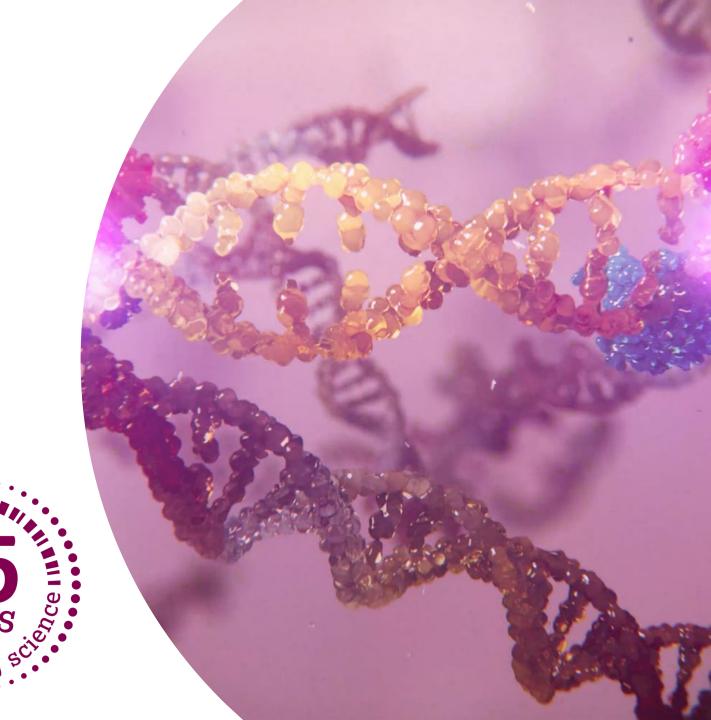


Full Year and Q4 2023 Results

Conference call and webcast for investors and analysts

08 February 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 Icosavax to complete the transactions contemplated by the merger agreement with Icosavax, including the parties' ability to satisfy the conditions to the consummation of the tender offer contemplated thereby and the other conditions set forth in the merger agreement with Icosavax; the ability of the Group and Gracell to complete the transactions contemplated by the merger agreement with Gracell, including the parties' ability to satisfy the conditions set forth in the merger agreement with Gracell; the Group's statements about the expected timetable for completing the acquisitions of Icosavax and Gracell; the Group's and Icosavax's beliefs and expectations and statements about the benefits sought to be achieved in the Group's pending acquisition of Icosavax; the Group's and Gracell's beliefs and expectations and statements about the benefits sought to be achieved in the Group's proposed acquisition of Gracell; the potential effects of the acquisition of Icosavax on both the Group and Icosavax and of the acquisition of Gracell on both the Group and Gracell; the possibility of any termination of the merger agreement with Icosavax or of the merger agreement with Gracell; the expected benefits and success of IVX-A12 and any combination product or GC012F and any combination product; the possibility that any milestone related to any contingent value right will not be achieved; 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 Icosavax will be satisfied on the expected timetable or at all or that IVX-A12 or any further vaccines using the VLP technology 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 Gracell will be satisfied on the expected timetable or at all or that GC012F will receive the necessary regulatory approvals or prove to be commercially successful if approved.



Q4 and FY 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	Sharon Barr	
Dioi narmaccaticais	EVP, BioPharmaceuticals Business	EVP, BioPharmaceuticals R&D	
Dava Diagona	Marc Dunoyer		
Rare Disease	Chief Executive Officer, Alexion		
CEO Clasina Damarka OS A	Pascal Soriot		
CEO Closing Remarks, Q&A	Chief Executive Officer		





CEO Opening Remarks

Pascal Soriot

CHIEF EXECUTIVE OFFICER



Delivered on our growth ambition

Total Revenue growth ambition to achieve >\$45bn in 2023 set in 2014



Source: Total Revenue Growth ambition as shown in 2014

Culture of science-led innovation unlocked a decade of industry-leading growth

Follow the science

Disciplined investment

Focus



Oncology

BioPharmaceuticals





Rare Disease



Poised to deliver through the next decade

Delivered on our upgraded 2023 guidance

Total Revenue to increase by mid single-digit %



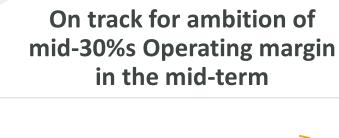
Total Revenue ex COVID-19¹ to increase by low teens %

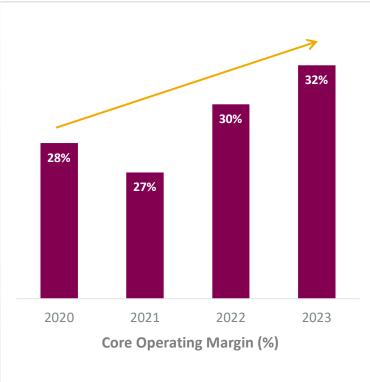
Core EPS to increase by low double-digit

to low teens %



+15%





Continued investment to drive sustainable long-term growth

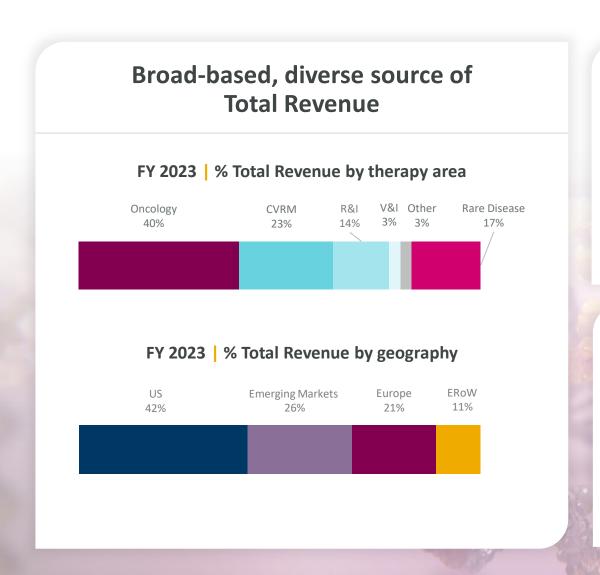
Driving near and mid-term growth across geographies and therapy areas

Building pipeline momentum

Shaping the future of medicine

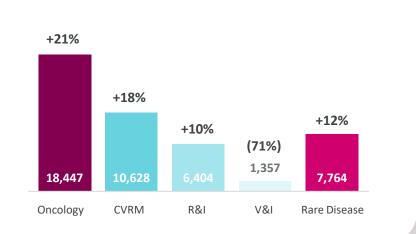


Driving strong growth across geographies and therapy areas



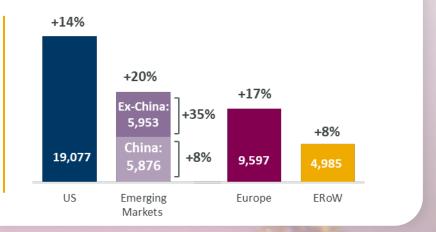


FY 2023 | Total Revenue



Growth across geographies

FY 2023 | Total Revenue ex COVID-19 ¹





Investing in new launches, near and mid-term pipeline

Phase III trials initiated across 18 medicines

>10 Phase III trials initiated with blockbuster potential

regulatory approvals in major markets

4 new medicines approved¹ and on track to deliver on ambition for at least 15 NME launches by 2030



