

*Sanofi Q1: robust 7% sales growth driven by launches, underpins full-year guidance*

Paris, April 25, 2024

**Sales growth of 6.7% at CER and business EPS<sup>(1)</sup> of €1.78**

- Dupixent sales up 24.9% to €2,835 million, on target to deliver ~€13bn in 2024
- Pharma launches up 90.5% to €606 million, led by Nexviazyme and ALTUVIIIIO
- Vaccines sales up 5.6%, boosted by Beyfortus
- Consumer Healthcare (CHC) up 9.0%, benefiting from the Qunol acquisition in Physical and Mental Wellness and strong Digestive Wellness performance
- R&D expenses grew 11.8%, reflecting strategic shift of resources into development
- SG&A expenses grew 2.9%, less than sales growth
- Business EPS<sup>(1)</sup> of €1.78, down 17.6% reported and 7.4% at CER
- IFRS EPS of €0.91, down 43.1% reported

**R&D transformation advanced further**

- Three regulatory approvals: new Dupixent indications in the US and Japan and Beyfortus in Japan
- Positive phase 3 results for rilzabrutinib in immune thrombocytopenia, a rare disease
- Reaffirming increasing pipeline news flow over 2024-2025, including 12 phase 3 data readouts

**Corporate Social Responsibility progress**

- Sanofi Global Health Unit: making a difference for patients in low- and middle-income countries
- Cancer and work: Sanofi supporting health and wellbeing in the workplace

**Business EPS guidance reiterated**

- Sanofi expects 2024 business EPS<sup>(1)</sup> to remain roughly stable excluding the impact of an expected effective tax rate increase to 21% and decrease low single-digit<sup>(2)</sup> at CER including the higher expected tax rate, barring unforeseen major adverse events. Applying average April 2024 exchange rates, the currency impact on 2024 business EPS is estimated between -5.5% to -6.5%.

***Paul Hudson, Chief Executive Officer, commented:***

*"We are off to an excellent start in 2024, delivering on our strategic priorities and a transformation of our portfolio of medicines and vaccines to become a development-driven, tech-powered biopharma company committed to serving patients and accelerating growth. Continued strong performance by Dupixent and our new launches drove sales growth of seven percent. In parallel, we are delivering on our promise of increased investments in our late-stage pipeline to fully realize its value for patients and Sanofi. We are awaiting regulatory decisions for Dupixent in COPD, a progressive disease with limited effective treatment options. If approved, Dupixent will be the first biologic treatment in COPD. With the robust progress in our portfolio transformation, we reaffirm our financial guidance for 2024."*

	Q1 2024	Change	Change at CER
IFRS net sales reported	€10,464m	+2.4%	+6.7%
IFRS net income reported	€1,133m	-43.2%	–
IFRS EPS reported	€0.91	-43.1%	–
Free cash flow <sup>(3)</sup>	€(309)m	-120.1%	–
Business operating income	€2,843m	-14.7%	-4.2%
Business net income <sup>(1)</sup>	€2,219m	-17.8%	-7.4%
Business EPS <sup>(1)</sup>	€1.78	-17.6%	-7.4%

*Changes in net sales are expressed at constant exchange rates (CER) unless otherwise indicated (definition in Appendix 7). (1) In order to facilitate an understanding of operational performance, Sanofi comments on the business net income statement. Business net income is a non-IFRS financial measure (definition in Appendix 7). The consolidated income statement for Q1 2024 is provided in Appendix 3 and a reconciliation of reported IFRS net income to business net income is set forth in Appendix 4; (2) 2023 business EPS was €8.11; (3) Free cash flow is a non-IFRS financial measure (definition in Appendix 7).*

## Q1 2024 summary

The performance shown in this press release covers the three-month period to 31 March 2024 (the quarter or Q1 2024) compared to the three-month period to 31 March 2023 (Q1 2023) respectively and all percentage changes in sales in this press release are at CER<sup>1</sup>, unless stated otherwise.

In Q1 2024, on a reported basis, Sanofi sales were €10,464 million, an increase of 2.4%. Exchange rate movements had a negative effect of 4.3 percentage points (pp). At CER, sales increased by 6.7%.

### Sales breakdown

Net sales (€ million)	Q1 2024	Change at CER	% of total sales
<b>Biopharma</b>	<b>8,939</b>	+6.3%	85.4%
Pharma	7,762	+6.4%	74.2%
Vaccines	1,177	+5.6%	11.2%
<b>CHC</b>	<b>1,525</b>	+9.0%	14.6%
<b>Total Sales</b>	<b>10,464</b>	<b>+6.7%</b>	<b>100 %</b>

### Business operating income

Business operating income (BOI) decreased by 14.7% to €2,843 million. At CER, BOI decreased by 4.2%. The ratio of BOI to net sales decreased 5.4pp to 27.2% (down 3.3pp to 29.3% at CER).

