

Q1 2024 Results

Conference call and webcast for investors and analysts



Cautionary statement regarding forward-looking statements

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A number of adjusted measures are used to report the performance of our business, which are non-IFRS measures. These measures are defined and reconciliations to the nearest IFRS measure are available in the Group's Q1 2024 Results and the Group's Annual Report on Form 20-F for FY 2023.

All expectations, guidance and outlooks regarding future performance and the dividend should be read together with the section "Guidance and outlooks, assumptions and cautionary statements on page 49 of our stock exchange announcement of GSK's Q1 2024 Results, the section "Assumptions and basis of preparation related to 2024 guidance" in the Appendix of this presentation and the statements on page 317 of GSK's Annual Report on Form 20-F for FY 2023.



Agenda

Strong start to 2024

Emma Walmsley

Performance: growth drivers

Luke Miels and Deborah Waterhouse

Q1 2024 performance and 2024 guidance

Julie Brown

Q&A

Emma Walmsley, Tony Wood, Luke Miels, Deborah Waterhouse, Julie Brown, and David Redfern



Strong start to 2024 with continued pipeline progress

Emma Walmsley, Chief Executive Officer



Strong start to 2024

Delivered 13%¹ sales growth, 35%¹ core operating profit growth

Sales growth across portfolio:

- Vaccines +22%¹
- Specialty Medicines +19%¹
- General Medicines +1%

Cash generated from operations exceeded £1 billion, with free cash flow of £0.3 billion

Q1 2024 performance

Sales

£7.4bn, +10%

+13%

Core EPS

43.1p, +28%

+37%

Core operating profit

£2.4bn, +27%

+35%

Dividend per

share

15p

Full-year 2024 guidance¹: upgraded

Sales growth: 5-7% (towards upper part of the range)

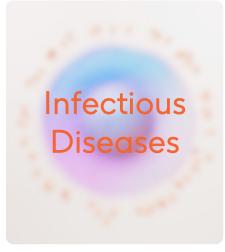
Core operating profit growth: 9-11%

Core EPS growth: 8-10%



Strong pipeline progress

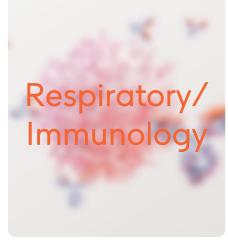
R&D priorities focused across four therapy areas



- gepotidacin: positive EAGLE-1 phase III data
- mRNA: positive influenza phase II data
- MenABCWY regulatory acceptance (US)
- bepirovirsen Fast Track designation (US)
- Arexvy: Priority Review for adults 50-59 (US)
- Shingrix: regulatory submission accepted for at-risk adults 18+ (CN)



- Cabenuva: positive LATITUDE phase III data; data presented at CROI
- ultra long-acting cabotegravir positive phase I data



Aiolos Bio acquisition completed



- Jemperli RUBY Part 1 & Part 2 data presented at SGO
- Jemperli US FDA Priority Review¹
- Blenrep: DREAMM-7 data presented at ASCO virtual; positive DREAMM 8 phase III data



Delivering health impact sustainably

For health impact, shareholder returns and thriving people

Six priority areas to build trust



Access



Environment



Product governance



Global health and health security



Diversity, equity and inclusion



Ethical standards

Key highlights

Access

- Announced \$35 per month cap on eligible US patient out-of-pocket costs for entire portfolio of asthma and COPD inhalers.
- Announced resourcing of up to \$2 million in 2024 in COiMMUNITY Initiative grants to help achieve higher adult vaccination rates and health equity in the US

Global health and health security

- Announced positive headline results from EAGLE-1 phase III trial for gepotidacin in gonorrhoea, a bacteria for which resistance to existing treatments is rising
- GSK-developed M72/AS01E tuberculosis (TB) vaccine candidate has entered phase III trials, sponsored by The Bill & Melinda Gates Medical Research Institute and Wellcome Foundation. Potentially the first new TB vaccine in over 100 years

Environment

- Action on sustainability recognised by CDP, scoring GSK an A- for tackling Climate Change, an A- for Water Security and Bs for Forests
- Announced long-term investment over next 15 years in Climate Asset Management's Nature Based Carbon Fund, which aims to invest in nature projects in developing economies