- Approval in asthma
- First-in-class inhaler



- Approval in HR+ 2L mBC
- First-in-class AKT inhibitor



- Approval in ATTRv-PN
- ATTR-CM Phase III ongoing



- Approval as add-on in PNH
- Oral to address significant EVH



Investing in new platforms and technologies

Shaping the future of medicine





Financial Results

Aradhana Sarin

CHIEF FINANCIAL OFFICER



FY 2023 – Reported profit and loss



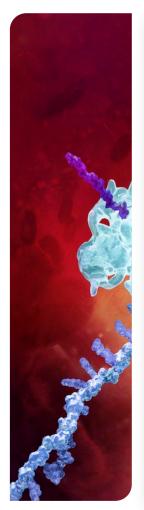
	FY 2023 \$m	CER change %	% Total Revenue	Q4 2023 \$m	CER change %	% Total Revenue
Total Revenue	45,811	6	100	12,024	8	100
- Product Sales	43,789	4	96	11,323	5	94
- Alliance Revenue	1,428	89	3	424	67	4
- Collaboration Revenue	594	(1)	1	277	74	2
Product Sales Gross Margin	81.1%	+10pp		79.6%	+6pp	
Total operating expense ¹	(30,690)	8	67	(8,589)	15	71
- R&D expense	(10,935)	13	24	(3,073)	15	26
- SG&A expense	(19,216)	6	42	(5,371)	16	45
Other operating income and expense	1,340	>2x	3	107	(42)	1
Operating profit	8,193	>2x	18	1,234	14	10
Tax rate	14%			(7%)		
Reported EPS	\$3.84	96		\$0.62	5	

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



FY 2023 – Core profit and loss



	FY 2023 \$m	CER change %	% Total Revenue	Q4 2023 \$m	CER change %	% Total Revenue
Total Revenue	45,811	6	100	12,024	8	100
- Product Sales	43,789	4	96	11,323	5	94
- Alliance Revenue	1,428	89	3	424	67	4
- Collaboration Revenue	594	(1)	1	277	74	2
Product Sales Gross Margin	81.7%	+2pp		79.8%	+2pp	
Total operating expense ¹	(24,545)	9	54	(7,093)	12	59
- R&D expense	(10,267)	9	22	(2,914)	14	24
- SG&A expense	(13,739)	9	30	(4,034)	12	34
Other operating income and expense	1,279	>2x	3	107	(15)	1
Operating profit	14,534	14	32	2,752	6	23
Tax rate	17%			10%		
Core EPS	\$7.26	15		\$1.45	7	

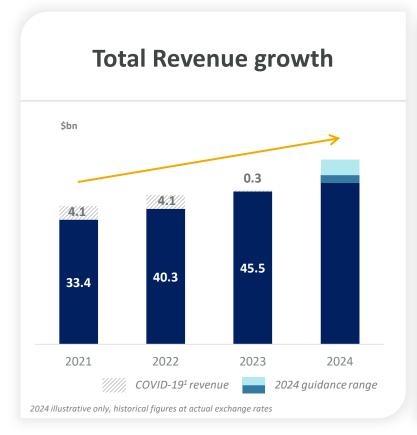
Due to rounding, the sum of a number of dollar values and percentages may not agree to totals.

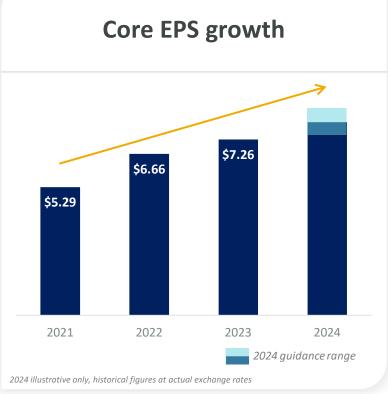




Track record and FY 2024 guidance (CER)

Underlying business momentum drives strong Total Revenue and Core EPS growth







Total Revenue

Low double-digit to low-teens percentage increase

Core EPS

Low double-digit to low-teens percentage increase

Strong Total Revenue momentum expected to continue into 2024

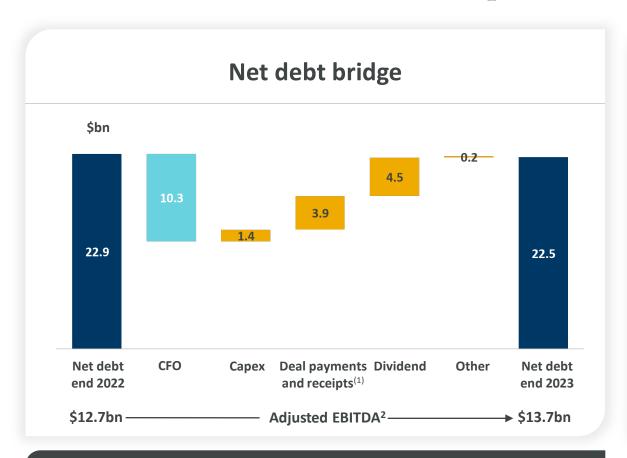
2021-2023 Core EPS CAGR of 17%

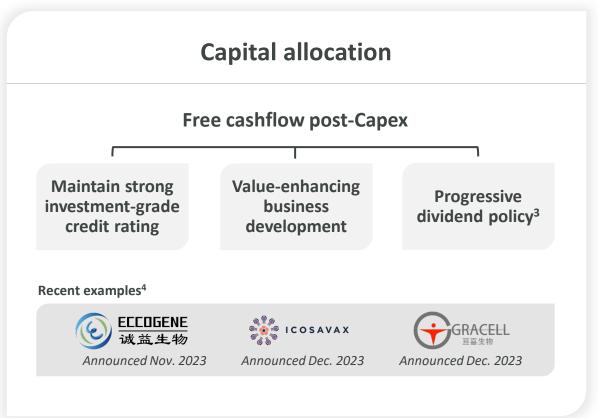
Low single-digit adverse FX impact anticipated on both Total Revenue and Core EPS²



Net debt and capital allocation

Delivered continued cash flow improvement





Net debt/Adjusted EBITDA 1.6x

Priority remains reinvesting in the business

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 acquisition of subsidiaries, net of cash acquired. 2. Rolling 12m EBITDA adding back the impact of unwind of inventory fair value uplift recognised on acquisition of Alexion of \$114m (FY 2022: \$3,484m). 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 or increasing dividend per share in US dollar terms. 4. Icosavax and Gracell acquisitions remains subject to customary closing conditions; all clinical development plans mentioned herein subject to deal closure.



Appendix: Glossary.

