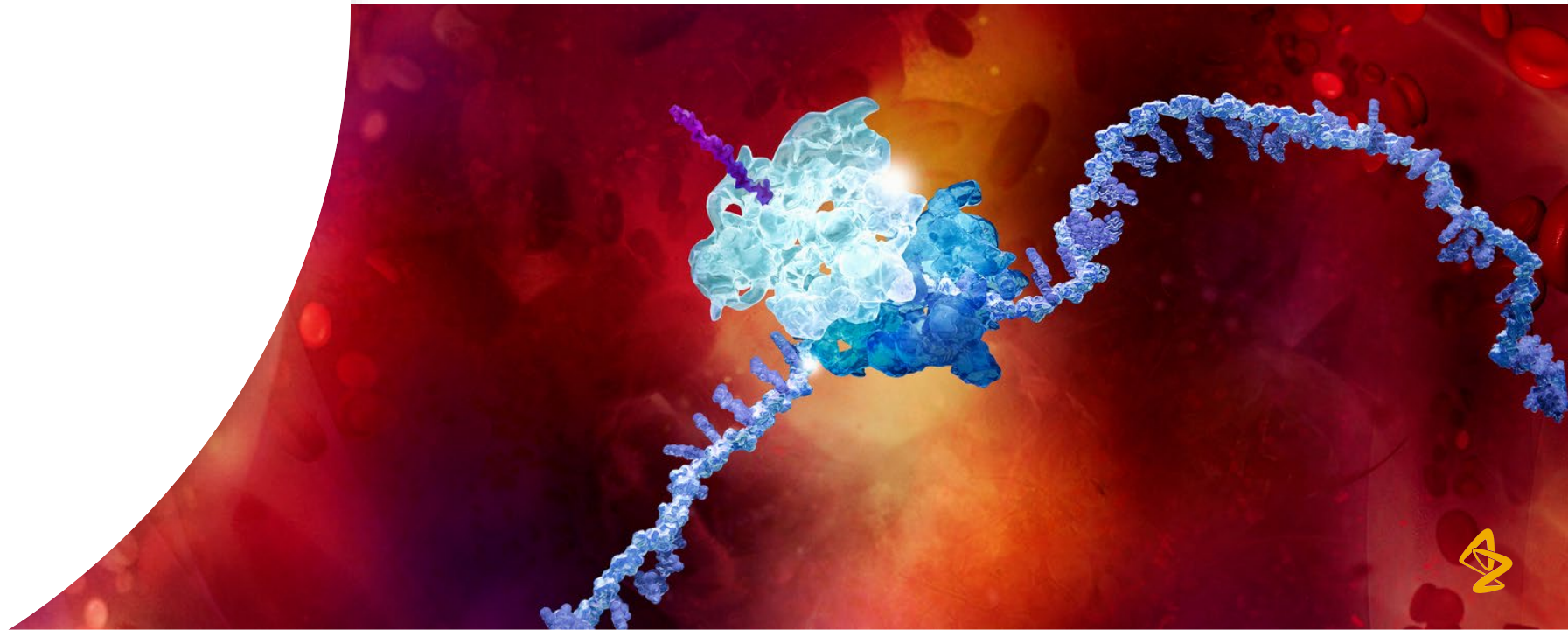
 **Voydeya**

 **AIRSUPRA**

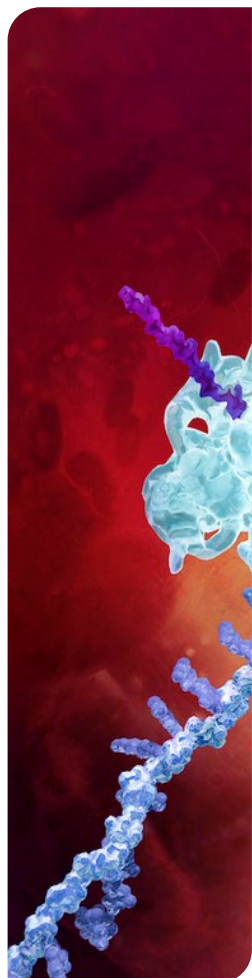


Financial Results

Aradhana Sarin
CHIEF FINANCIAL OFFICER



Q1 2024 – Reported profit and loss



	Q1 2024 \$m	CER change %	% Total Revenue
- Product Sales	12,177	18	96
- Alliance Revenue	457	59	4
- Collaboration Revenue	45	66	-
Total Revenue	12,679	19	100
Product Sales Gross Margin	81.8%	-	
Total operating expense ¹	(7,413)	10	58
- R&D expense	(2,783)	7	22
- SG&A expense	(4,495)	12	35
Other operating income and expense	67	(83)	1
Operating profit	3,115	31	25
Tax rate	22%		
Reported EPS	\$1.41	30	

Due to rounding, the sum of a number of dollar values and percentages may not agree to totals.

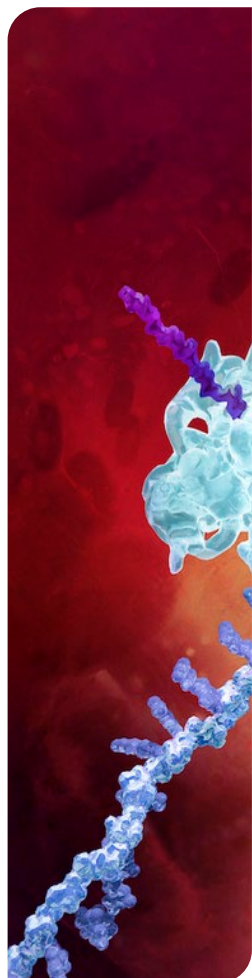
Absolute values at actual exchange rates; changes at CER. Product Sales Gross Margin excludes the impact of Alliance and Collaboration Revenue and any associated costs, thereby reflecting the underlying performance of Product Sales.

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

Appendix: [Glossary](#).



Q1 2024 – Core profit and loss



	Q1 2024 \$m	CER change %	% Total Revenue
- Product Sales	12,177	18	96
- Alliance Revenue	457	59	4
- Collaboration Revenue	45	66	-
Total Revenue	12,679	19	100
Product Sales Gross Margin	82.0%	-1pp	
Total operating expense ¹	(6,246)	15	49
- R&D expense	(2,698)	18	21
- SG&A expense	(3,413)	13	27
Other operating income and expense	65	(80)	1
Operating profit	4,310	15	34
Tax rate	21%		
Core EPS	\$2.06	13	

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. Product Sales 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 expense includes distribution, R&D and SG&A expenses.

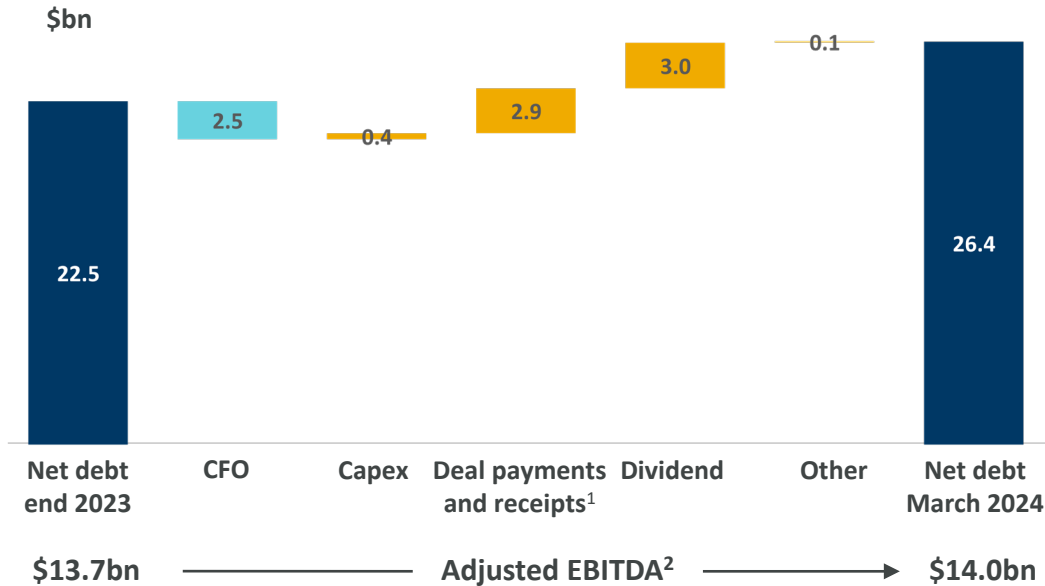
Appendix: [Glossary](#).



Net debt and FY guidance

Reiterating FY 2024 guidance

Net debt bridge



Net debt/Adjusted EBITDA 1.9x

FY 2024 guidance reiterated (CER)

Total Revenue

- Low double-digit to low teens percentage increase

Core EPS

- Low double-digit to low teens percentage increase

Anticipated FX impact: low single-digit adverse impact on Total Revenue and mid single-digit impact on Core EPS³

Due to rounding, the sum of a number of dollar values and percentages may not agree to totals. 1. Comprises disposal of intangible assets, movement in profit participation liability, purchase of intangible assets, payment of contingent consideration on business combinations, purchase and disposal of non-current asset investments, payment of Acerta Pharma share purchase liability and acquisitions of subsidiaries, net of cash acquired. The Company uses Debt issuance to finance new Business Development opportunities

11 2. Rolling 12m EBITDA adding back the impact of unwind of inventory fair value uplift recognised on acquisition of Alexion of \$78m (FY 2023: \$114m). AstraZeneca credit ratings: Moody's: short-term rating P-1, long-term rating A2, outlook stable. S&P Global Ratings: short-term rating A-1, long-term rating A, outlook stable. 3. If foreign exchange rates for April to December were to remain at the average rates seen in March 2024. Appendix: [Glossary](#).