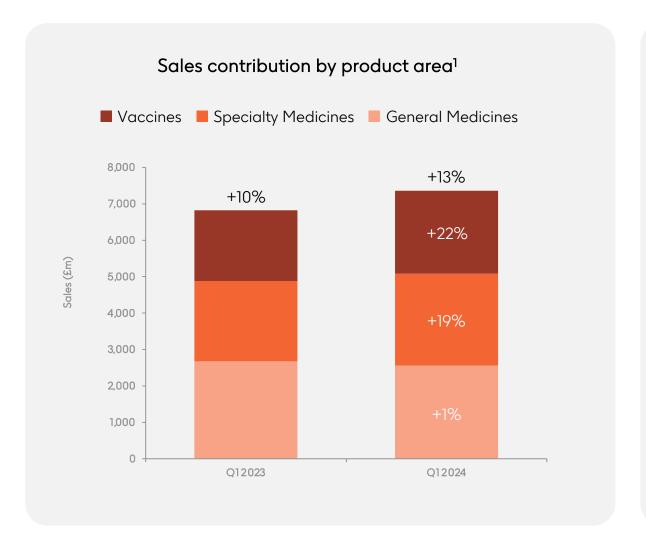
Performance: growth drivers

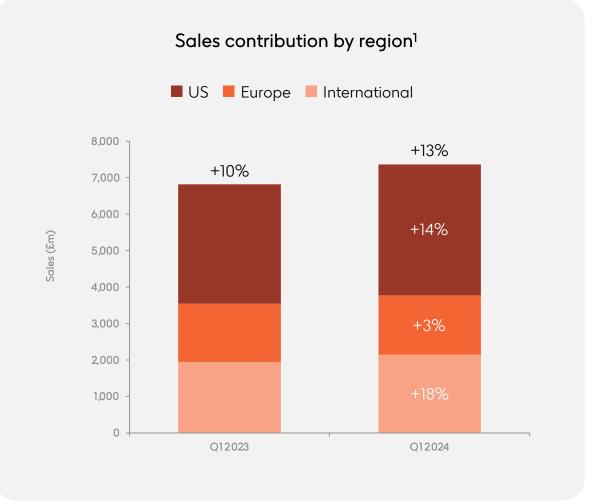
Luke Miels, Chief Commercial Officer

Deborah Waterhouse, CEO, ViiV Healthcare and President, Global Health



Strong start to 2024 with growth in all product areas and regions

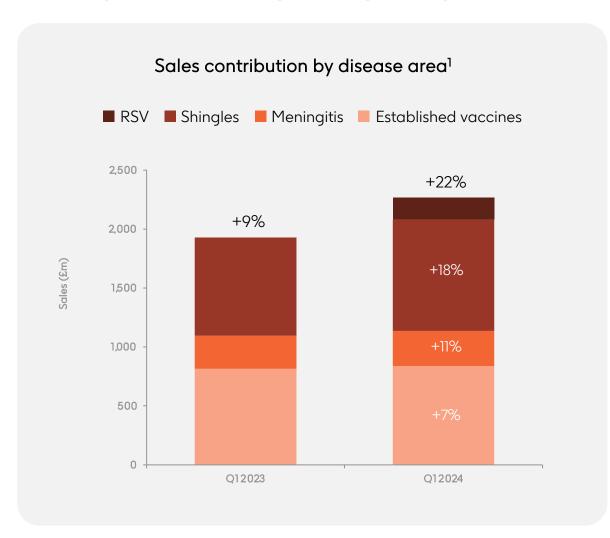






Vaccines: +22%¹ led by *Arexvy* and record sales of *Shingrix*

2024 guidance: high-single digit to low-double digit % growth¹



RSV (Arexvy) £182m²

Leading market share with 2/3 retail prescriptions

Shingles (Shingrix) +18%

- Growth driven by public funding expansion and early supply to Zhifei
- Launched in 39 markets ex-US, majority <5% penetration
- 37% of US adults recommended to receive Shingrix now vaccinated
- >£4bn in peak year sales by 2026
- 82% vaccine efficacy in adults ≥50 at year 11

Meningitis +11%

- Bexsero +3% driven by Australia performance and launch in Vietnam
- Menveo +41% driven by Brazil performance and phasing
- Combined meningitis portfolio to reach ~£2bn in peak year sales
- MenABCWY file acceptance by FDA

Influenza +8%

Established vaccines +7%



Arexvy performance

>1.3m people vaccinated with *Arexvy* in Q1 2024¹, >7m people since launch

International rollout

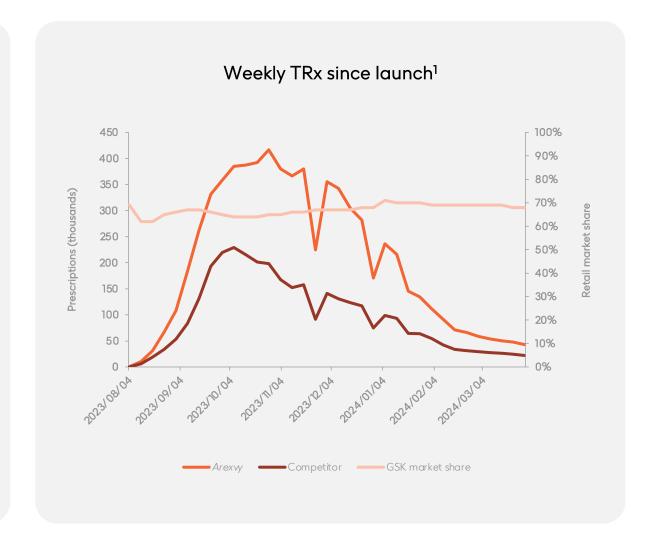
- 68% market share in retail at the end of Q1 2024¹
- Majority of 2024 sales to be in the US, weighted to H2
- Launched in 24 markets to date
- >£3bn in global peak year sales potential

Competitive profile

- Label expansion file accepted in at risk individuals aged 50-59
- PDUFA on 7 June 2024

Season 3 duration of protection and revaccination

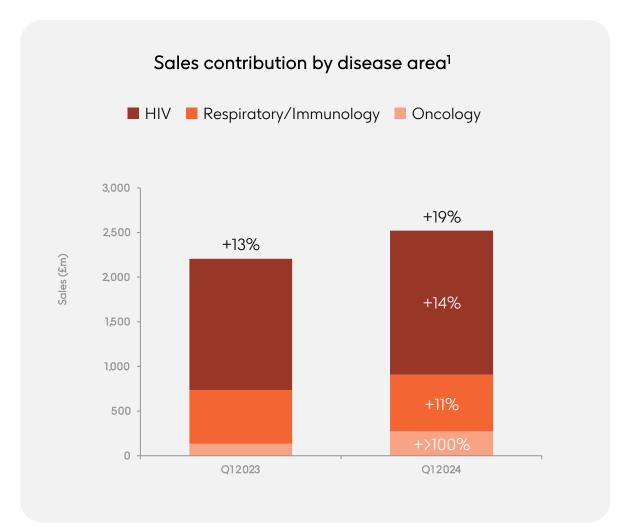
- Vaccine efficacy and immunogenicity data due H2 2024
- Revaccination decision not expected before 2025





Specialty Medicines: +19%1 with strong growth in all therapy areas

2024 guidance: low-double digit % growth¹



HIV +14%

Momentum in long-acting innovation

Respiratory/Immunology +11%

- Nucala +13% reflecting high patient demand for treatments addressing severe eosinophilic asthma, CRSwNP² and EGPA³; COPD⁴ data readout in H2 2024
- Benlysta +8% representing good underlying demand, with biopenetration and volume uptake in International

Oncology +>100%

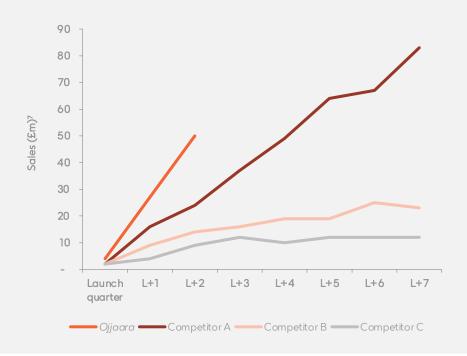
- Ojjaara/Omjjara £52m⁵ strong US launch in 2023, launched in the UK and Germany in Q1 2024
- Jemperli £80m⁵ +>100% continued momentum and growth
- Zejula £141m⁵ +27% strong growth across all regions from increased patient demand, higher volumes, and new launches in International