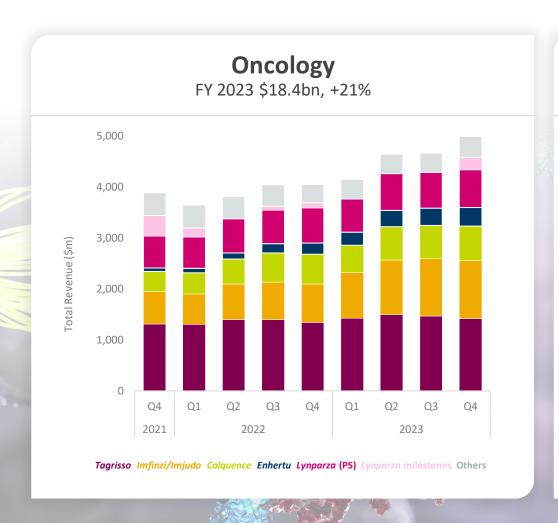
Dave Fredrickson
ONCOLOGY BUSINESS

Susan Galbraith
ONCOLOGY R&D



Oncology – FY and Q4 2023

Total Revenue +21% in FY 2023 fuelled by strong global demand growth

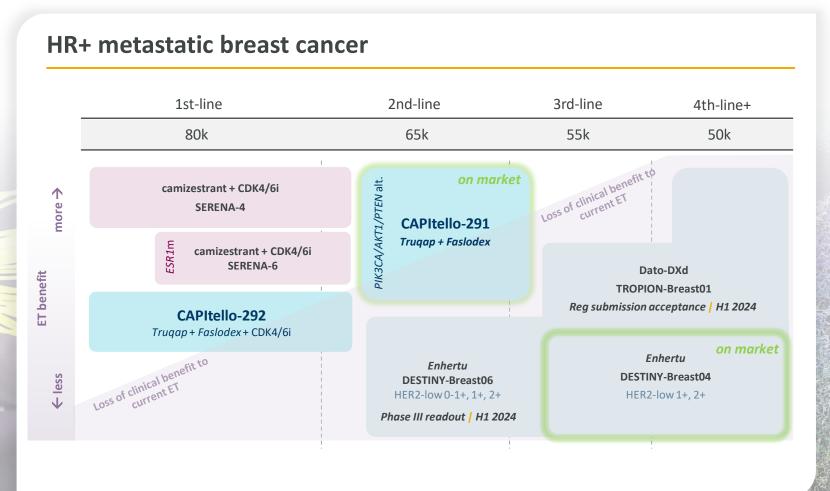


Q4 2023: key dynamics

- *Tagrisso* +6%, demand growth in US and EU, offset by JP pricing, ERoW rebate reclassification and hospital ordering dynamic in CN
- Lynparza PS +8%, growth supported by continued PARPi leadership
- Imfinzi/Imjudo +52%, fueled by BTC (TOPAZ-1), HCC (HIMALAYA)
- Calquence +14%, BTKi NPS leadership in CLL across US and EU
- *Enhertu* +68%, clear standard-of-care in HER2+ (DB03) and HER2-low (DB04), sequential NPS growth in HER2+ in US and DE
- New indications: US (*Truqap* HR+, HER2- mBC), CN (*Imfinzi* BTC)
- Regulatory/payer: US (Enhertu HER2+ tumour agnostic Priority Review), CN (adjuvant (ADAURA) NRDL inclusion)



Oncology – Truqap launch reinforces breast cancer leadership



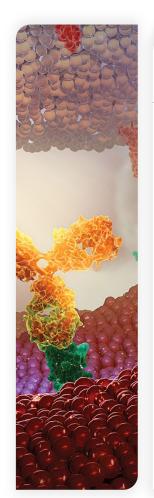
Trugap poised to redefine 2L HR+, potential SoC for altered tumours ~28k patients diagnosed, treated with 21 HR+ disease in the US¹ 85% of patients received ET in 1L Up to 50% with alterations in PIK3CA, AKT1 or PTEN

Strong US launch momentum, filings underway in EU, JP



Oncology – R&D highlights

Gracell acquisition furthers our position in CAR-T and haematology





Gracell cell therapy manufacturing platform

Shorter manufacturing time

Increased manufacturing capacity

Lower cell dose required

May improve safety

Enhanced T-cell fitness

Potential to improve outcomes

GC012F

potential best-in-class BCMA/CD19 dual-targeted CAR-T

Differentiated clinical activity in newly diagnosed multiple myeloma¹



100%

95%+

ORR at all dose levels

MRD negativity 6-12 mo. after infusion

Strengthening haematology portfolio and pipeline

Calquence | AZD0486 | AZD0305 | GC012F | AZD9829 | AZD3470



Ruud Dobber

BIOPHARMACEUTICALS BUSINESS

Sharon Barr

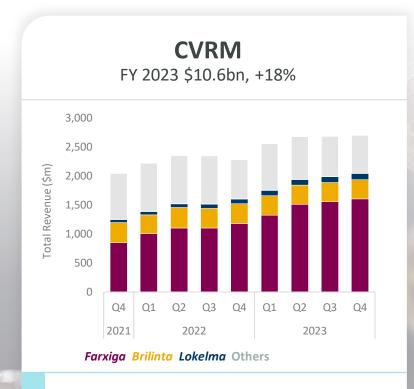
BIOPHARMACEUTICALS R&D



BioPharmaceuticals

BioPharmaceuticals – FY and Q4 2023

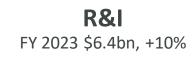
Double-digit growth from CVRM and R&I, strong Beyfortus launch in V&I

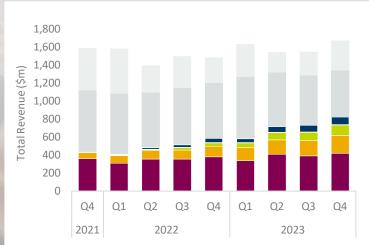


Farxiga +35%, demand growth outpacing SGLT2i

Lokelma +38%, K+ Binder leadership in US

roxadustat +27%, increased demand





Fasenra Breztri Tezspire Saphnelo Symbicort Others



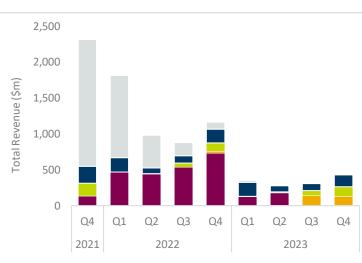
24 dynamics



Breztri +72%, global market share gains

Tezspire >2x, strong launches continue

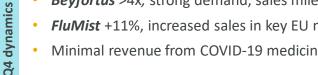




COVID-19 mAbs¹ Beyfortus FluMist Synagis Vaxzevria

Beyfortus >4x, strong demand, sales milestone FluMist +11%, increased sales in key EU markets

Minimal revenue from COVID-19 medicines





Appendix: Glossary.