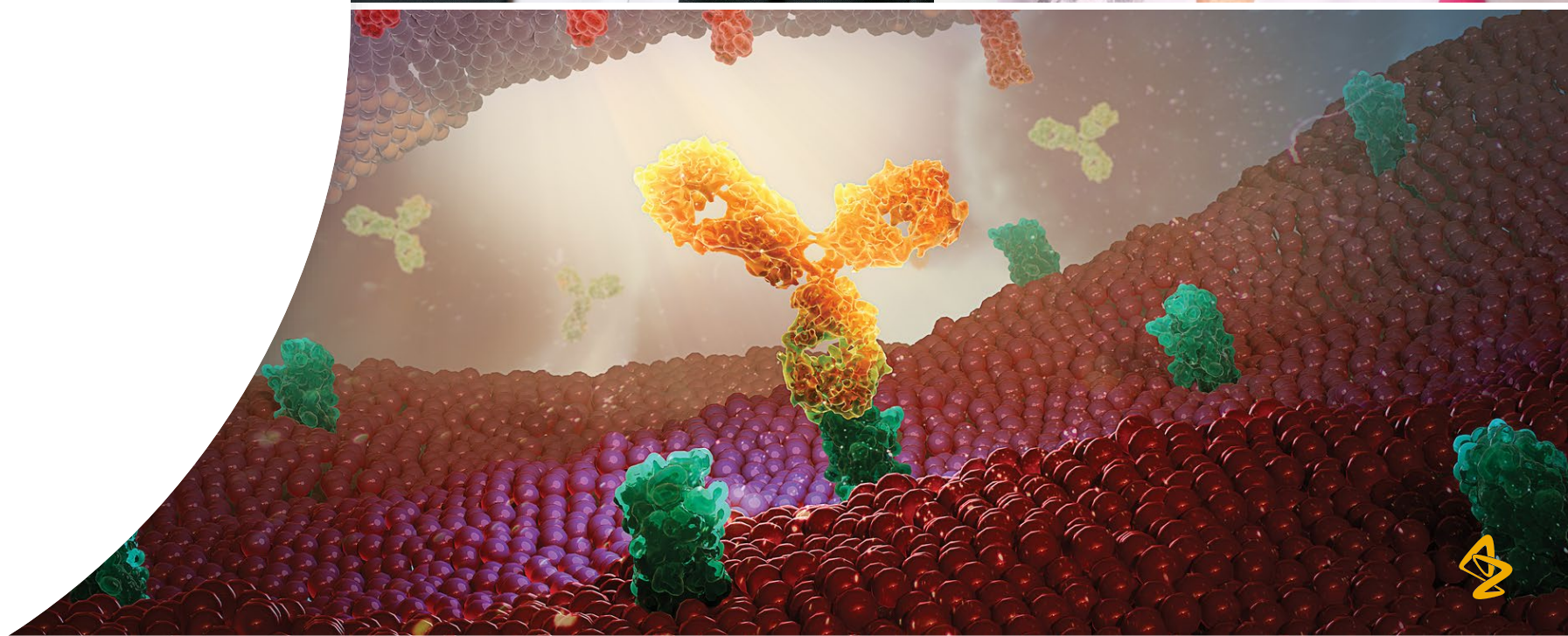


Oncology



Dave Fredrickson
ONCOLOGY BUSINESS

Susan Galbraith
ONCOLOGY R&D

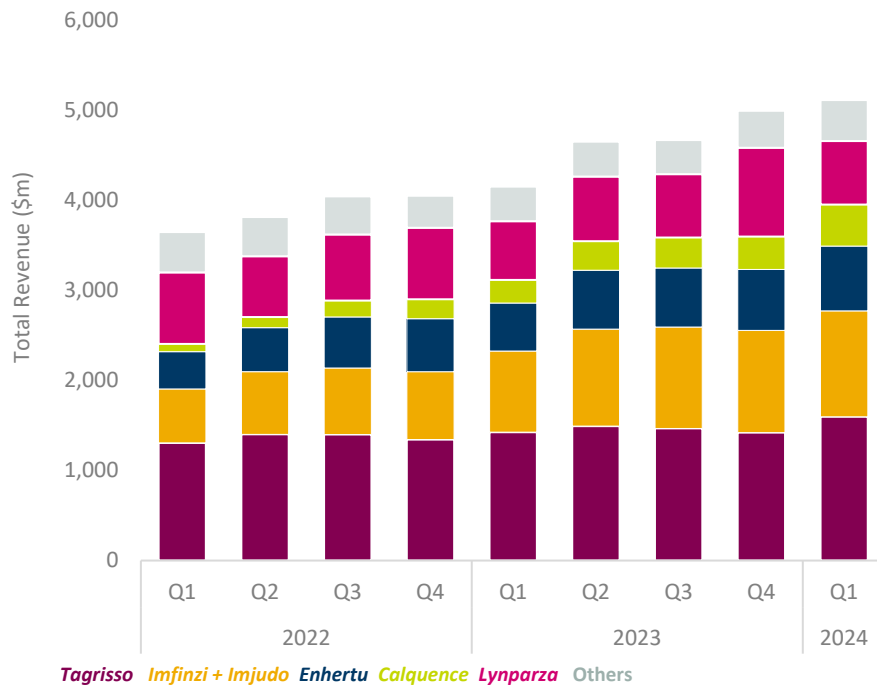


Oncology – Q1 2024

Total Revenue +26% with strong double-digit growth across all regions

Oncology

Q1 2024 \$5.1bn, +26%



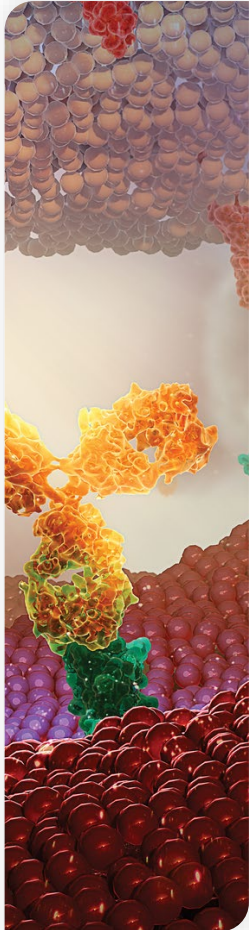
Q1 2024: key dynamics

- **Tagrisso** +15%, continued ADAURA and FLAURA demand, strong FLAURA2 awareness and early uptake in target patient segments
 - **Lynparza PS** +11%, continued PARPi leadership
 - **Imfinzi** +33%, achieved TOPAZ-1 (BTC) peak penetration in US, EU; JP repricing effective from February 1st
 - **Imjudo** +70%, HIMALAYA (HCC) acceleration, durable POSEIDON (NSCLC) demand
 - **Calquence** +35%, sustained BTKi leadership in 1L CLL
 - **Enhertu** +79%, NPS growth in HER2+ (DB03), one-time EU pricing benefit, strong initial mBC uptake in Emerging Markets
 - **Truqap** n/m, rapid adoption in core biomarker-altered population
-
- New indications: US (*Tagrisso* FLAURA2, *Enhertu* DPT02), JP (*Truqap* CAPitello-291)
 - ASCO 2024 Plenaries: *Imfinzi* ADRIATIC, *Tagrisso* LAURA



Oncology – R&D highlights

Fusion Pharmaceuticals acquisition expands next-gen Radioconjugate capabilities



30-50% of patients receive conventional radiotherapy¹

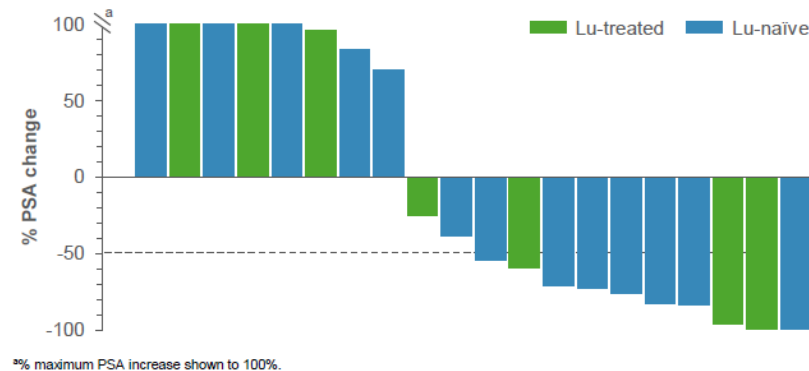
Clinical-stage portfolio, combination potential with next-gen IO and DDR

Accelerates AstraZeneca's Radioconjugate research and manufacturing to commercial build

FPI-2265 in Prostate cancer

PSMA-Actinium RC with potential post-Pluvicto and Pluvicto-naïve

Maximum percent PSA change from baseline during the treatment period (weeks 0-28) by prior Lu treatment²



- PSA50 achieved in 43% of Lu-treated participants
- PSA50 achieved in 54% of Lu-naïve participants
- No discontinuations from Xerostomia

Multi-blockbuster opportunity in mCRPC with FPI-2265

1. KANTAR data, US 2025 estimates (across lung, head and neck, prostate, bladder, endometrial, ovarian, colon, rectal and gastric cancers) 2. TATCIST data presented at AACR 2024.