Progress in oncology driven by Ojjaara launch and lifecycle innovation

Ojjaara: fastest US launch uptake in value for a JAKi¹ in MF²

- Driven by strong execution
- US share in patients with anaemia: 14%³ in 1L⁴ and 28%³ in 2L⁵
- 56% of US physicians expect to increase prescribing *Ojjaara* in the next six months⁶



Jemperli: a potential backbone 108 therapy

- 1L⁴ dMMR⁹ EC¹⁰ new patient share up to ~33%¹¹, nearly doubled vs Q4 2023¹²
- RUBY Part 1: only IO⁸ combination to show statistically significant and clinically meaningful OS¹³ in the overall EC¹⁰ population; sBLA¹⁴ granted priority review by FDA
- RUBY Part 2: significantly improved PFS¹⁵ in 1L⁴, OS¹³ data due end of 2024

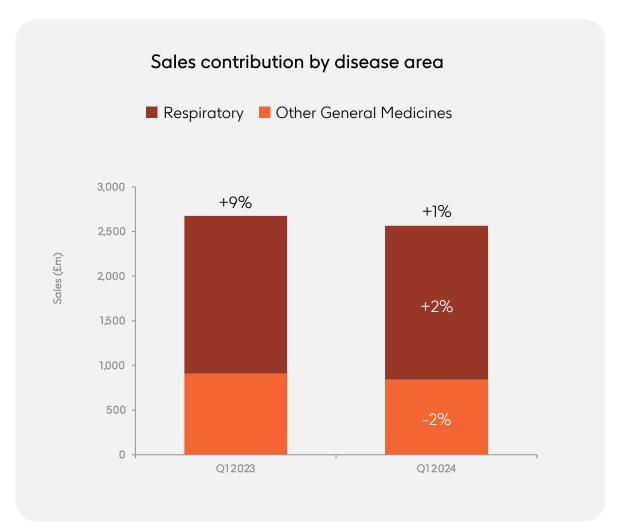
Blenrep: potential to redefine the treatment of relapsed or refractory multiple myeloma

- Encouraging DREAMM-7 and DREAMM-8 PFS¹⁵ data
- Strong OS¹³ trends, trials continue to follow up
- DREAMM-7 presented at ASCO¹⁶ Plenary Series in February 2024
- DREAMM-8 to be presented at ASCO¹⁶ in June
- Regulatory filing in H2 2024



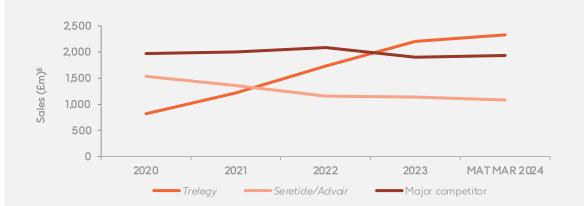
General Medicines: +1% driven by *Trelegy* and EP¹ in Emerging Markets

2024 guidance: mid-single digit % decrease



Respiratory +2%

 Trelegy +33%: most prescribed SITT² worldwide³, helping ~9.6 million patients since launch⁴



• GSK retaining share of voice and continuing strong competitive execution, including co-promotion with *Arexvy*

Other General Medicines -2%

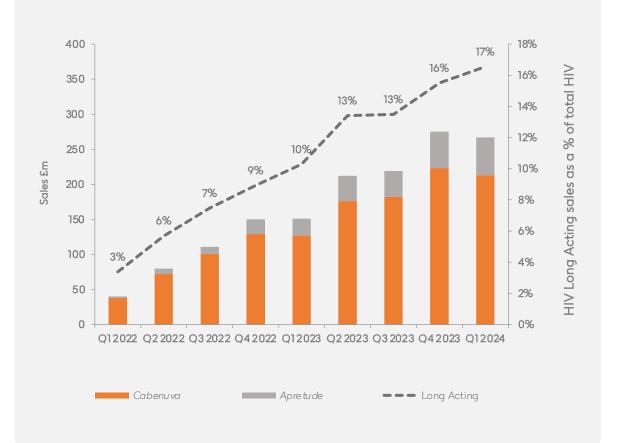
- Augmentin +10%,
- Up to \$550m sales exposure in 2024 related to US AMP⁶ Cap removal



Absolute values at AER; changes at CER for full year, unless stated otherwise 1. Established Products 2. Single inhaler triple therapy 3. Global geography analysed: Algeria, Argentina, Australia, Australia, Australia, Belarus, Belgium, Brazil, Bulgaria, Canada, Chile, China, Colombia, Costa Rica, Croatia, Czech, Denmark, Ecuador, Egypt, El Salvador, Finland, France, Germany, Greece, Guatemala, Honduras, Hong Kong, Hungary, India, Indonesia, Ireland, Italy, Japan, Jordan, Kuwait, Latvia, Lebanon, Luxembourg, Malaysia, Mexico, Morocco, Netherlands, New Zealand, Nicaragua, Norway, Pakistan, Panama, Peru, Philippines, Poland, Portugal, Puerto Rico, Romania, Russian Federation, Saudi Arabia, Serbia, Slovakia, Slovenia, South Korea, Spoth Korea, Spoth Korea, Spoth, Chilada, Tunisia, Turkey, United Arab Emirates, United Kingdom, United States, Uruguay, Vietnam, West Africa 4. Based on internal analysis by GSK using data from the following source: IQVIA MIDAS® global* monthly data, since launch up to (and including) February 2024, reflecting estimates of real-world activity. Measure: volume sales (units); ATC: R03L3. All rights reserved. Copyright IQVIA 5. GSK Annual Reports 2020-23; GSK Q1 2024 financial results; competitor financial results; FactSet exchange rates

HIV: Growing 14% in Q1 2024, with momentum in long-acting (LA) innovation

Positive momentum across LA portfolio



Performance driven by continued momentum of LA portfolio and strong delivery on *Dovato*

- Q1 2024 global sales of £1.6bn driven by 2% increase in market share
- LAI¹ delivered £267m, growing 83%
 - Cabenuva sales of £213m, growing 73%
 - Apretude sales of £54m, >100%
- Dovato sales of £483m, growing 27% leading oral 2DR²
- CROI³:
 - LATITUDE study indicated superiority of Cabenuva compared to daily oral therapy in adherence challenged populations
 - Positive phase I for Cab ULA⁴ Q4M⁵