Q4 dynamics

BioPharmaceuticals

Pipeline success leads to multiple launches of differentiated medicines

CVRM



R&I



V&I



First-and-only self-admin auto-injector for the treatment of ATTRV-PN

First-and-only rescue inhaler that also reduces risk of asthma exacerbations

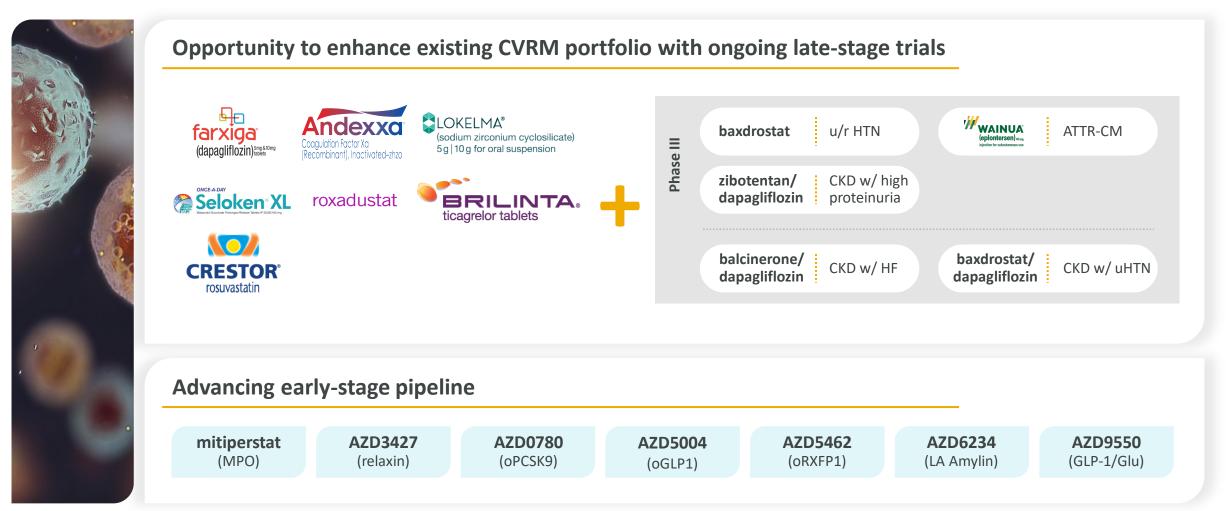
First-and-only RSV mAb approved for the broad infant population



CEO Opening Remarks Financial Results Oncology BioPharmaceuticals Rare Disease CEO Closing Remarks

BioPharmaceuticals

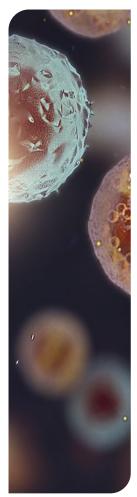
Future expansion of CVRM portfolio with numerous ongoing late-stage, early-stage trials

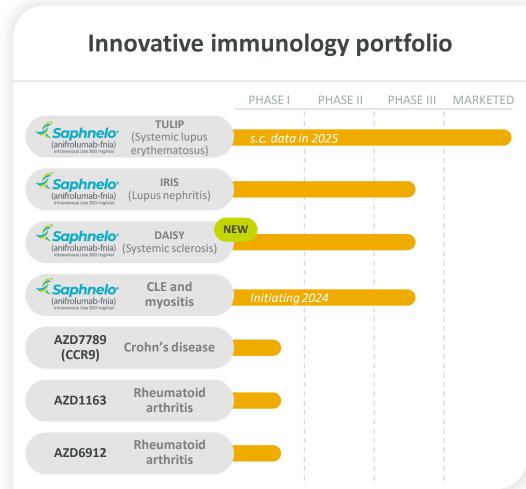


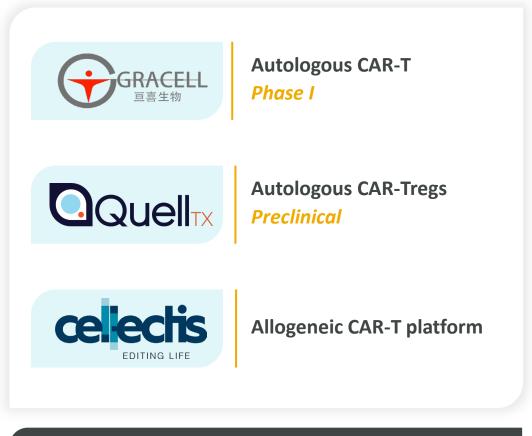


BioPharmaceuticals – R&D highlights

Accelerating our ambition in immune-mediated diseases







Investing in transformational cell therapies with curative potential



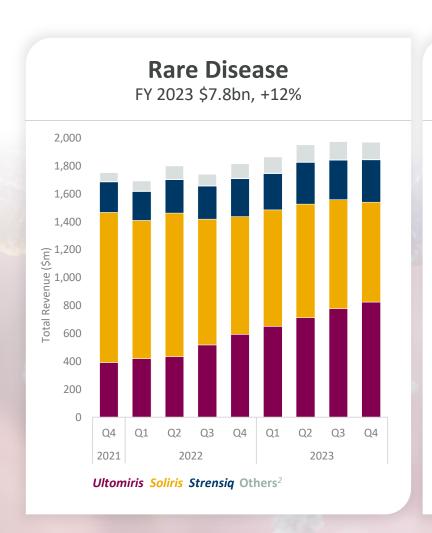
Marc Dunoyer

CHIEF EXECUTIVE OFFICER, ALEXION



Rare Disease – FY and Q4 2023

Total Revenue +12% in FY 2023 driven by neurology and patient demand

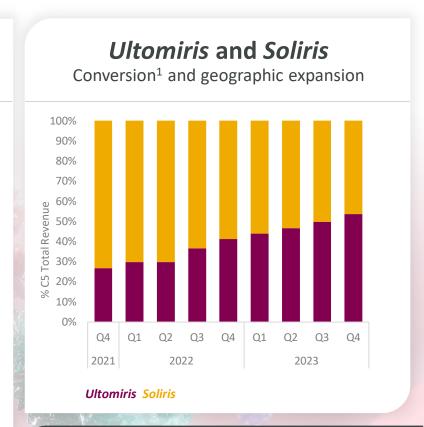


Q4 2023: key dynamics

Sustainable, durable growth of C5 Franchise

- Ultomiris, +38% driven by neurology expansion
- Soliris, (13%) due to conversion, partly offset by Emerging Markets growth

Strensiq, +13% and **Koselugo**, +48% driven by continued patient demand



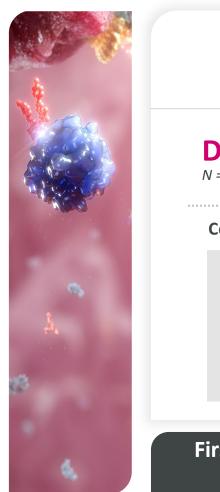
Sustained C5 leadership with durable *Ultomiris* and *Soliris* growth