BioPharmaceuticals

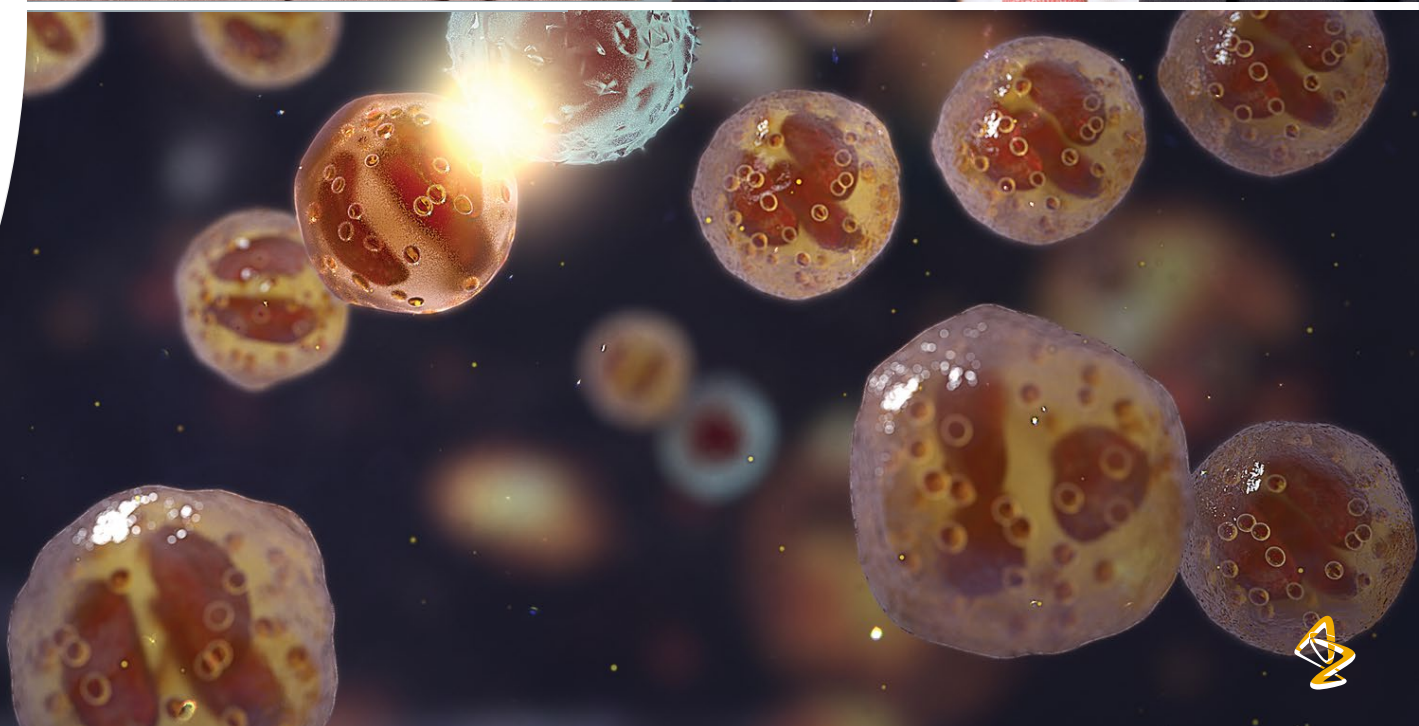


Ruud Dobber

BIOPHARMACEUTICALS BUSINESS

Sharon Barr

BIOPHARMACEUTICALS R&D

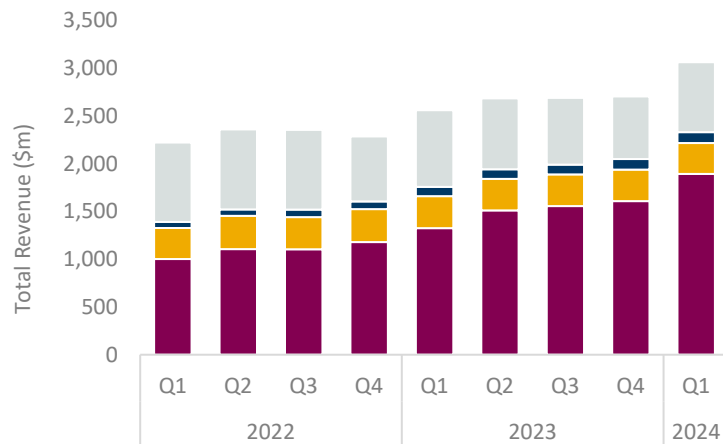


BioPharmaceuticals – Q1 2024

Total Revenue \$5.2bn, +16% – demand growth, accelerating new launch momentum

CVRM

Q1 2024 \$3.1bn, +23%



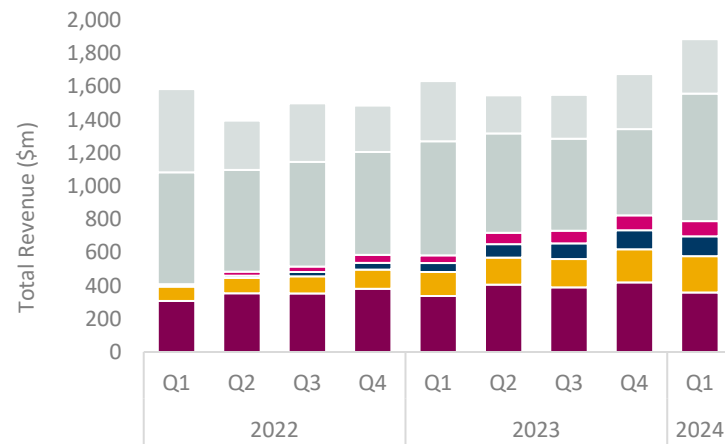
Farxiga Brilinta Lokelma Other

Q1 dynamics

- **Farxiga** +45%, demand growth outpacing SGLT2i
- **Lokelma** +19%, K+ Binder leadership in US
- **roxadustat** +28%, demand growth

R&I

Q1 2024 \$1.9bn, +17%



Fasenna Breztri Tezspire Saphnelo Symbicort Other

Q1 dynamics

- **Fasenna** +6%, continued IL-5 class leadership
- **Breztri** +54%, global market share gains
- **Tezspire** >2x, strong global launch demand



- >18k unique prescribers¹
- 65k TRx in 1Q24²



- Strong initial ATTRv-PN launch uptake



- 90% effective in preventing infant hospitalisation³

All growth rates at CER.

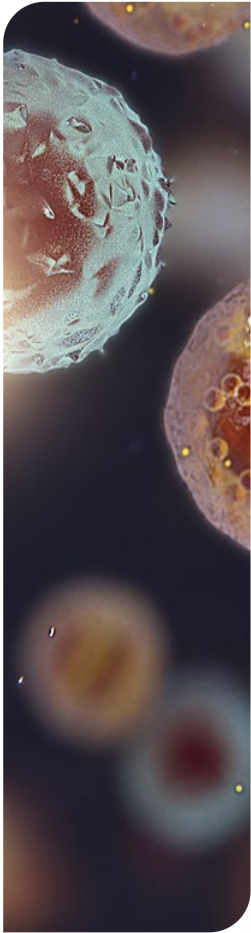
1. IQVIA Xponent unique prescribers data, week ending 29 March 2. IQVIA NPA Data Monthly Ending 24 March. 3. Moline, Tannis, Toepfer et. al. Centres for Disease Control and Prevention, "Early Estimate of Nirsevimab Effectiveness for Prevention of Respiratory Syncytial Virus–Associated Hospitalization Among Infants Entering Their First Respiratory Syncytial Virus Season — New Vaccine Surveillance Network, October 2023–February 2024." March 2024. Collaboration partners: Amgen (*Tezspire*); Sanofi (*Beyfortus*).

Appendix: [Glossary](#).



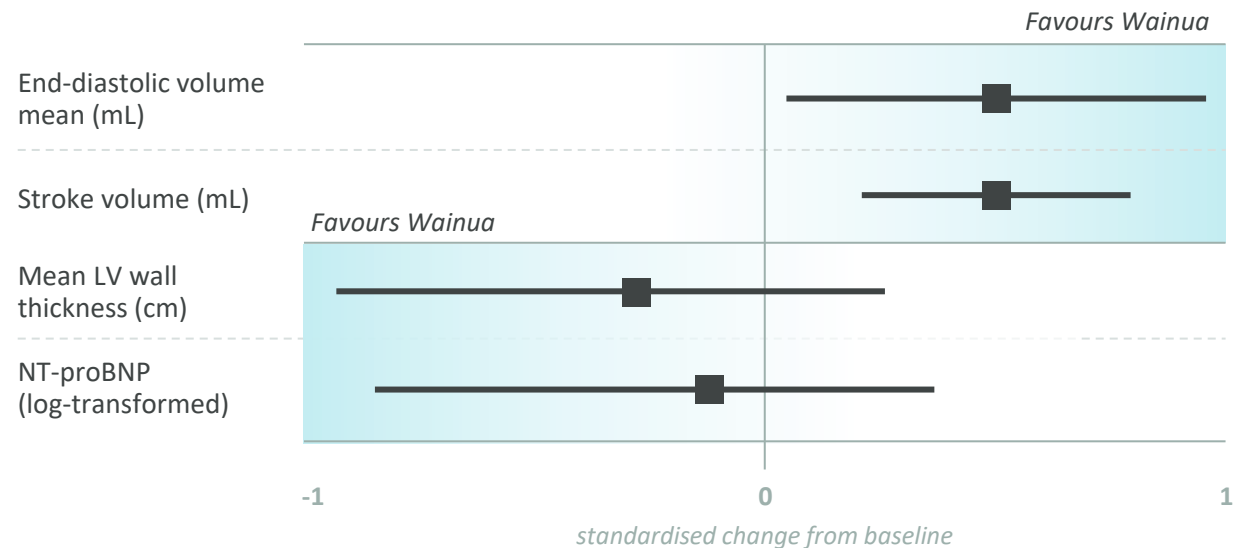
BioPharmaceuticals – R&D highlights

Exploratory endpoints support confidence in upcoming ATTR-CM opportunity



Phase III NEURO-TTRansform

Cardiomyopathic exploratory endpoints¹



Exploratory data supports potential efficacy in ATTR-CM

CARDIO-TTRansform largest and only Phase III trial to incorporate cardiovascular mortality endpoints

Primary endpoint: composite of CV mortality and recurrent CV events

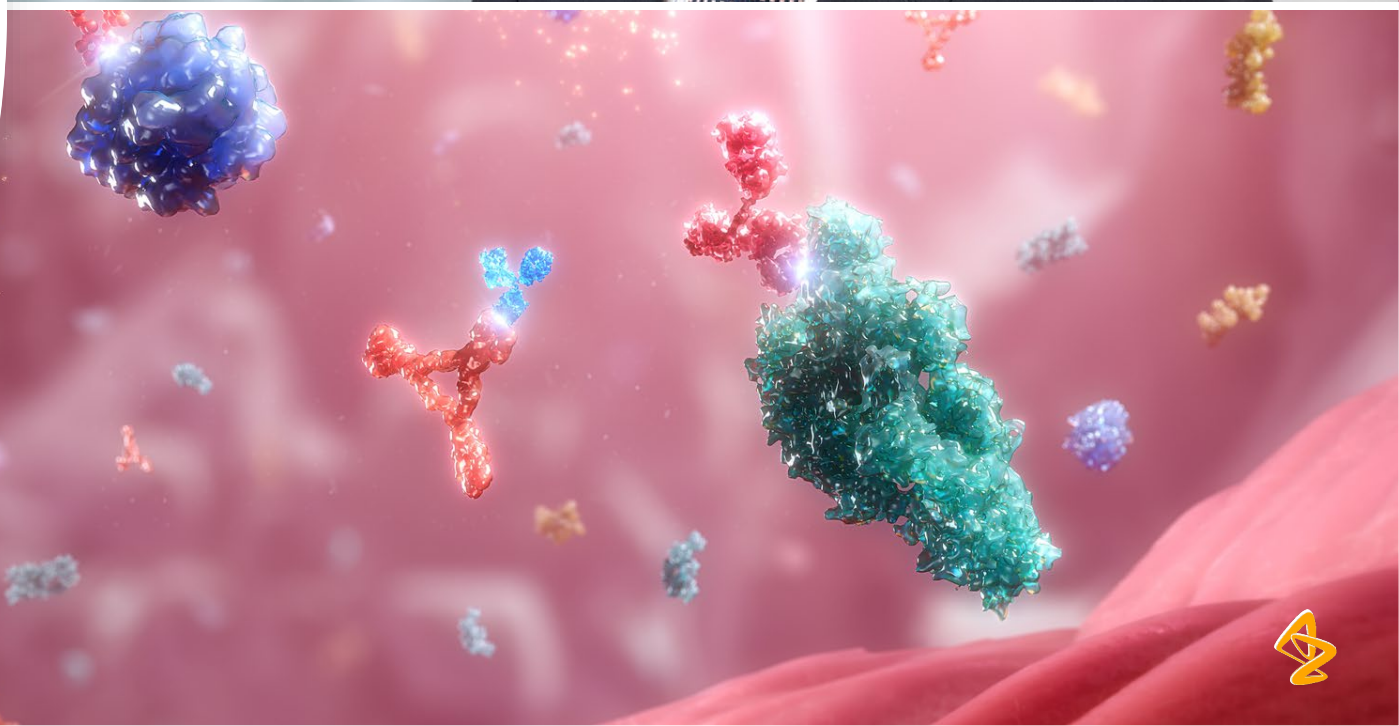
Secondary endpoint: change in 6MWT and KCCQ scores, rate of CV events, CV and all-cause mortality