Q1 2024 performance and 2024 guidance

Julie Brown, Chief Financial Officer



Strong start to 2024

	Q1 2023	Q1 2024	AER	CER	Key commentary on CER basis
Core results	£m	£m	%	%	
Sales	6,951	7,363	6	10	Sales grew +13% (excl. COVID-19 solutions)
Cost of sales	(1,752)	(1,733)	(1)	-	Mix benefits
Gross profit	5,199	5,630	8	14	
Gross profit margin	74.8%	76.5%	+170 bps	+250 bps	Improved +250 bps (excl. COVID-19 solutions)
SG&A	(2,065)	(1,979)	(4)	(2)	LSD growth ¹ excluding 6 ppt favourability from successful <i>Zejula</i> royalty dispute appeal
Research and development	(1,222)	(1,359)	11	14	Vaccines, Respiratory and Infectious Diseases investment
Royalties	180	151	(16)	(16)	Impact of lower Gardasil royalties
Operating profit	2,092	2,443	17	27	Grew +35% (excl. COVID-19 solutions)
Operating profit margin	30.1%	33.2%	+310 bps	+460 bps	Improved +580 bps (excl. COVID-19 solutions)
Earnings per share	37.0p	43.1p	16	28	EPS grew +37% (excl. COVID-19 solutions)
	Q1 2023	Q1 2024	AER	CER	
<u>Total results</u>	£m	£m	%	%	
Total operating profit	2,082	1,490	(28)	(18)	Total profit decrease primarily due to ViiV CCL remeasurement
Total operating profit margin	30.0%	20.2%	-970 bps	-780bps	
Total earnings per share	36.8p	25.7p	(30)	(19)	



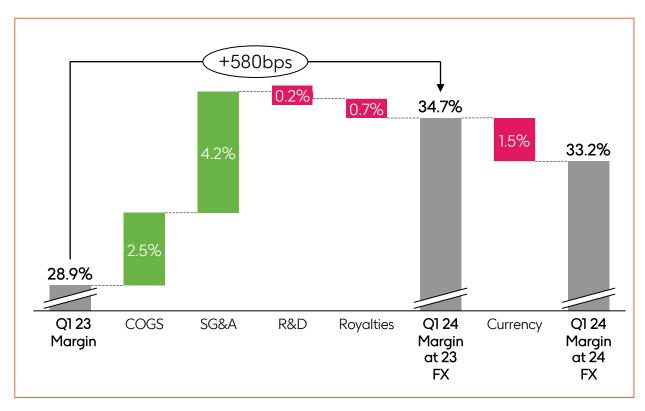
Q1 2024 core operating margin significantly higher

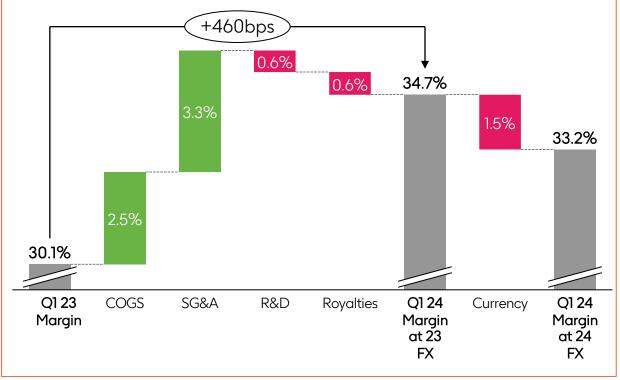
Excluding COVID-19 solutions +580 bps CER

Margin benefits driven by SG&A and gross margin

Including COVID-19 solutions +460 bps CER

Margin benefits driven by SG&A and gross margin







Note: Charts may not sum due to rounding

Q1 2024 free cash flow of £0.3bn

Cash generated from operations of £1.1bn

	£m Q1 2023	£m Q1 2024
Core operating profit	2,092	2,443
Decrease/(Increase) in working capital	(840)	(311)
Contingent consideration paid	(290)	(306)
Other CGFO	(675)	(700)
Cash generated from operations (CGFO ¹)	287	1,126
Taxation paid	(234)	(168)
Net capex ²	(518)	(535)
Other ³	(224)	(134)
Free cash flow	(689)	289

Key drivers of cash flow

Q12024

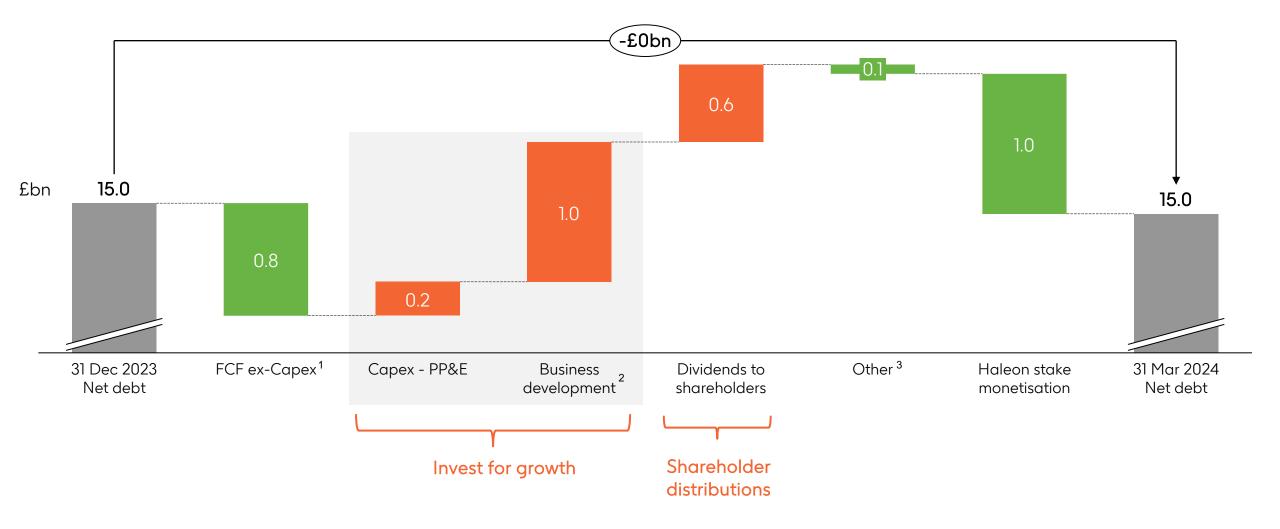
£0.8bn increase in cash generated from operations, mainly driven by:

- Higher Core Operating profit
- Improvement in working capital driven by the timing of US Vaccines receivables' collections

Net capital investment increased, offset by lower taxation and net interest paid



Capital deployment supports business growth and shareholder returns





2024 guidance at CER and excl. COVID-19 solutions

Upgraded guidance

Sales¹

5-7%

Towards upper part of range (Previously 5-7%)

Core operating profit¹

9-11%

(Previously 7-10%)

Core earnings per share¹

8-10%

(Previously 6-9%)

Modelling considerations

- Sales growth benefits from new product launches, which annualise in H2. Expect H1 growth to be higher than H2.
- Core operating profit growth benefits from higher operating leverage and product mix relative to H2. Expect H1 growth to be higher than H2.