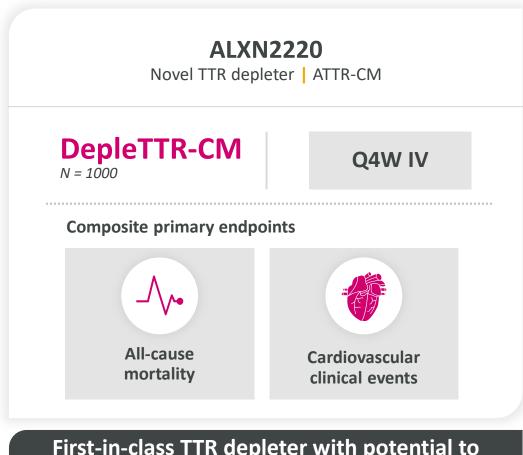
All growth rates at CFF

1. Patients converting their treatment from Soliris to Ultomiris 2. Includes Kanuma and Koselugo.

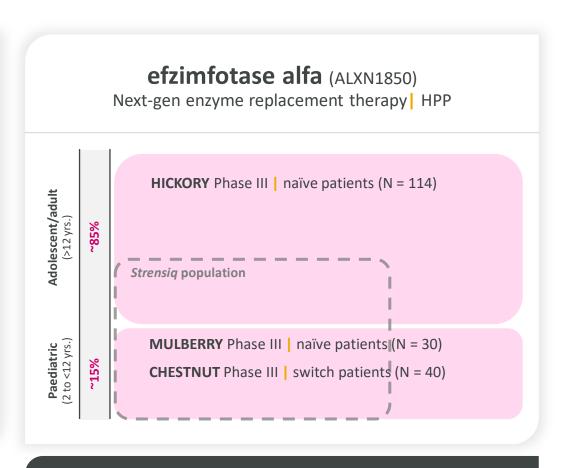
Rare Disease – R&D

New Phase III trial starts with blockbuster potential





First-in-class TTR depleter with potential to reverse the course of disease



3x Strensiq addressable population¹



Appendix: Glossary.

CEO Closing Remarks

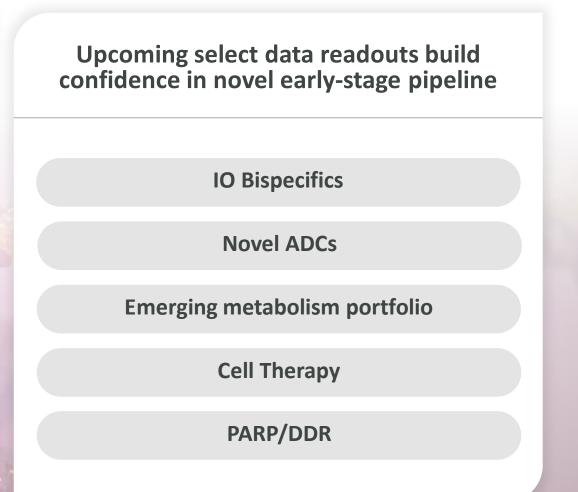
Pascal Soriot

CHIEF EXECUTIVE OFFICER



Industry-leading pipeline, significant catalysts in 2024

Strong near-term Phase III catalyst volume select opportunities include: **LAURA** EGFRm NSCLC (unresectable Stg. III) **Tagrisso DESTINY-Breast06** HER2-low breast cancer (2L) Enhertu **TROPION-Breast02 TNBC** H₂ 2024 (locally rec. inop./met.) Dato-DXd **WAYPOINT Chronic Rhinosinusitis** H₂ 2024 with Nasal Polyps *Tezspire* **EMERALD-2** Liver cancer (adjuvant) *Imfinzi*





EO Opening Remarks Financial Results Oncology BioPharmaceuticals Rare Disease CEO Closing Remarks

Investor Day to feature strategic progress and pipeline

Roadmap to delivering industry-leading growth through 2030





21 May 2024

The Discovery Centre (DISC)
Cambridge, UK

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 BarrEXECUTIVE VICE PRESIDENT,
BIOPHARMACEUTICALS R&D



Ruud Dobber
EXECUTIVE VICE PRESIDENT,
BIOPHARMACEUTICALS
BUSINESS



Iskra Reic
EXECUTIVE VICE PRESIDENT,
VACCINES AND IMMUNE
THERAPIES

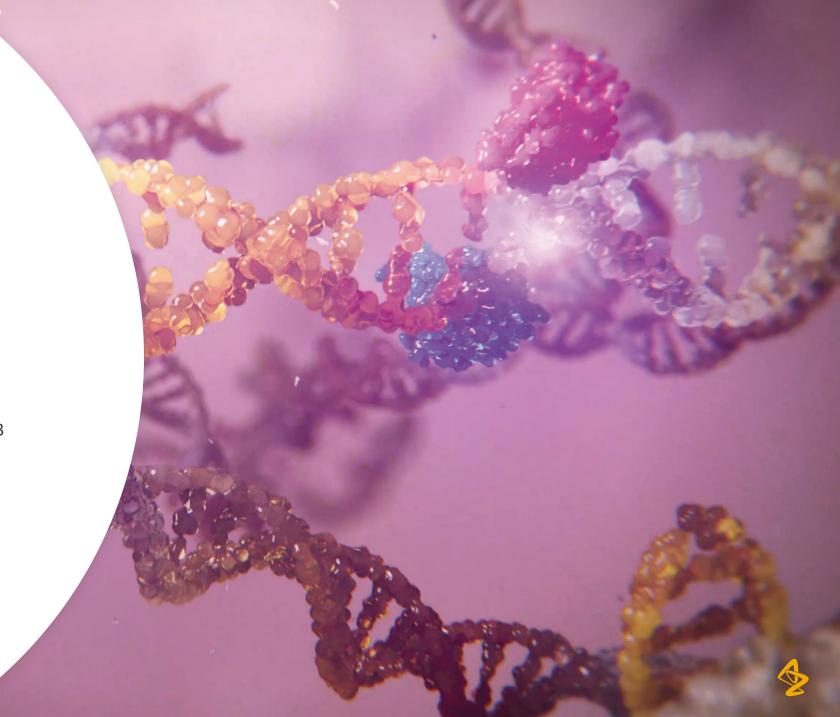


Leon Wang
EXECUTIVE VICE PRESIDENT,
INTERNATIONAL





- Glossary
- ESG summary of sustainability progress in 2023
- Oncology tumour maps
- Emerging Markets performance
- Key medicines performance by therapy area



Glossary – abbreviations (1 of 2)

1L = first line

2L = second-line

ADCs = antibody drug conjugates

AI = aromatase inhibitor

AKT1 = Ak strain transforming 1

AKTi = Ak strain transforming inhibitor

ATTR-CM = transthyretin amyloid cardiomyopathy

ATTRv-PN = polyneuropathy of hereditary transthyretin-mediated amyloidosis

BTC = biliary tract cancer

BTKi = bruton tyrosine kinase inhibitor

CAGR = compound annual growth rate

CapEx = capital expenditure

CAR-T = chimeric antigen receptor

CAR-Tregs = chimeric antigen receptor regulator

CD19/BCMA = cluster of differentiation 19/B cell maturation antigen

CDK4/6i = cyclin-dependent kinase 4 and 6 inhibitor

CER = constant exchange rate

CFO = cash flow from operations

CKD = chronic kidney disease

CLE = cutaneous lupus erythematous

CLL = chronic lymphocytic leukemia

CN = China

CRSwNP = chronic rhinosinusitis with nasal polyps

CRT = chemoradiotherapy

CTx = chemotherapy

CVRM = cardiovascular, renal and metabolism

Dato-DXd = datopotamab deruxtecan

DE = Germany

DHP = docetaxel, trastuzumab and pertuzumab

EBITDA = earnings before interest, depreciation and amortisation

EGFRm = epidermal growth factor receptor mutation

EM = emerging markets

ERoW = established rest of world

ERT = estrogen replacement therapy

Est epi (G7) = estimated epidemiology across G7 (US, EU5, JP)