Anticipate **CARDIO-TTRansform** trial readout 2025+

Rare Disease

Marc Dunoyer

CHIEF EXECUTIVE OFFICER,
ALEXION

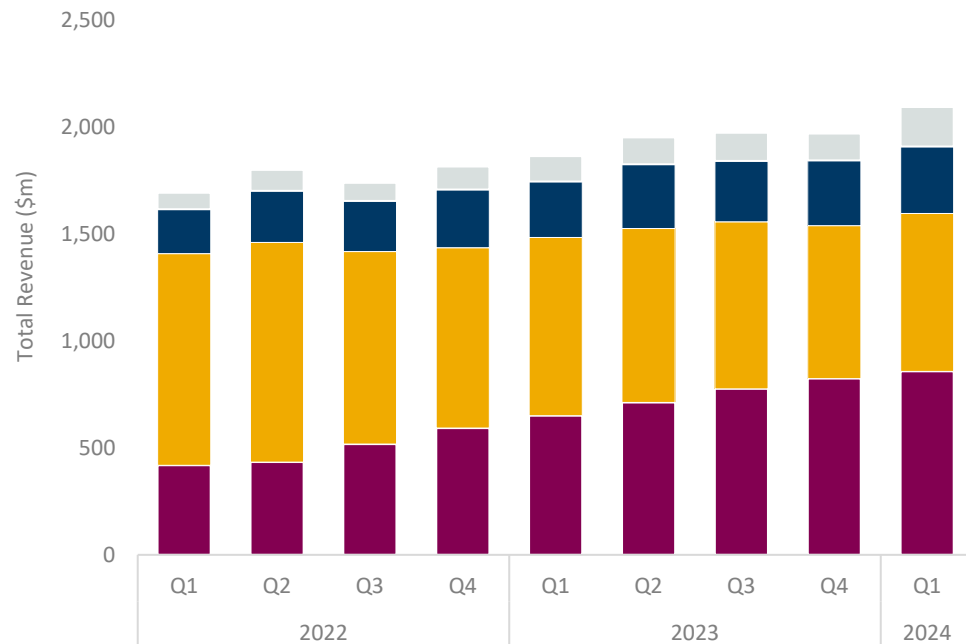


Rare Disease – Q1 2024

Total Revenue +16% in Q1 2024 driven by neurology

Rare Disease

Q1 2024 \$2.1bn, +16%



Ultomiris Soliris Strensiq Other¹

Q1 2024: key dynamics

Sustainable, durable growth of C5 Franchise

- **Ultomiris** +34%, driven by demand growth in neurology indications (gMG, NMOSD)
- **Soliris** (8%), continued conversion to *Ultomiris*, partly offset by growth in Emerging Markets

Strensiq +21% and **Koselugo** +82%, driven by continued global demand and order timing in certain Emerging Markets

-
- New indications: US (*Ultomiris* NMOSD, *Voydeya* PNH-EVH), EU (*Voydeya* PNH-EVH)

All growth rates at CER.

1. Includes *Kanuma* and *Koselugo*.

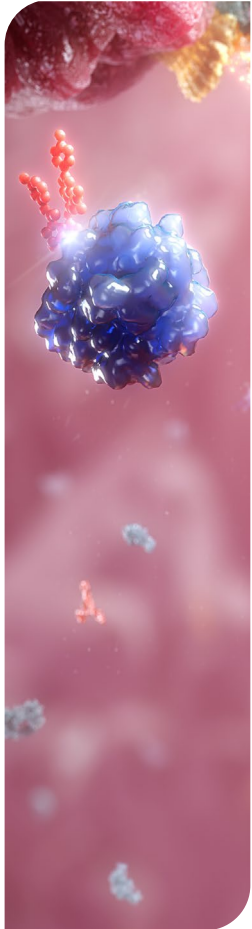
Collaboration partners: Merck & Co., Inc. (*Koselugo*).

Appendix: [Glossary](#).



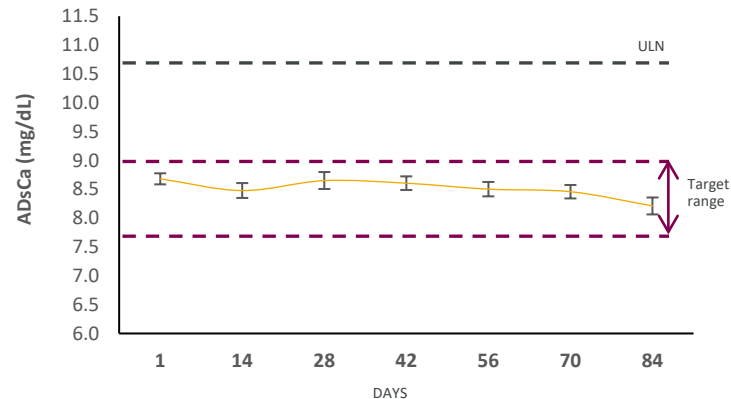
Rare Disease – R&D

Blockbuster opportunity with proposed Amolyt Pharma acquisition



eneboparatide | Phase IIa hypoparathyroidism

Serum calcium (mean ADsCa, mg/dL)



Clinical priorities:

- Normalising serum calcium levels
- Decreasing urinary calcium excretion
- Preserving bone mineral density

Potential **best-in-class therapy** with differentiated mechanism of action

Strong strategic fit, furthering commitment to rare endocrinology

Large unmet need:

US:
106k

EU5:
105k

JP:
42k

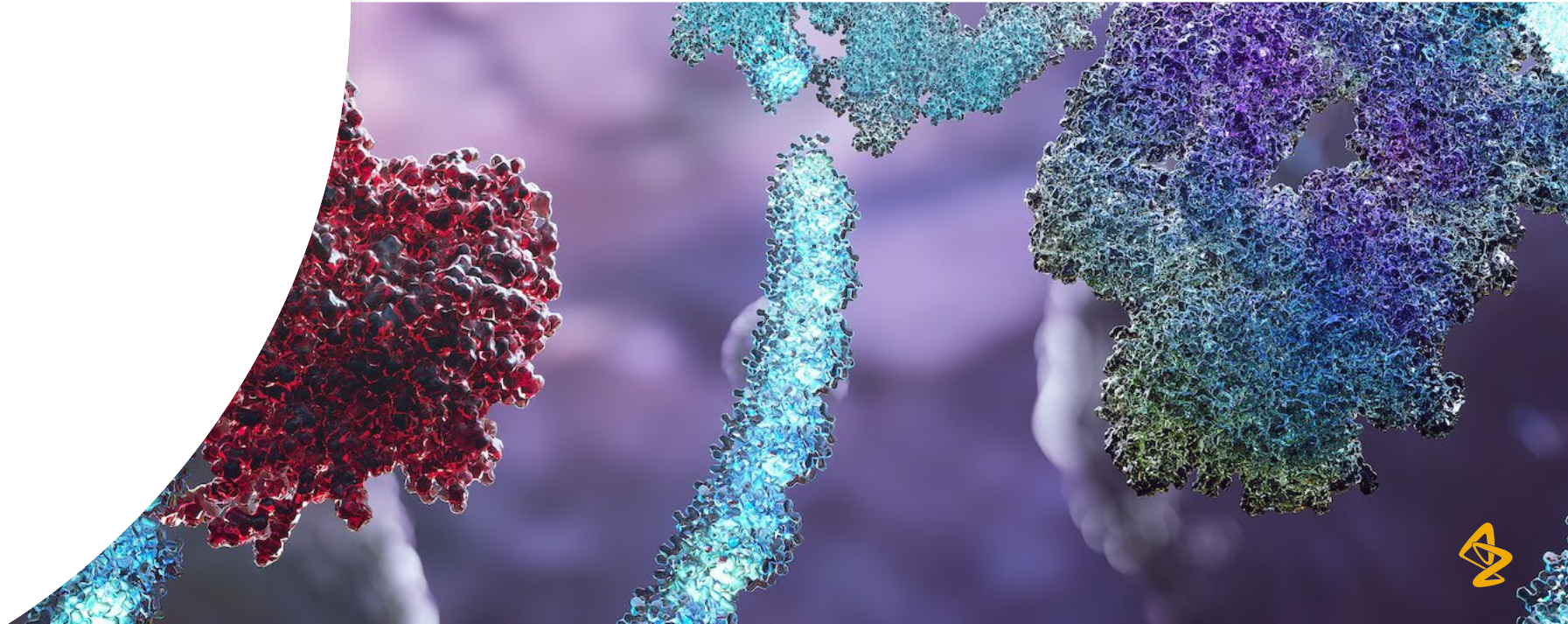
Anticipated Phase III CALYPSO data in 2025



CEO Closing Remarks

Pascal Soriot

CHIEF EXECUTIVE OFFICER



Increasing momentum of high-value pipeline

Significant pipeline catalysts over the remainder of 2024

DESTINY-Breast06 <i>Enhertu</i>	2L HER2-low breast cancer	H1 2024
CAPitello-290 <i>Truqap</i>	Triple Negative Breast Cancer	H1 2024
CAPitello-281 <i>Truqap</i>	PTEN deficient mHSPC	H2 2024
TROPION-Breast02 Dato-DXd	Triple Negative Breast Cancer	H2 2024
WAYPOINT <i>Tezspire</i>	Chronic Rhinosinusitis with Nasal Polyps	H2 2024

Phase III trial initiations since FY2023

TROPION-Lung10 <i>Nonsq PDL1-high</i>	Dato-DXd ± rilvegostomig	TROPION-Lung14¹ <i>1L EGFRm NSCLC</i>	Dato-DXd ± Tagrisso
CLARITY-Gastric01 <i>2L+ Gastric</i>	AZD0901	BalanceD-HF <i>HF with renal imp</i>	balcinrenone / dapagliflozin FDC
ICAN <i>IgAN</i>	Ultomiris	BaxDuo-Arctic <i>CKD with HTN</i>	baxdrostat / dapagliflozin FDC
THARROS <i>COPD</i>	Breztri	CHESTNUT <i>HPP</i>	efzimfotase alfa

Pipeline enhanced with recent BD



Q&A 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



Sharon Barr
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 & Glossary

- Glossary
- ESG – summary of sustainability progress
- Oncology tumour maps
- Emerging Markets performance
- Key medicines performance by therapy area