IR Roadmap 2024 to 2025

	H1 2024	H2 2024*	2025
Execution	Full year 2023 resultsGuidance 2024Q1 2024 results	 Half-year 2024 results Q3 2024 results 	 Full-year 2024 results Guidance 2025 Q1 2025 results Half-year 2025 results Q3 2025 results
Pipeline Phase III and regulatory decisions ¹	Ojjaara/Omjjara: MOMENTUM, myelofibrosis (JP) Ojjaara/Omjjara: MOMENTUM, myelofibrosis (EU) Nucala: severe asthma (CN) Arexvy, RSV, 50-59 YoA² (US)	 Arexvy, RSV, 50-59 YoA (EU, JP) Nucala, CRSwNP³ (JP) Jemperli RUBY Part 1, 1L⁴ EC⁵ (US) 	 gepotidacin uUTI⁶, GC⁷ (US) MenABCWY 1st gen (US, EU)⁸ Shingrix adults 18+ years (CN) depemokimab ANCHOR-1/2 CRSwNP³, (US, JP) depemokimab SWIFT-1/2 SA⁹ (US, JP) Nucala MATINEE COPD¹⁰ (US) Nucala CRSwNP³ (CN) Blenrep DREAMM-7/8, 2L+ MM¹¹ (US, EU, JP) Jemperli RUBY (Part 1) 1L⁴ EC⁵ (EU) linerixibat cholestatic pruritus (US)
	• gepotidacin EAGLE-1, GC ⁷ • depemokimab SWIFT-1/2, SA ⁹ • Blenrep DREAMM-7, 2L+MM ^{1,1} • Jemperli RUBY, 1L dMMR/MSI-H ¹² EC ⁵ (EU) • Jemperli RUBY Part 1, 1L OS ¹³ EC ⁵ • Jemperli RUBY Part 2, 1L EC ⁵	 Bexsero infants (US) depemokimab ANCHOR-1/2, CRSwNP3 Nucala MATINEE, COPD¹⁰ Blenrep DREAMM-8, 2L+¹⁴ MM¹¹ Zejula FIRST 1L maintenance OC¹⁵ Zejula ZEAL, 1L maintenance NSCLC¹⁶ linerixibat GLISTEN, PBC¹⁷ 	 tebipenem PIVOT-PO, cUTI¹⁸ camlipixant CALM 1/2, RCC¹⁹ depemokimab OCEAN, EGPA²⁰ cobolimab COSTAR, 2L NSCLC¹⁶
Capital Allocation	Full-year 2023 dividend declarationDividend expectation 2024		 Full-year 2024 dividend declaration Dividend expectation 2025
Investor	Meet the management, Oncology	Meet the management, Early pipeline	



engagement

HIV long-acting combination decision 1. Includes phase III data readouts and regulatory decisions with the applicable geography denoted in brackets (United States (US), Europe (EU), Japan (JP), and China (CN) 2. Years of age 3. Chronic rhinosinusitis with nasal polyps 4. First-line treatment. 5. Endometrial cancer 6. Uncomplicated Urinary Tract Infections (EAGLE 2/3) 7. Urogenital gonorrhoea (EAGLE-1) 8. Regulatory submission and acceptance 9. Severe asthma with an eosinophilic phenotype 10. Chronic Obstructive Pulmonary Disease 11. Multiple Myeloma. Not included in the updated outlook 12. mismatch repair deficient (dMMR) or microsatellite instability-high (MSI-H) 13. Overall survival 14. Second-line and later treatment of relapsed or refractory multiple myeloma 15. Ovarian cancer

16. Non-Small Cell Lung Cancer 17. Cholestatic pruritus in primary biliary cholangitis 18. Complicated urinary tract infection 19. Refractory chronic cough 20. Eosinophilic granulomatosis with polyangiitis

Roadshows and Medical congresses

Focused on prevention and changing the course of disease



R&D based on science of the immune system and use of new platform and data technologies Leaders in development of new Vaccines and Specialty Medicines, for Infectious Diseases, HIV Respiratory/Immunology and Oncology Products that improve the health of millions of people, and sector leaders in ESG performance Strong momentum and improving outlook for sustained growth through the decade





Appendix



2024 full year outlook considerations to support modelling

	2023 growth excl. Covid	2024 Guidance	2024 Assumptions
Turnover	+14%	5-7%	Towards upper part of the range
- Vaccines	+24%	HSD – LDD	
- Specialty	+15%	LDD	
- HIV	+13%	HSD – LDD	
- Gen Meds	+5%	MSD decrease	Largely AMP Cap removal
Core Operating Profit	+16%	9-11%	SG&A: LSD increase R&D: increase broadly in-line with sales Royalties: £550-£600m; minimal Gardasil royalties
Core Op. Profit margin	28.6%	n/a	
Core EPS	+ 22%	8-10%	Interest: slightly lower than 2023 Core tax rate: around 17% Non-controlling interest: ViiV is main ongoing NCI
Dividend	58p	60p	

2021 – 2026 BIU 2021	2021 – 2026 BIU 2024
>5% CAGR	>7% CAGR
HSD CAGR	LDD CAGR
DD CAGR	DD CAGR
MSD CAGR	6-8% ¹
Broadly Stable	Broadly Stable
>10% CAGR	>11% CAGR
>30%	>31%



All guidance, outlooks and expectations regarding future performance should be read together with the section "Guidance and outlooks, assumptions and cautionary statements" on page 49 of GSK's Q1 2024 stock-exchange announcement. 2024 guidance growth at CER, unless stated otherwise. All outlook statements are given on a CER basis and use 2023 average exchange rates as a base. All values excluding COVID-19 solutions. CAGR is defined as the compound annual growth rate and shows the annualised average rate for growth in sales and Core operating profit between 2021 to 2026, assuming growth takes place at an exponentially compounded rate during those years.