ET = endocrine therapy

EU = Europe

EVH = extravascular haemolysis

FX = foreign exchange

GLP1/Glu = glucagon like peptide 1/glucagon

HER2m = human epidermal growth factor 2 mutated

HER2- = human epidermal growth factor receptor 2 negative

HER2+ = human epidermal growth factor receptor 2 positive

HER2-low = human epidermal growth factor receptor 2 low

HF = heart failure

HLR = high level results

HPP = hypophosphatasia

HR+ = hormone receptor positive

HSR = huge seller repricing

IHC = immunohistochemistry

i.v. = intravenous

IO = immuno-oncology

JP = Japan

LA = long acting

mAb = monoclonal antibody

mBC = metastatic breast cancer

MPO = myeloperoxidase

MRD = minimal residual disease

NME = new molecular entity

NPS = new patient share

Glossary – abbreviations (2 of 2)

NRDL = national reimbursement drug list

NSCLC = non-small cell lung cancer

NST = neoadjuvant systemic treatment

oGLP1 = oral glucagon-like receptor peptide 1

oPCSK9 = oral protein convertase subtilisin/kexin type 9

ORR = overall response rate

oRXFP1 = oral relaxin family peptide receptor 1

PARPi = poly(ADP-ribose) polymerase-1

PD1 = Programmed cell death protein 1

PD-L1 = Programmed cell death ligand 1

PIK3CA = phosphatidylinositol-4,5-biphosphate 3-kinase catalytic subunit

PNH = paroxysmal nocturnal haemoglobinuria

PS = product sales

PTEN = phosphatase and TENsin homolog deleted on chromosome 10

Q4W = every 4 weeks

QoQ = quarter on quarter

R&D = research and development

R&I = respiratory and immunology

RSV = respiratory syncytial

SBRT = Stereotactic brain radiotherapy

SG&A = sales, general and administrative

SoC = standard of care

Stg I/II/III = Stage 1/2/3

TIGIT = T-cell immunoreceptor with immunoglobulin and ITIM domains

TKI = tyrosine kinase inhibitor

TIM-3 = T-cell immunoglobulin and mucin domain-containing protein

TNBC = triple negative breast cancer

TTR = transthyretin

u/r HTN = uncontrolled or treatment resistant hypertension

V&I = vaccines and immune therapies

Continued leadership in Sustainability

Accelerating bold, scalable action through collaboration





Demonstrating our commitment to nature by bringing nature more prominently into decision-making



Early adopter

Accelerating the transition to net zero health systems through an industry-first

Unlocking green power in China together with four global healthcare leaders

comparable to taking 25,000 cars off the road

Collaboration platforms include:

agreement



WØRLD ECONOMIC FORUM

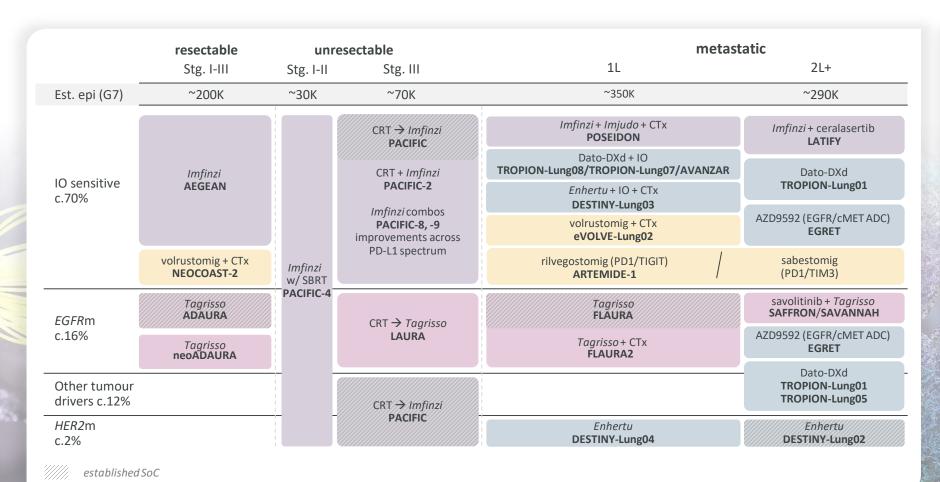


Innovating for the health of people, society and the planet



AstraZeneca in Lung Cancer

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



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

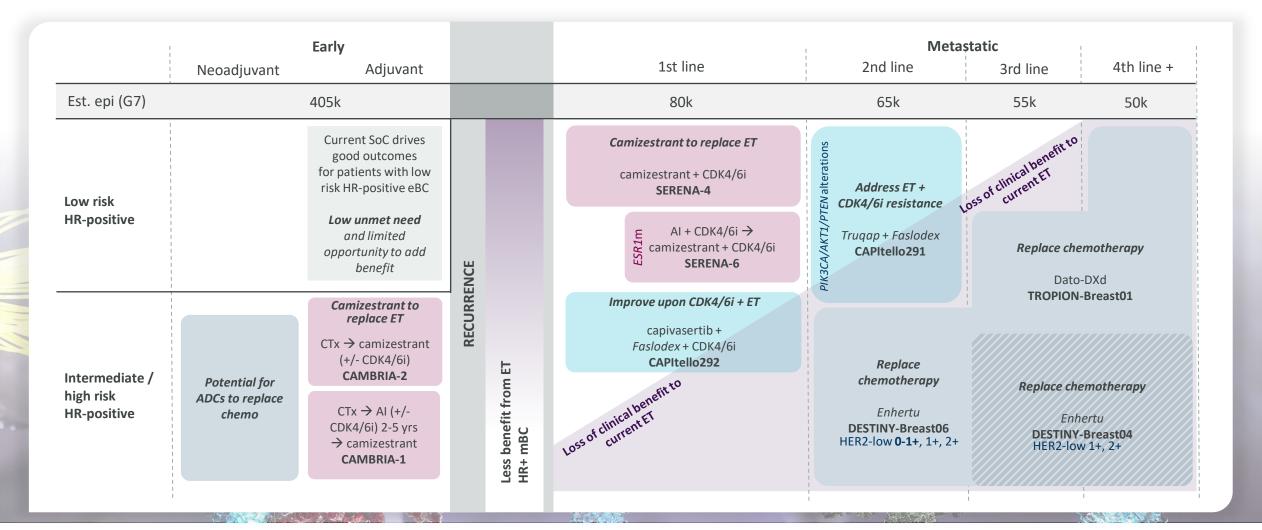
Ambition to eliminate breast cancer as a cause of death