Glossary – abbreviations

1L	=	First-line or Front-line	hMPV/RSV	=	Human metapneumovirus/respiratory syncytial virus
2L	=	Second-line	HPP	=	Hypophosphatasia
6MWT	=	6-minute walk test	HtN	=	Hypertension
ADsCa	=	Albumin-adjusted serum calcium	IgAN	=	Immunoglobulin A Nephropathy
ASCO	=	American society of clinical oncology	IL5	=	Interleukin 5
ATTR-CM	=	Transthyretin amyloid cardiomyopathy	IO	=	Immuno-oncology
ATTR-PN	=	Transthyretin amyloid polyneuropathy	JP	=	Japan
BTC	=	Biliary tract cancer	K+	=	Potassium
BTKi	=	Bruton's tyrosine kinase	KCCQ	=	Kansas City Cardiomyopathy Questionnaire
C5	=	Complement component 5	LS-SCLC	=	Limited stage small-cell lung cancer
Capex	=	Capital expenditure	LV	=	Left ventricular
CER	=	Constant exchange rates	mBC	=	Metastatic breast cancer
CFO	=	Cash flow from operations	mg/dL	=	Miligrams per decilitre
CKD	=	Chronic kidney disease	mCRPC	=	Metastatic castration-resistant prostate cancer
CLL	=	Chronic lymphocytic leukemia	mHSPC	=	Metastatic hormone sensitive prostate cancer
CLDN18.2	=	Claudin 18 isoform 2	mL	=	Milliliter
cm	=	Centimeter	n/m	=	Not material
COPD	=	Chronic obstructive pulmonary disease	NMOSD	=	Neuromyelitis optica spectrum disorder
CTx	=	Chemotherapy	Nonsq	=	Non-squamous
CV	=	Cardiovascular	NSCLC	=	Non-small cell lung cancer
CVRM	=	Cardiovascular, renal and metabolism	NT-proBNP	=	N-terminal pro-B-type natriuretic peptide
DDR	=	DNA damage response	PARPi	=	Poly-ADP ribose polymerase inhibitor
EBITDA	=	Earnings before interest, tax, depreciation and amortisation	PDL1	=	Programmed cell death ligand 1
EGFRm	=	Epidermal growth factor receptor mutation	PNH-EVH	=	Paroxysmal nocturnal hemoglobinuria with extravascular haemolysis
EPS	=	Earnings per share	PSA	=	Prostate-specific antigen
ERoW	=	Established rest of world	PSA50	=	Prostate-specific antigen 50
EU	=	Europe	PTEN	=	Phosphatase and TENsin homolog deleted on chromosome 10
FDC	=	Fixed dose combination	R&I	=	Respiratory and immunology
FX	=	Foreign exchange	RC	=	Radioconjugates
gMG	=	Generalised myasthenia gravis	Renal imp	=	Renal impairment
HCC	=	Hepatocellular carcinoma	SGLT2i	=	Sodium/glucose cotransporter 2 inhibitor
HF	=	Heart Failure	Stg. III u/r NSCLC	=	Stage III unresectable non-small cell lung cancer
HLR	=	High-level results	TKI	=	Tyrosine kinase inhibitor
			ULN	=	Upper limit of normal
			V&I	=	Vaccines and immune therapies



2023 Sustainability highlights

Progress on our overall strategy includes:

15

public and private sector organisations convened by AstraZeneca CEO through the Sustainable Markets Initiative 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 Annex are “on plan”

Access to Healthcare

127,384

healthcare workers and others trained¹ (cumulative)

By 2025: 170,000

>66.4m

people reached through Access to Healthcare programmes (cumulative)¹

By 2025: 50m

>13.6m

people reached through our patient assistance programmes (cumulative)

Environmental Protection

67.6%

reduction in Scope 1 and Scope 2 greenhouse gas emissions

By 2026: 98% from 2015 base year

19.5%

reduction in our water use

By 2025: 20% below 2015 baseline

13.2%

reduction in our waste

By 2025: 10% below 2015 baseline

Ethics and Transparency

50.1%

senior middle management roles held by women

By 2025: reach gender equality in management positions

11 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



AstraZeneca in Lung Cancer

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</p> <p><i>Enhertu</i> + IO + CTx DESTINY-Lung03</p> <p>volrustomig + CTx eVOLVE-Lung02</p> <p>rilvegostomig (PD1/TIGIT) ARTEMIDE-1</p>	<p><i>Imfinzi</i> + ceralasertib LATIFY</p> <p>Dato-DXd TROPION-Lung01</p> <p>AZD9592 (EGFR/cMET ADC) EGRET</p> <p>sabestomig (PD1/TIM3)</p>	
EGFRm c.16%	<p><i>Tagrisso</i> ADAURA</p> <p><i>Tagrisso</i> neoADAURA</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 tumour drivers c.12%						<p>Dato-DXd TROPION-Lung01 TROPION-Lung05</p>
HER2m c.2%				<p>CRT → <i>Imfinzi</i> PACIFIC</p>	<p><i>Enhertu</i> DESTINY-Lung04</p>	<p><i>Enhertu</i> DESTINY-Lung02</p>

/// established SoC


Leading the future of lung cancer treatment

- Establishing *Tagrisso* as backbone TKI 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 + ADCs
- Investing behind new technologies and platforms, including cell therapy and testing/screening



AstraZeneca in Breast Cancer

Ambition to eliminate breast cancer as a cause of death

 established SoC	Early		1st line	Metastatic	3rd line	4th line +
	Noadjuvant	Adjuvant		2nd line		
Est. epi (G7)	540k		125k	90k	65k	55k
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-Breast02	
HR-positive 65-75% --- HER2-low 1+, 2+ 60%		Low risk Good outcomes with current SoC CTx → camizestrant (± CDK4/6i) CAMBRIA-2 CTx → AI (± CDK4/6i) 2-5 yrs → camizestrant CAMBRIA-1	RECURRENCE camizestrant + CDK4/6i SERENA-4 ESR1m AI + CDK4/6i → camizestrant + CDK4/6i SERENA-6 <i>Truqap</i> + <i>Faslodex</i> + CDK4/6i CAPitello292	PIK3CA/ AKT1/ PTEN alt. <i>Truqap</i> + <i>Faslodex</i> CAPitello291 <i>Enhertu</i> DESTINY-Breast06 HER2-low IHC 0-1+, 1+, 2+	Dato-DXd TROPION-Breast01 <i>Enhertu</i> DESTINY-Breast04 HER2-low IHC 1+, 2+	
TNBC 10-15% --- HER2-low 1+, 2+ 35%	Dato-DXd + <i>Imfinzi</i> TROPION-Breast04	NST → residual disease → Dato-DXd ± <i>Imfinzi</i> TROPION-Breast03	<i>Truqap</i> + paclitaxel CAPitello290 PD-L1+ 40% Dato-DXd + <i>Imfinzi</i> TROPION-Breast05 PD-L1- 60% Dato-DXd TROPION-Breast02	HER2-Low		
gBRCAm 5% of HR-positive 15% of TNBC		CTx → <i>Lynparza</i> OlympiA	<i>Lynparza</i> OlympiAD			

All numbers are approximate. Illustrative settings and populations, not to scale.

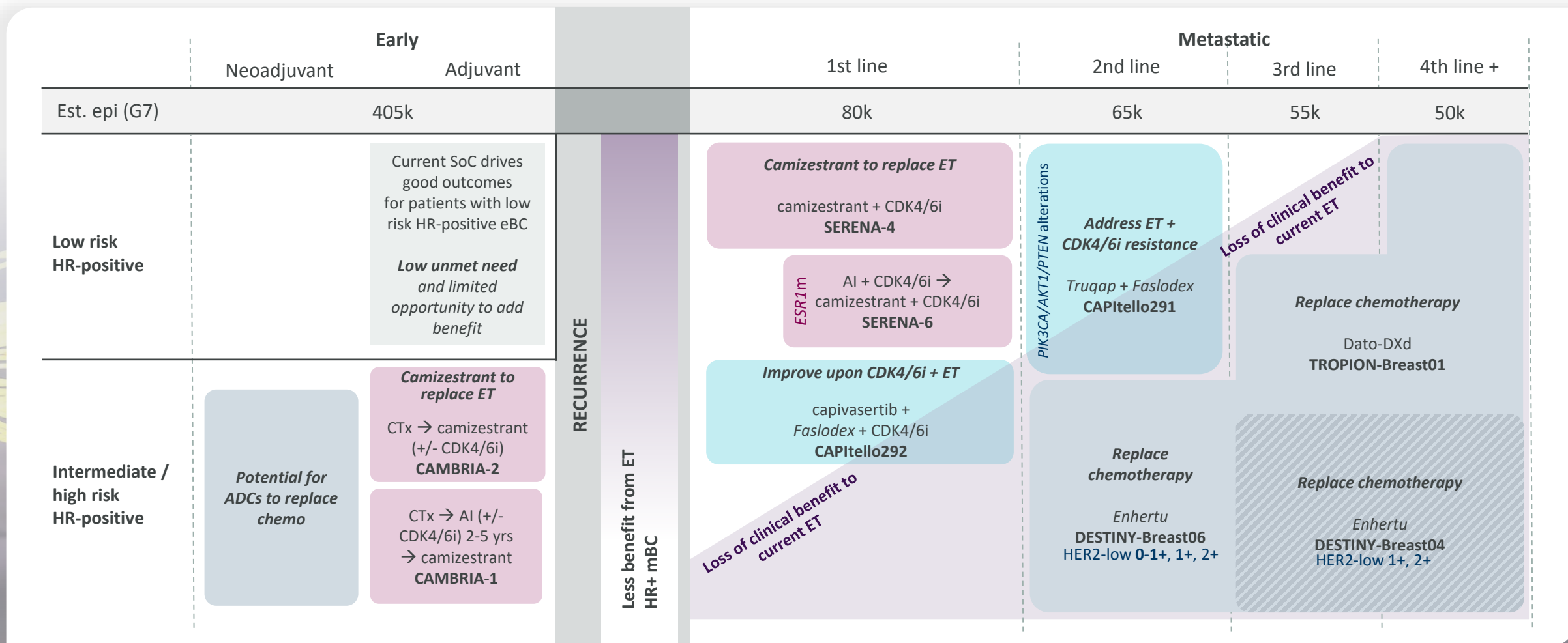
28 Collaboration partners: Daiichi Sankyo (*Enhertu*, Dato-DXd), Merck & Co., Inc. (*Lynparza*).

Appendix: [Glossary](#).



AstraZeneca in Breast Cancer

Ambition to eliminate breast cancer as a cause of death



All numbers are approximate. Illustrative settings and populations, not to scale.