1. As per HIV Meet The Management event, 28 September 2023

Q1 Total to Core operating profit reconciliation

	Q1 2023	Q1 2024	Key commentary on CER basis
	Operating profit (£m)	Operating profit (£m)	
Total results	2,082	1,490	-18% at CER
Intangible amortisation	169	196	
Intangible impairment	16	54	
Major restructuring	108	57	~£1.1bn benefits to date¹
Transaction-related	(271)	704	Primarily ViiV CCL ² movements
Divestments, significant legal and other	(12)	(58)	Fair value gains on investments, milestone and equity investment income, significant legal charges
Core results	2,092	2,443	+27% incl. COVID; +35% excl. COVID-19 solutions



Improved core earnings per share with +28% growth at CER

	Q1 2023 £m	Q1 2024 £m	Key commentary on CER basis
Core operating profit (OP)	2,092	2,443	+27% incl. COVID; +35% excl. COVID-19 solutions
Net finance expense	(170)	(132)	Lower bond interest costs and higher interest income
Share of associates	(2)	(1)	
Tax	(303)	(404)	
Tax rate	15.8%	17.5%	Broadly in-line with guidance of 17% in full-year
Non-controlling interests	(121)	(154)	Higher NCI related to ViiV
Core Profit attributable to shareholders	1,496	1,752	+28% incl. COVID
Core earnings per share (EPS)	37.0p	43.1p	+28% incl. COVID, +37% excl. COVID-19 solutions
Total EPS	36.8p	25.7p	-19% at CER
Weighted average number of shares (millions)	4,044	4,069	



Core results unless stated otherwise.

Quarterly summary of results

	2023					2024				
	Q1	Q2	Q3	Q4	FY	Qī	Q2	Q3	Q4	FY
Including COVID-19 solutions										
Sales (£m)	6,951	7,178	8,147	8,052	30,328	7,363				
Operating profit (£m)	2,092	2,170	2,772	1,752	8,786	2,443				
Operating margin	30.1%	30.2%	34.0%	21.8%	29.0%	33.2%				
Earnings per share (pence) post-share consolidation	37.0	38.8	50.4	28.9	155.1	43.1				
COVID-19 solutions impact										
Sales (£m)	132	41	1	20	194	1				
Operating profit (£m)	118	57	(4)	8	179	(1)				
Earnings per share (pence) post-share consolidation	2.5	1.2	(0.1)	0.2	3.8	0.0				
Excluding COVID-19 solutions impact										
Sales (£m)	6,819	7,137	8,146	8,032	30,134	7,362				
Operating profit (£m)	1,974	2,113	2,776	1,744	8,607	2,444				
Operating margin	28.9%	29.6%	34.1%	21.7%	28.6%	33.2%				
Earnings per share (pence) post-share consolidation	34.5	37.6	50.5	28.7	151.3	43.1				



Core results; some figures may not sum due to rounding.

Currency

2023 currency sales e	xposure ¹	
US\$	52%	
Euro €	19%	
Japanese ¥	4%	
Other ²	25%	

2024 core operating profit³

US \$: 10 cents movement in the average exchange rate for full year impacts core operating profit by approx. +/- 9.0%

Euro €: 10 cents movement in the average exchange rate for full year impacts core operating profit by approx. +/- 0.5%

Japanese ¥: 10 Yen movement in the average exchange rate for full year impacts core operating profit by approx. +/- 1.0%

			2023					2024		
Historical average exchange rates quarterly	Q1	Q2	Q3	Q4	FY 23	Qī	Q2	Q3	Q4	FY 24
US\$	1.22	1.25	1.26	1.25	1.24	1.27				
Euro €	1.14	1.15	1.16	1.15	1.15	1.16				
Japanese ¥	162	173	182	183	175	187				
Historical period end exchange rates										
US\$	1.24	1.26	1.23	1.27		1.26				
Euro €	1.14	1.17	1.16	1.15		1.17				
Japanese ¥	165	183	183	180		191				



Upcoming pipeline catalysts: 2024 and 2025

H2 2024

Regulatory decision

Arexvy: 50-59 YoA1

Omjjara: myelofibrosis

US

JP

H1 2024

 Arexvy: 50-59 YoA¹
 EU, JP

 Nucala: CRSwNP²
 JP

 Jemperli³: RUBY (Part 1)⁴. 1L EC⁵
 US

Regulatory submission acceptance aepotidacin: EAGLE-2/3, uUTI6 US MenABCWY vaccine 1st Gen EU depemokimab: SWIFT-1/2, asthma US depemokimab: ANCHOR-1/2, CRSwNP² US Nucala: MATINEE, COPD7 US Blenrep: DREAMM-7/8, 2L+ MM8 US, EU, JP Blenrep: DREAMM-7, 2L+ MM8 CN FU Jemperli³: RUBY (Part 1)⁴, 1L EC⁵

Late-stage phase III readouts depemokimab: SWIFT-1/2, asthma

1. Years of age 2. Chronic rhinosinusitis with nasal polyps 3. Tesaro asset 4. Overall population 5. Endometrial cancer 6. Uncomplicated urinary tract infection 7. Chronic obstructive pulmonary disorder 8. Multiple myeloma 9. Ovarian cancer 10. Non-small cell lung cancer 11. Primary biliary cholangitis 12. Urogenital gonorrhoea 13. Complicated urinary tract infection 14. Refractory chronic cough 15. Eosinophilic granulomatosis with polyangiitis polyps

Bexsero (infants US)

depemokimab: ANCHOR-1/2, CRSwNP²

Nucala: MATINEE, COPD7

Zeiula³: FIRST. 1L maintenance OC⁹

Zejula³: ZEAL, 1L maintenance NSCLC¹⁰

linerixibat: GLISTEN, cholestatic pruritus in PBC¹¹

gepotidacin: EAGLE-2/3, uUTI6 US gepotidacin: EAGLE-1, GC12 US MenABCWY vaccine 1st Gen US Shingrix: 18+ YoA CN depemokimab: SWIFT-1/2, asthma US. JP depemokimab: ANCHOR-1/2, CRSwNP² US, JP Nucala: CRSwNP2 CN Nucala: MATINEE, COPD7 US Blenrep: DREAMM-7/8, 2L+ MM8 US. EU. JP Jemperli³: RUBY (Part 1)⁴. 1L EC⁵ EU linerixibat: GLISTEN, cholestatic pruritus in PBC¹¹ US US Bexsero (infants US) gepotidacin: EAGLE-1, GC12 US .JP aepotidacin: EAGLE-J. uUTI6 tebipenem pivoxil: PIVOT-PO, cUTI¹³ US camlipixant: CALM-1/2, RCC14 US. EU EU. CN. JP depemokimab: SWIFT-1/2, asthma depemokimab: ANCHOR-1/2, CRSwNP² EU. CN. JP Nucala: MATINEE, COPD7 EU. CN Blenrep: DREAMM-8, 2L+ MM8 CN cobolimab3: COSTAR, 2L NSCLC10 US. EU linerixibat: GLISTEN, cholestatic pruritus in PBC¹¹ US, EU, CN, JP tebipenem pivoxil: PIVOT-PO, cUTI¹³ camlipixant: CALM-1/2, RCC14

depemokimab: OCEAN, EGPA¹⁵

cobolimab3: COSTAR, 2L NSCLC10

2025

Infectious diseases

Respiratory/Immunology

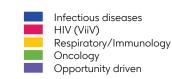
HIV (ViiV)