established SoC	Neoadjuvant	Early Adjuvant		1st line	Metastatic 2nd line	3rd line	4th line +
Est. epi (G7)	54			125k	90k	65k	55k
HER2-positive 15-20%	Enhertu ± THP DESTINY-Breast11	NST→ residual disease → Enhertu DESTINY-Breast05		Enhertu ± pertuzumab DESTINY-Breast09	Enhertu DESTINY-Breast03	Enhertu DESTINY-Breast02	
HR-positive 65-75% HER2-low 1+, 2+ 60%		ৰ Good outcomes with current SoC		camizestrant + CDK4/6i SERENA-4	HR3CA/ AKT1/ Faslodex Faslodex CAPItello291	Dato-DXd TROPION-Breast01	
		CTx → camizestrant (± CDK4/6i) CAMBRIA-2	RENCE	E AI + CDK4/6i → camizestrant + CDK4/6i SERENA-6	Enhertu	: 	
	CTx → AI (± CDK4/6i) 2-5 yrs → camizestrant CAMBRIA-1	RECURRENCE	Truqap + Faslodex + CDK4/6i CAPItello292	DESTINY-Breast06 HER2-low IHC 0-1+, 1+, 2+	Enhertu DESTINY-Breast04 HER2-low IHC 1+, 2+		
TNBC 10-15%	Dato-DXd+	NST		Truqap + paclitaxel CAPItello290	HER2- Low		
 HER2-low 1+, 2+	Imfinzi → residual disease TROPION- → Dato-DXd ± Imfinzi			PD-L1+ Dato-DXd + <i>Imfinzi</i> 40% TROPION-Breast05			
35%	Breast04	TROPION-Breast03		PD-L1- Dato-DXd 60% TROPION-Breast02			
gBRCAm 5% of HR-positive 15% of TNBC	 	CTx → <i>Lynparza</i> OlympiA			Lynparza OlympiAD		



AstraZeneca in Breast Cancer

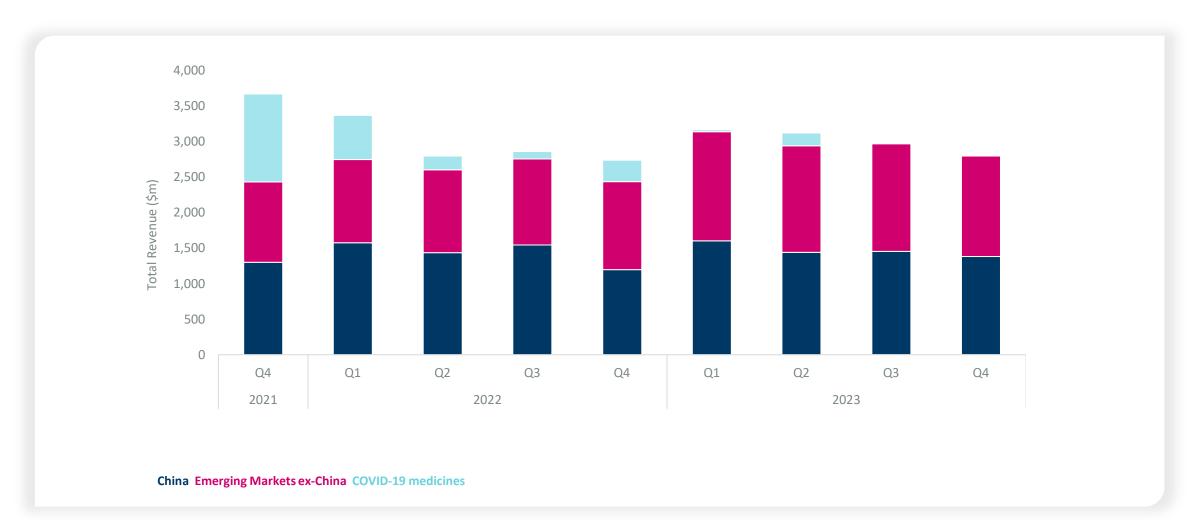
Ambition to eliminate breast cancer as a cause of death