29 Collaboration partners: Daiichi Sankyo (Enhertu, Dato-DXd).

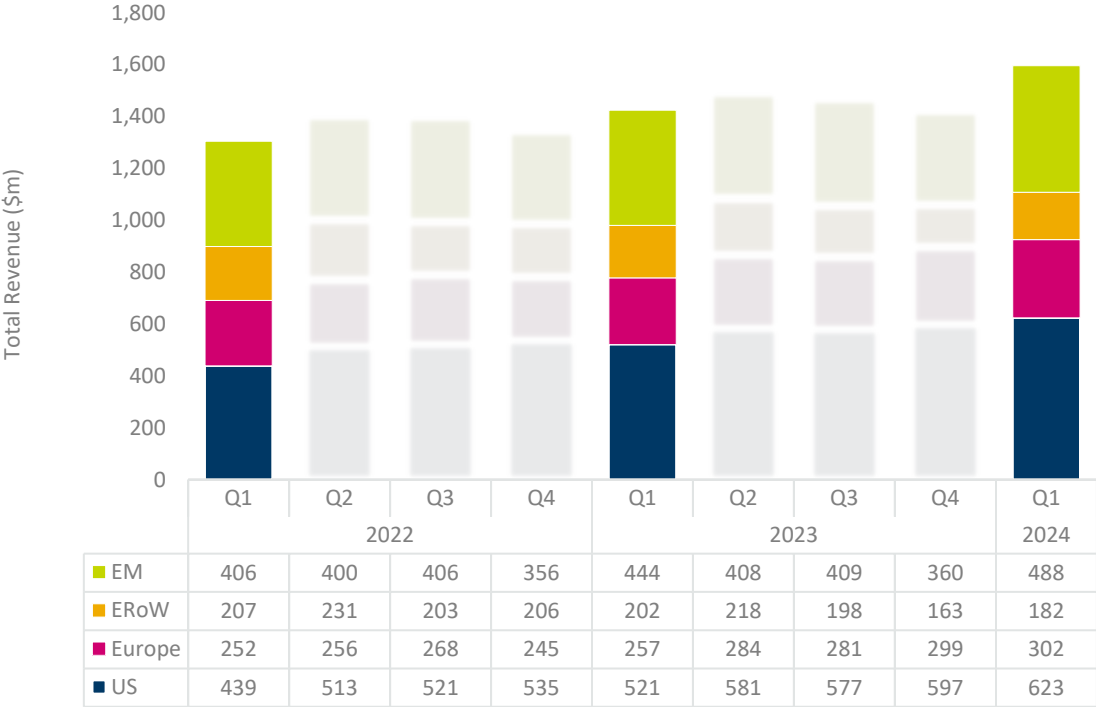
Appendix: [Glossary](#).