Oncology Opportunity driven

72 potential new vaccines and medicines in pipeline

Phase III / Registration – 18 assets

Arexvy (RSV vaccine)	Recombinant protein, adjuvanted*	RSV older adults (50-59 YoA)^
gepotidacin (GSK2140944)	BTI inhibitor*	Uncomplicated UTI**
bepirovirsen (GSK3228836)	Antisense oligonucleotide*	Chronic HBV infection**
Bexsero (MenB vaccine)	Recombinant protein, OMV	Meningitis B (infants US)
MenABCWY vaccine (GSK3536819)	Recombinant protein, OMV, conjugated vaccine	MenABCWY, 1st Gen^
tebipenem pivoxil (GSK3778712)	Antibacterial carbapenem*	Complicated UTI
ibrexafungerp (GSK5458448)	Antifungal glucan synthase inhibitor*	Invasive candidiasis
Nucala (mepolizumab)	Anti-IL5 antibody	COPD
depemokimab (GSK3511294)	Long-acting anti-IL5 antibody*	Asthma**
latozinemab (GSK4527223)	Anti-sortilin antibody*	Frontotemporal dementia ^{1**}
camlipixant (GSK5464714)	P2X3 receptor antagonist	Refractory chronic cough
Low carbon version of MDI ² , Ventolin (salbutamol)	Beta 2 adrenergic receptor agonist	Asthma ³
Ojjaara/Omjjara (momelotinib)	JAK1, JAK2 and ACVR1 inhibitor*	Myelofibrosis ^{A4}
Jemperli (dostarlimab)	Anti-PD-1 antibody*	Endometrial cancer^**
Zejula (niraparib)	PARP inhibitor*	Ovarian cancer**
Blenrep (belantamab mafodotin)	Anti-BCMA ADC*	Multiple myeloma
cobolimab (GSK4069889)	Anti-TIM-3 antibody*	Non-small cell lung cancer
linerixibat (GSK2330672)	IBAT inhibitor	Cholestatic pruritus in primary biliary cholangitis





72 potential new vaccines and medicines in pipeline

Phase II – 33 assets

GSK3437949	Recombinant protein, adjuvanted*	
GSK4406371	Live, attenuated	
GSK3536852	GMMA*	
GSK3528869	Viral vector with recombinant protein, adjuvanted*	
GSK4023393	Recombinant protein, OMV, conjugated vaccine	
GSK4178116	Live, attenuated	
GSK5101956	MAPS Pneumococcal 24-valent*	Adu
GSK5101955	MAPS Pneumococcal 24-valent paed*	Paediatr
GSK4106647	Recombinant protein, adjuvanted*	
GSK4348413	GMMA	
GSK4382276	mRNA*	
GSK4396687	mRNA*	
GSK3993129	Adjuvanted recombinant subunit	
GSK3943104	Recombinant protein, adjuvanted*	Therap
GSK5637608	Hepatitis B virus-targeted siRNA*	
GSK4077164	Bivalent GMMA	Invasive na
ganfeborole (GSK3036656)	Leucyl t-RNA synthetase inhibitor*	
sanfetrinem cilexetil (GV118819)	Serine beta lactamase inhibitor*	
alpibectir (BVL-GSK3729098)	Ethionamide booster*	
VH3810109	Broadly neutralizing antibody*	
VH3739937	Maturation inhibitor	
VH4004280	Capsid protein inhibitor	
VH4011499	Capsid protein inhibitor	
VH4524184	Integrase inhibitor*	
Benlysta (belimumab)	Anti-BLys antibody	Systemic sclerosis associate
GSK3858279	Anti-CCL17 antibody*	
GSK1070806	Anti-IL18 antibody	
GSK4527226 (AL-101)	Anti-sortilin antibody*	
GSK3915393	TG2 inhibitor*	
GSK5784283	TSLP monoclonal antibody*	
belrestotug (GSK4428859)	Anti-TIGIT antibody*	No
nelistotug (GSK6097608)	Anti-CD96 antibody*	
GSK4532990	HSD17B13 siRNA*	Non

Infectious diseases
HIV (ViiV)
Respiratory/Immunology
Oncology
Opportunity driven

Malaria fractional dose MMRV new strain Shigella

Chronic HBV infection^{1**}

MenABCWY, 2nd Gen¹

Varicella new strain

Adult pneumococcal disease

Paediatric pneumococcal disease

Human papillomavirus¹

Gonorrhoea¹

Seasonal flu

COVID-19

. . . .

Cytomegalovirus¹

Therapeutic herpes simplex virus¹

Chronic HBV infection

non-typhoidal salmonella**

Tuberculosis

Tuberculosis

_

Tuberculosis

HIV

HIV

HIV

HIV

HIV

temic sclerosis associated interstitial lung disease

Osteoarthritis pain**

Atopic dermatitis

Alzheimer's disease

Pulmonary fibrosis²

Asthma³

Non-small cell lung cancer**

Cana

Non-alcoholic steatohepatitis

^{*}In-license or other alliance relationship with third party ** Additional indications or candidates also under investigation