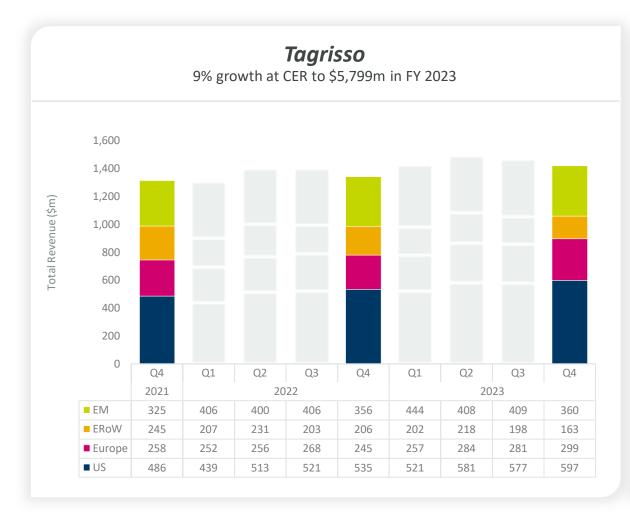


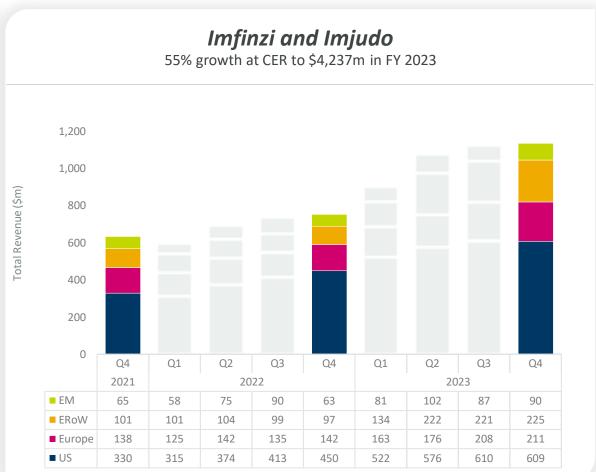
Emerging Markets – FY 2023

Total Revenue +9% at CER to \$12.0bn, +20% at CER ex-COVID-19 medicines

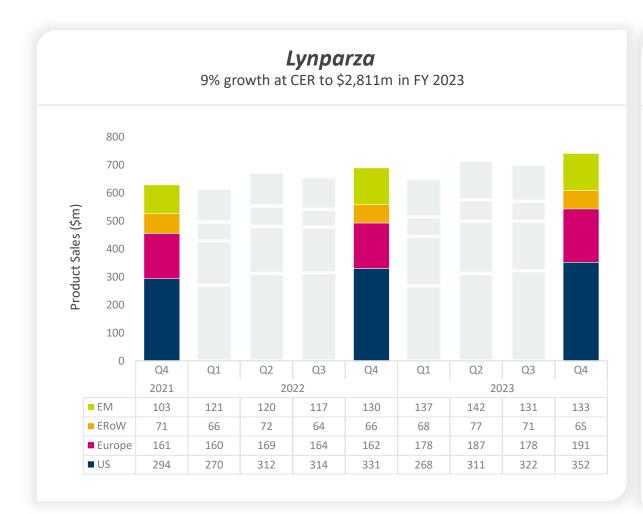


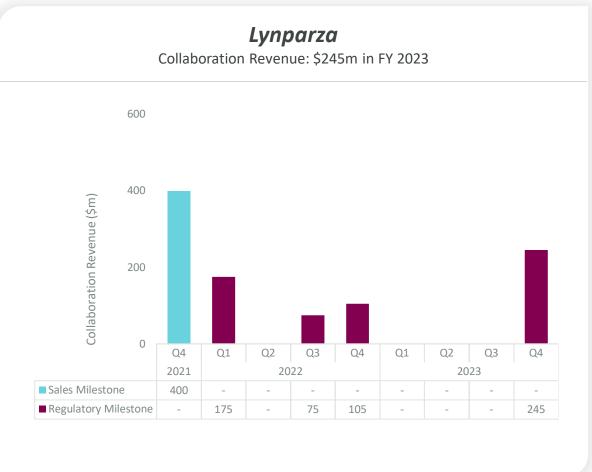




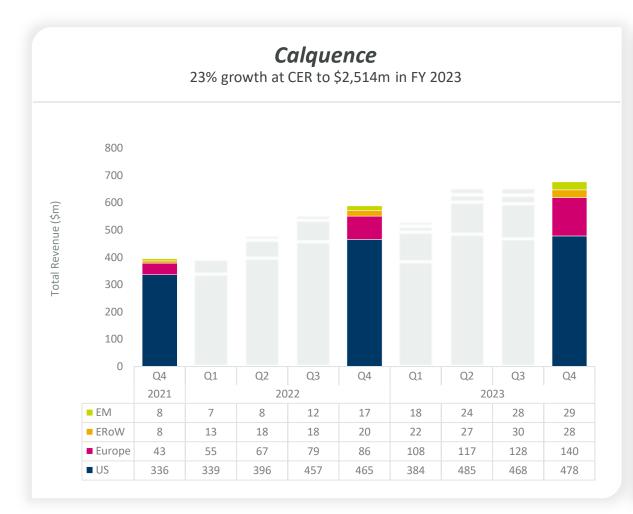


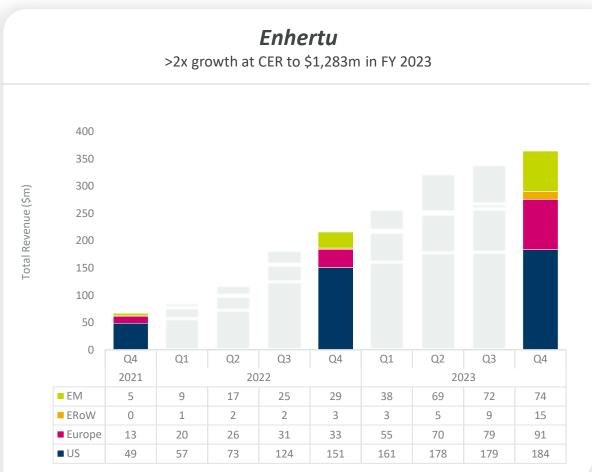










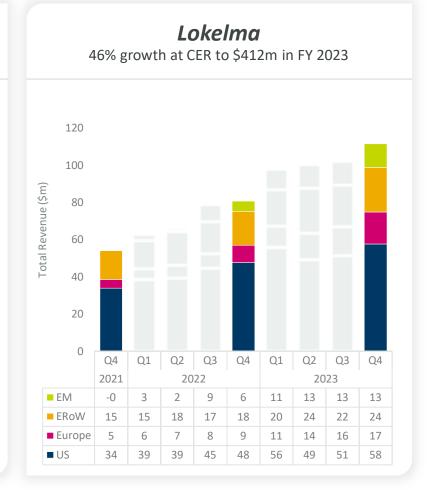




BioPharmaceuticals: Cardiovascular, Renal & Metabolism





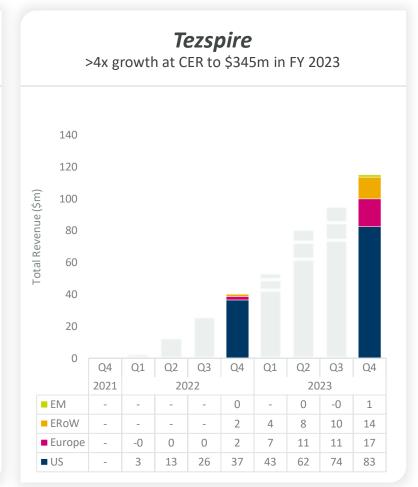




BioPharmaceuticals: Respiratory & Immunology

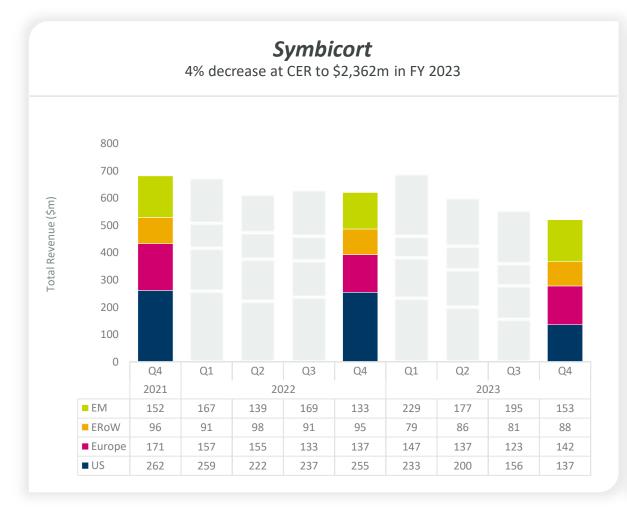


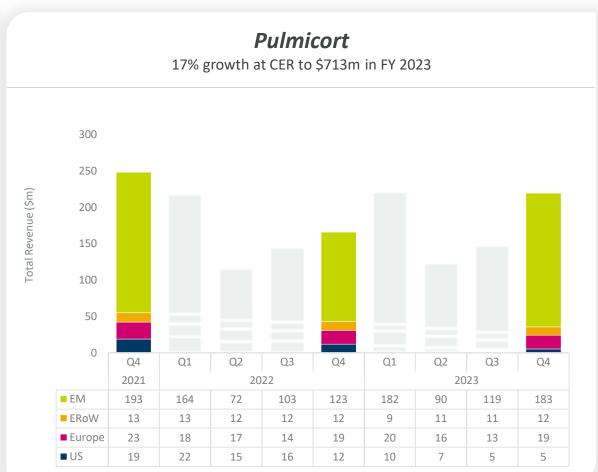






BioPharmaceuticals: Respiratory & Immunology



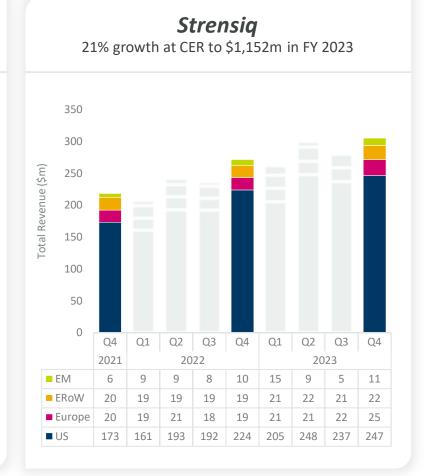




Rare Disease









Appendix: Glossary.