Oncology

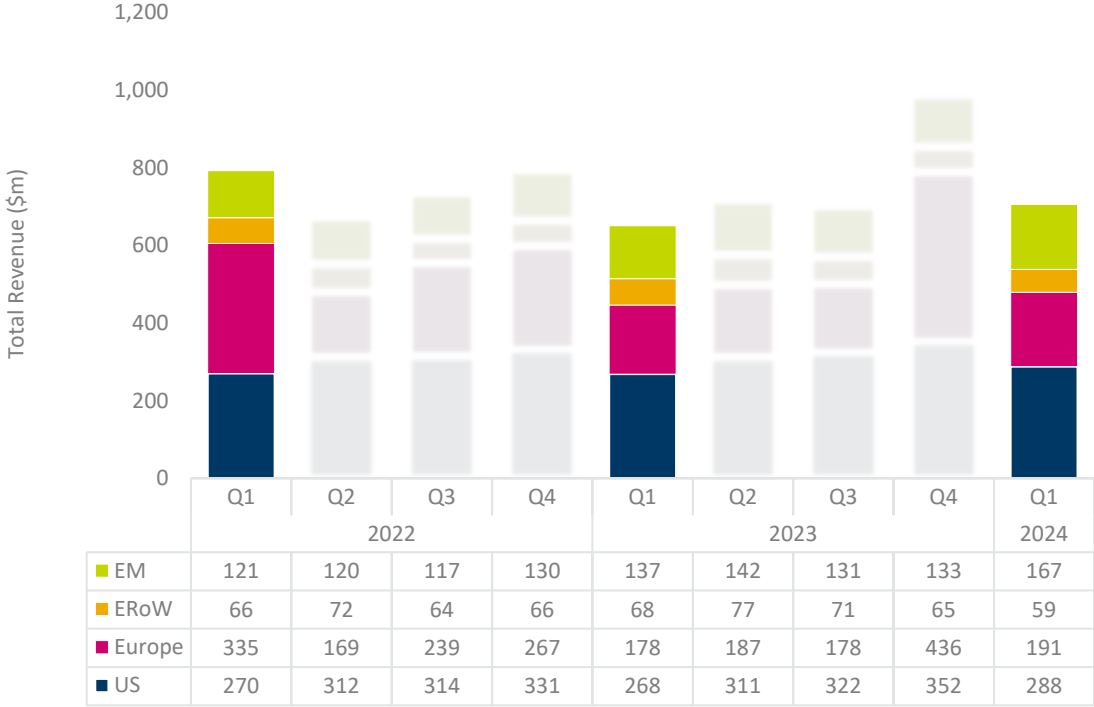
Tagrisso

15% growth at CER to \$1,595m in Q1 2024



Lynparza

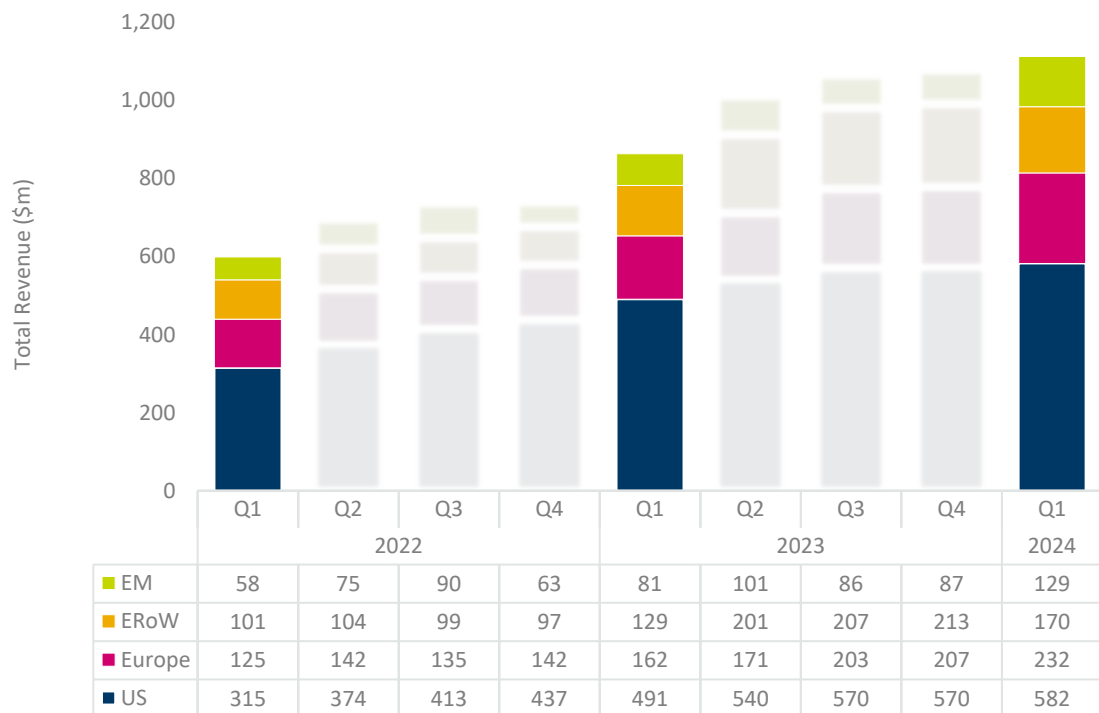
11% growth at CER to \$705m in Q1 2024



Oncology

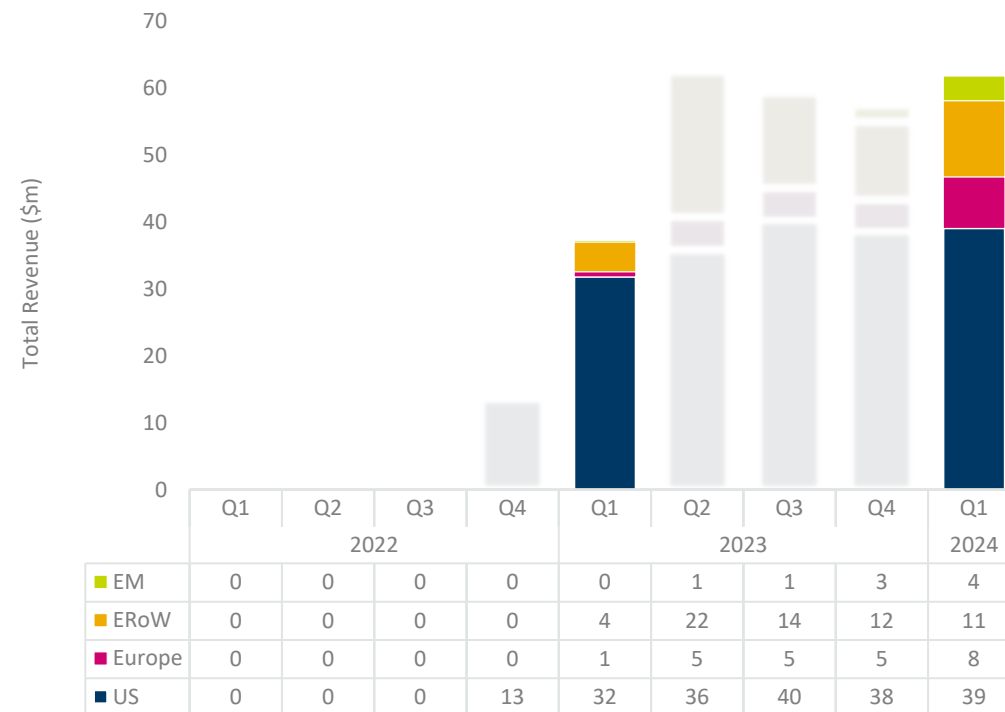
Imfinzi

33% growth at CER to \$1,113m in Q1 2024



Imjudo

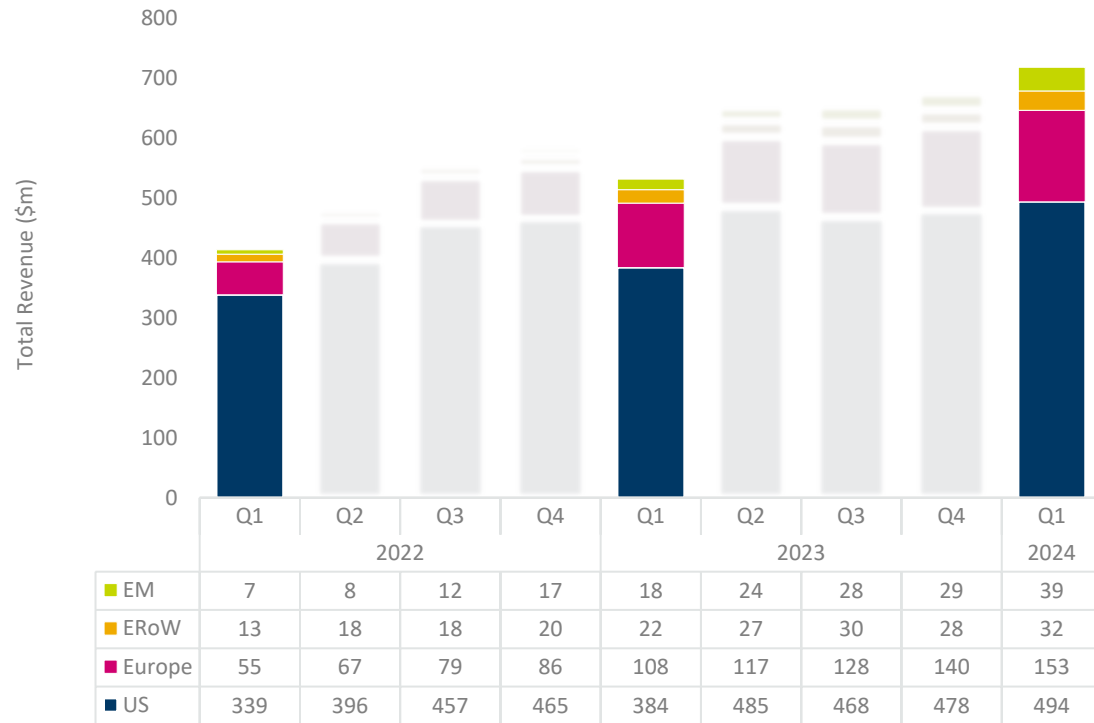
70% growth at CER to \$62m in Q1 2024



Oncology

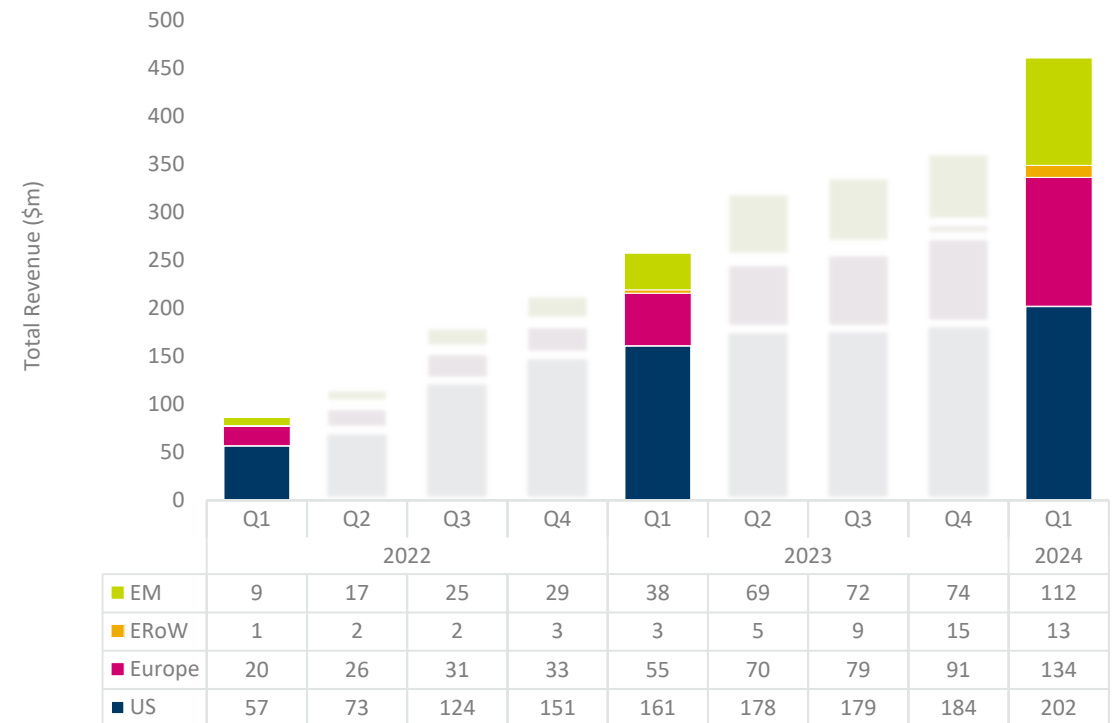
Calquence

35% growth at CER to \$718m in Q1 2024



Enhertu

79% growth at CER to \$461m in Q1 2024



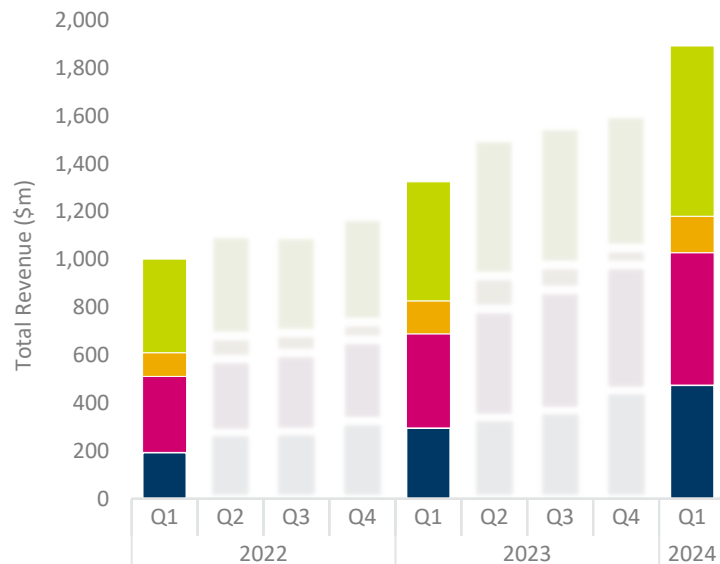
Due to rounding, the sum of a number of dollar values and percentages may not agree to totals.



BioPharmaceuticals: Cardiovascular, Renal & Metabolism

Farxiga

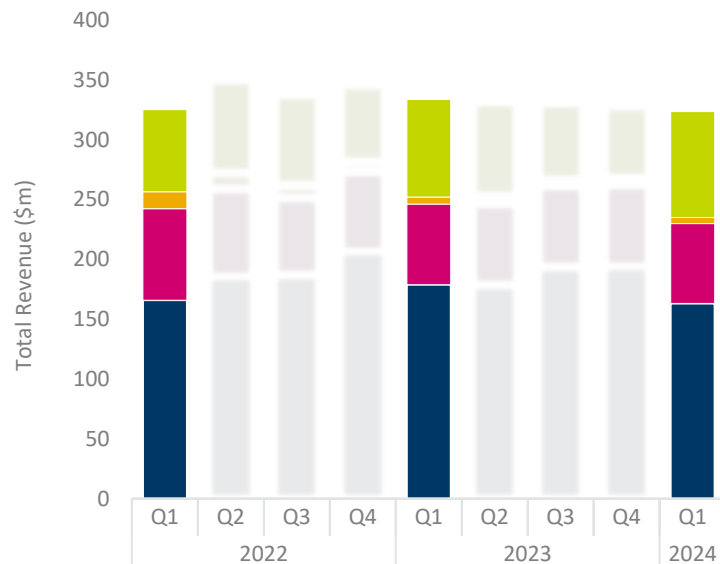
45% growth at CER to \$1,892m in Q1 2024



	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1
	2022				2023				2024
EM	391	423	410	441	498	578	579	559	711
ERoW	99	98	84	72	138	136	105	73	152
Europe	318	309	329	342	393	456	506	525	553
US	193	275	279	323	296	339	366	451	475

Brilinta

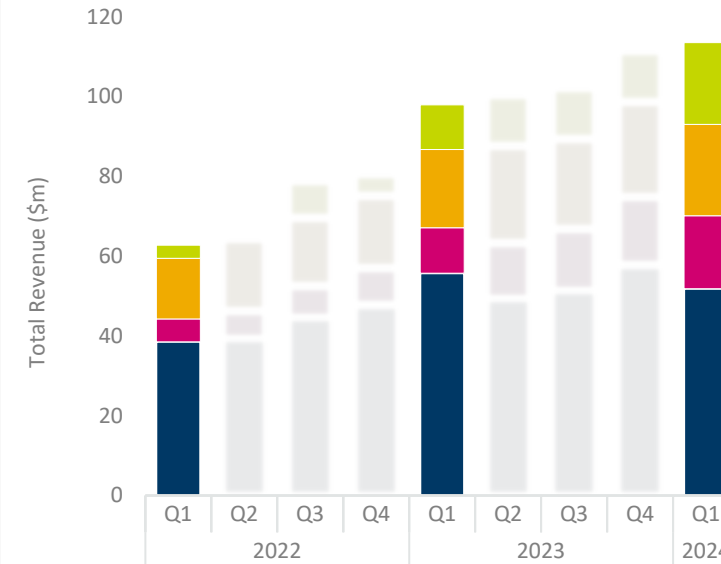
1% decrease at CER to \$323m in Q1 2024



	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1
	2022				2023				2024
EM	69	78	76	64	82	79	64	61	88
ERoW	14	14	10	8	6	6	6	6	5
Europe	76	73	65	67	67	68	68	68	67
US	166	185	187	206	179	178	193	194	163

Lokelma

19% growth at CER to \$114m in Q1 2024



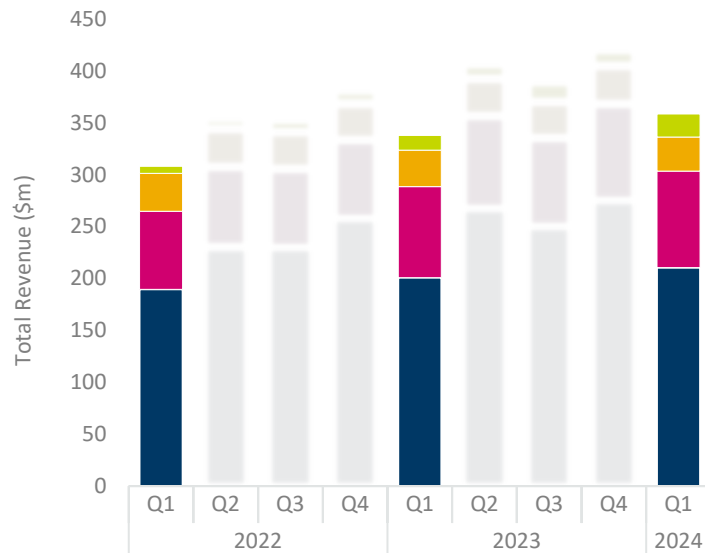
	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1
	2022				2023				2024
EM	3	2	9	6	11	13	13	13	21
ERoW	15	18	17	18	20	24	22	24	23
Europe	6	7	8	9	11	14	16	17	18
US	39	39	45	48	56	49	51	58	52