^{1.} In phase I/II study 2. Phase II study start imminent

^{3.} Phase II start expected in 2025

72 potential new vaccines and medicines in pipeline

Phase I - 21 assets

GSK3536867	Bivalent conjugate*	Salmonella (typhoid + paratyphoid A)
GSK2556286	Mtb cholesterol dependent inhibitor*	Tuberculosis
GSK3494245	Proteasome inhibitor*	Visceral leishmaniasis
GSK3772701	P. falciparum whole cell inhibitor*	Malaria
GSK4024484	P. falciparum whole cell inhibitor*	Malaria
GSK3882347	FimH antagonist*	Uncomplicated UTI
GSK3923868	PI4K beta inhibitor	Rhinovirus disease
GSK3965193	PAPD5/PAPD7 inhibitor	Chronic HBV infection ¹
GSK5251738	TLR8 agonist*	Chronic HBV infection
cabotegravir (GSK1265744)	Integrase inhibitor	HIV
GSK3888130	Anti-IL7 antibody*	Autoimmune disease
GSK3862995	Anti-IL33 antibody	COPD
GSK5462688	RNA-editing oligonucleotide*	Alpha-1 antitrypsin deficiency
GSK4347859	Interferon pathway modulator	Systemic lupus erythematosus
GSK4381562	Anti-PVRIG antibody*	Cancer
XMT-2056 ² (wholly owned by Mersana Theraprutics)	STING agonist ADC*	Cancer
belantamab (GSK2857914)	Anti-BCMA antibody	Multiple myeloma
GSK4524101	DNA polymerase theta inhibitor*	Cancer ¹
GSK5764227	ADC-targeting B7-H3*	Solid tumors
GSK5733584	ADC-targeting B7-H4*	Gynecologic malignancies
GSK4172239	DNMTI inhibitor*	Sickle cell disease





Changes since Q4 2023

Infectious diseases HIV (ViiV) Respiratory/Immunology Oncology Opportunity driven

Changes on pipeline

New to Phase I

GSK5764227: ADC targeting B7-H3, solid tumors

Removed from Phase I

GSK3186899: CRK-12 inhibitor, visceral leishmaniasis

Progressed from Phase I to Phase II

GSK3915393: TG2 inhibitor, pulmonary fibrosis nelistotug (GSK6097608): anti-CD96 antibody, cancer

New to Phase II

GSK5784283: TSLP monoclonal antibody, asthma

Achieved pipeline catalysts

Regulatory submission acceptances



Other events

bepirovirsen: Chronic HBV infection – FDA Fast Track Designation
gepotidacin: EAGLE-1, urogenital gonorrhoea — Positive phase III data readout
mRNA Seasonal flu — Phase II data readout
Cabenuva (cabotegravir + rilpivirine): LATITUDE positive phase III readout
latozinemab: Frontotemporal dementia³ – FDA Breakthrough Therapy Designation
Blenrep: DREAMM-8, 2L+ MM – Positive phase III data readout



Glossary

ADC	Antibody drug conjugate
AE	Adverse event
AESI	Adverse event of special interest
AIR	At increased risk
AUC	Area under curve
ВСМА	B-cell maturation antigen
BICR	Blinded Independent Central Review
BRCA	Breast cancer
CAE	Corneal adverse events
CBR	Clinical benefit rate
cCR	Complete clinical response
CKD	Chronic kidney disease
CfB	Change from baseline
CMV	Cytomegalovirus
CN	China
COPD	Chronic obstructive pulmonary disease
СР	Cholestatic pruritus
CRR	Complete response rate
CRSwNP	Chronic rhinosinusitis with nasal polyps
cUTI	Complicated urinary tract infection
CV	Cardiovascular
DDI	Drug-drug interaction
DFS	Disease-freee survival
DL	Dose level
DLT	Dose-limiting toxicity
dMMR	Deficient mismatch repair
DoR	Duration of response
DPNP	Diabetic peripheral neuropathic pain
EASI	Eczema Area and Severity Index
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EGPA	Eosinophilic granulomatosis with polyangiitis
FVC	Forced vital capacity
GC	Urogenital gonorrhea
GMMA	Generalised Modules for Membrane Antigens
GSI	Gamma secretase inhibitor
НА	Healthy adults
HBV	Hepatitis B virus
HES	Hypereosinophilic syndrome
Hgb	Hemoglobin
hSBA	Human serum bactericidal assay
HZ	Herpes zoster
IC	Immunocompromised
ICR	Independent central review
iNTS	Invasive non-typhoidal salmonella
ITT	Intention-to-treat
JP	Japan
LLOQ	Lower limit of quantitation
LRTS	Lower respiratory tract symptoms
MAD	Multiple ascending dose
MAE	Medical attended events
MDI	Metered dose inhaler
MAPS	Mulitple Antigen Presenting System
MM	Multiple myeloma
MMR	Measles, mumps and rubella
MMRV	Measles, mumps, rubella and varicella
MRD	Multiple rising dose
MSI-H	Microsatellite instability high
NASH	Nonalcoholic steatohepatitis
NRS	Numeric Rating Scale

NSCLC	Non-small cell lung cancer
OMV	Outer membrane vesicle
ORR	Overall response rate
OS	Overall surival
PBC	Primary biliry cholangitis
PFS	Progression-free survival
PFS2	Time to second disease progression or death
PK	Pharmacokinetic
PMF	Primary myelofibrosis
Post-PV/ET MF	Post-essential thrombocythemia myelofibrosis
RCC	Refractory chronic cough
RL	Repeat dose level
RRMM	Relapsed/refractory multiple myeloma
RSV	Respiratory syncytial virus
SAD	Single ascending dose
SAE	Serious adverse event
siRNA	Small interfering RNA
SoC	Standard of care
SSc-ILD	Systemic sclerosis associated interstitial lung disease
TOC	Test of cure
TTBR	Time to best response
TTD	Time to treatment discontinuation
TTP	Time to tumour progression
TTR	Time to treatment response
UTI	Urinary tract infection
uUTI	Uncomplicated urinary tract infection
VGPR	Very good partial remission
VSP	Vital sign parameters
YoA	Years of age



Assumptions and basis of preparation related to 2024 guidance

In outlining the guidance for 2024, the Group has made certain assumptions about the healthcare sector (including regarding possible governmental legislative and regulatory reform), the different markets and competitive landscape in which the Group operates and the delivery of revenues and financial benefits from its current portfolio, its development pipeline and restructuring programmes.

These planning assumptions as well as operating profit and earnings per share guidance and dividend expectations assume no material interruptions to supply of the Group's products, no material mergers, acquisitions or disposals, no material litigation or investigation costs for the Company (save for those that are already recognised or for which provisions have been made) and no change in the Group's shareholdings in ViiV Healthcare. The assumptions also assume no material changes in the healthcare environment or unexpected significant changes in pricing as a result of government or competitor action. The 2024 guidance factors in all divestments and product exits announced to date.

Notwithstanding our guidance, outlooks and expectations there is still uncertainty as to whether our assumptions, guidance, outlooks and expectations will be met.



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