BioPharmaceuticals: Respiratory & Immunology

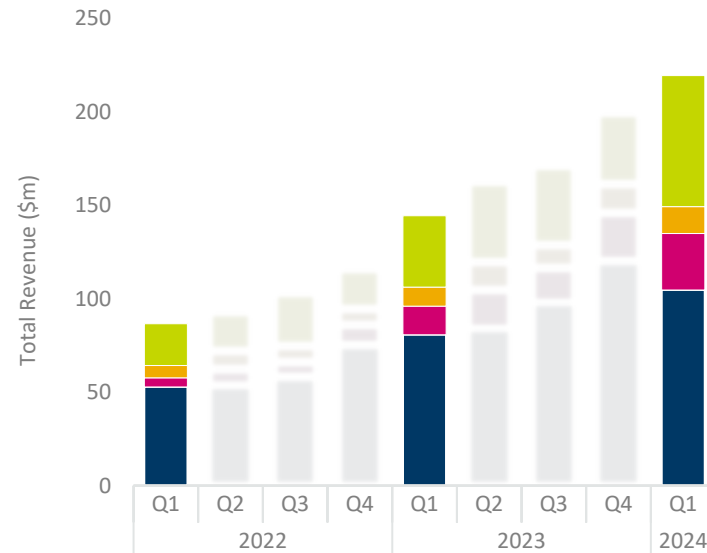
Fasenra

6% growth at CER to \$358m in Q1 2024



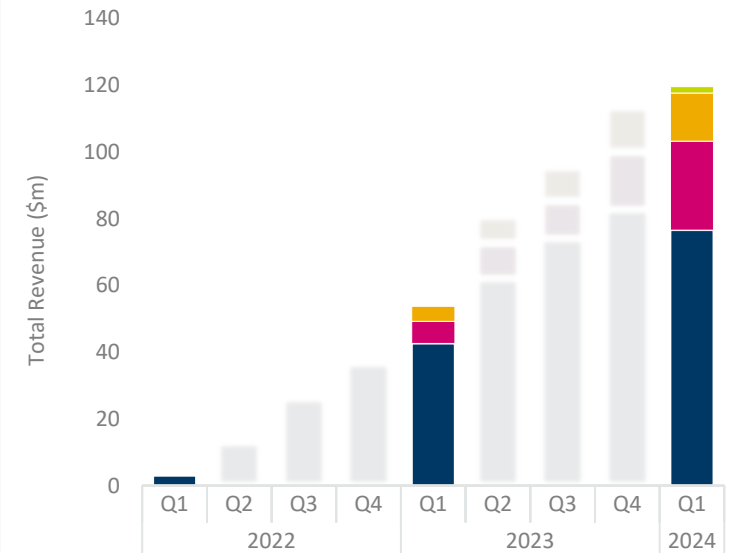
Breztri

54% growth at CER to \$219m in Q1 2024



Tezspire

>2x growth at CER to \$120m in Q1 2024



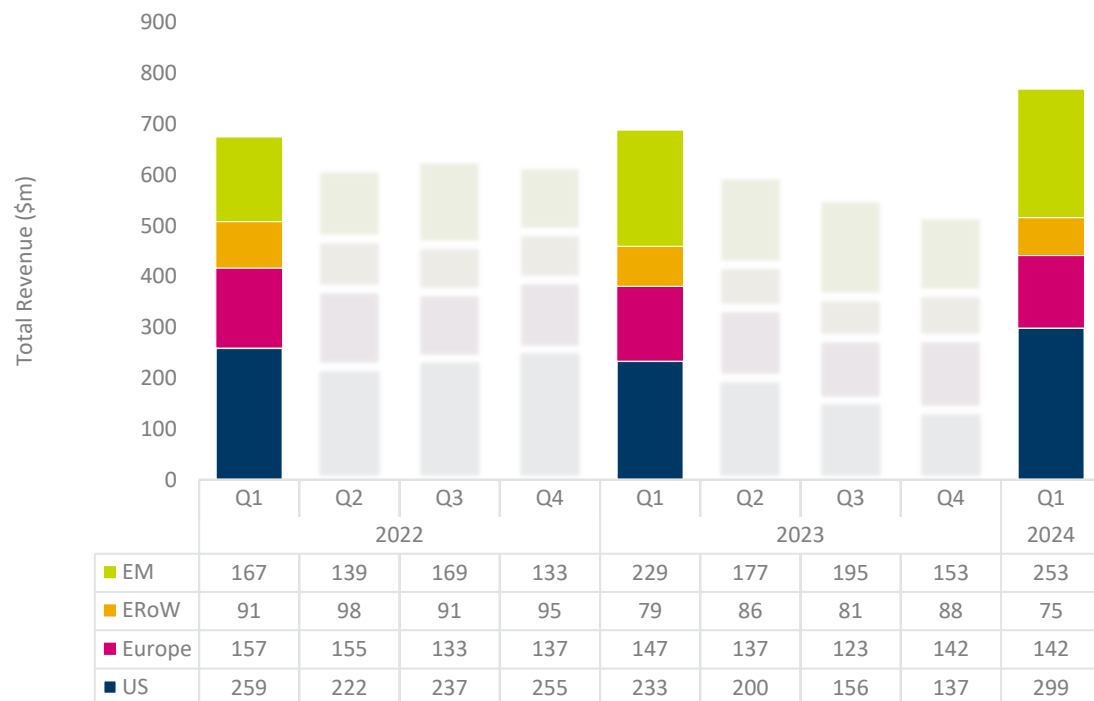
Due to rounding, the sum of a number of dollar values and percentages may not agree to totals.



BioPharmaceuticals: Respiratory & Immunology

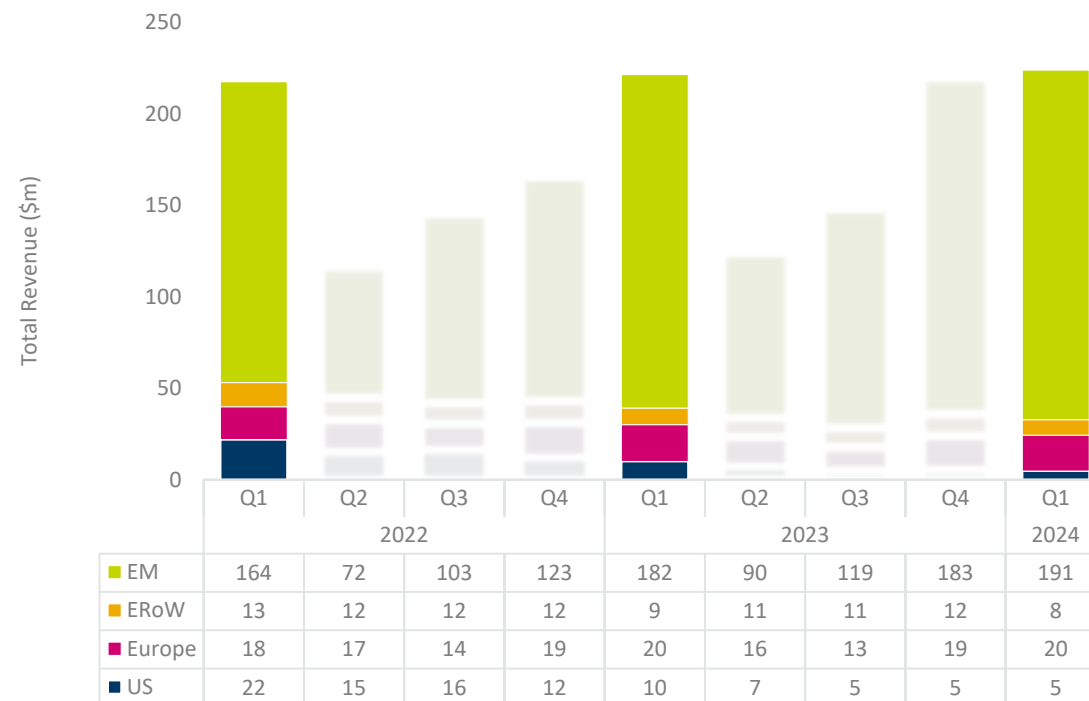
Symbicort

14% growth at CER to \$769m in Q1 2024



Pulmicort

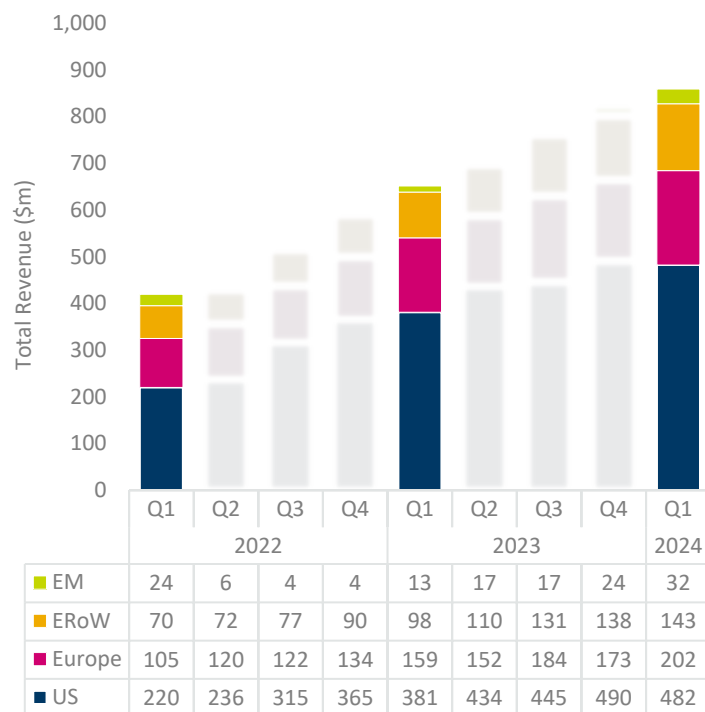
5% growth at CER to \$224m in Q1 2024



Rare Disease

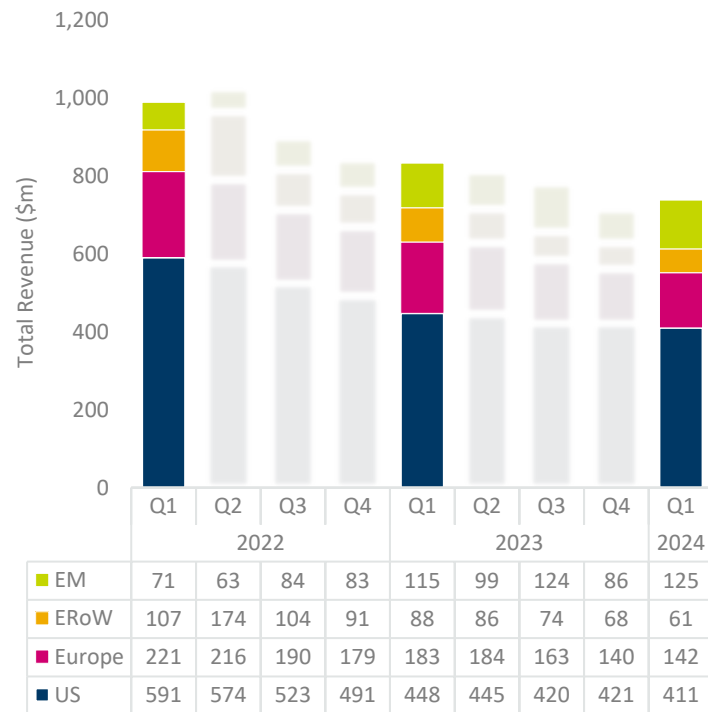
Ultomiris

34% growth at CER to \$859m in Q1 2024



Soliris

8% decrease at CER to \$739m in Q1 2024



Strensiq

21% growth at CER to \$313m in Q1 2024