### Acquisitions and major collaborations

On January 23, Sanofi and Inhibrx, Inc., entered into an agreement under which Sanofi has agreed to acquire Inhibrx following the spin-off of non-INBRX-101 assets into New Inhibrx for \$30.0 in cash, a contingent value right of \$5.0 and 0.25 shares of a new publicly traded company that will retain Inhibrx's non-INBRX-101 assets. INBRX-101 is a human recombinant protein that holds the promise of allowing alpha-1 antitrypsin deficiency patients to achieve normalization of serum AAT levels with less frequent dosing. The acquisition supports Sanofi's strategy and complements 30-years of heritage in rare diseases and proven industry leadership in immunology and inflammation. This acquisition is subject to customary closing conditions and is expected to close in Q2 2024.

## Sales by geographic region

Net sales (€ million)	Q1 2024	Change at CER
<b>United States</b>	<b>4,316</b>	+7.3%
<b>Europe</b>	<b>2,480</b>	-4.0%
<b>Rest of World</b>	<b>3,668</b>	+13.8%
of which China	757	+6.2%

**US** sales increased by 7.3% to €4,316 million. The strong performance of Dupixent as well as the Pharma launches and Beyfortus late-season deliveries were partially offset by the impact of generic competition on Aubagio as well as lower sales of Lantus.

**Europe** sales decreased by 4.0% to €2,480 million reflecting the impact of generic competition on Aubagio, lower sales of Lovenox and a high base of comparison for Vaccines due to COVID-19 vaccine sales recorded in Q1 2023 which more than offset the strong performance of Dupixent.

**Rest of World** sales increased by 13.8% to €3,668 million, mainly driven by Dupixent, Beyfortus launch in two Southern Hemisphere countries and CHC. Sales in **China** increased by 6.2% to €757 million driven by Dupixent, Toujeo and Plavix.

<sup>1</sup> See Appendix 9 for definitions of financial indicators.

## Biopharma

The Biopharma segment includes **Pharma** and **Vaccines**. Sales increased by 6.3% to €8,939 million, driven by continued strong performance of Dupixent, Pharma launches and Vaccines. The divestments of medicines/portfolio streamlining had a negative impact of 0.7pp.

## Pharma

### Dupixent

Net sales (€ million)	Q1 2024	Change at CER
Dupixent	2,835	+24.9%

**Dupixent** (collaboration with Regeneron) sales increased by 24.9% to 2,835 million. In the US, in-market performance was strong with total prescriptions up by 25% (year-over-year) and new-to-brand prescriptions up by 22%, impacted by the customary dynamics of the annual reset of insurance plans. Sales increased by 16.8% to €2,030 million and were driven by continued strong demand in the approved indications: atopic dermatitis (AD), asthma, chronic rhinosinusitis with nasal polyposis (CRSwNP), eosinophilic esophagitis and prurigo nodularis. In Europe, Dupixent sales increased by 33.6% to €371 million reflecting continued growth in AD, asthma and CRSwNP. In the Rest of World region, sales reached €434 million, an increase of 67.5%, driven mainly by sales in Japan and China.

### Pharma launches

Net sales (€ million)	Q1 2024	Change at CER
Nexviazyme/Nexviadyne	152	+96.3%
ALTUVIIIIO	122	—%
Sarclisa	106	+28.7%
Rezurock	93	+40.3%
Cablivi	59	+3.4%
Xenpozyme	35	+100.0%
Enjaymo	29	+87.5%
TZIELD	10	—%
<b>Total Pharma launches</b>	<b>606</b>	<b>+90.5%</b>

**Nexviazyme/Nexviadyne** sales were €152 million, including €85 million in the US, and increased by 96.3% driven by the conversion from Myozyme/Lumizyme in the eligible late-onset Pompe disease population and by new patients. Pompe franchise (Nexviazyme/Nexviadyne + Myozyme/Lumizyme) sales were at €343 million, an increase of 15.2%. Nexviazyme/Nexviadyne sales now represent 44% of the Pompe franchise.

**ALTUVIIIIO** (hemophilia A) was launched at the end of March 2023 and generated sales of €122 million, mainly in the US where ALTUVIIIIO growth was driven by patient switches, the majority coming from medicines other than Eloctate. The hemophilia A franchise (ALTUVIIIIO + Eloctate) sales were €208 million and increased by 79.0%, improving Sanofi's share in the factor and the overall hemophilia A market.

**Sarclisa** (multiple myeloma) sales were €106 million, an increase of 28.7%, driven by strong growth in the US and the Rest of World.

**Rezurock** (chronic graft-versus-host disease) sales were €93 million, an increase of 40.3%, driven by improved patient adherence and new patients in the US as well as launches in China and UK.

**Cablivi** (acquired thrombotic thrombocytopenic purpura) sales increased by 3.4% to €59 million, primarily driven by use in the US.

**Xenpozyme** (acid sphingomyelinase deficiency) sales were €35 million, an increase of 100.0%, driven by use in the US and Europe.

**Enjaymo** (cold agglutinin disease) sales were €29 million, mainly generated in the US and Japan.

**TZIELD** (delay onset of type 1 diabetes) sales were €10 million driven by increased screenings and infusions. The launch continues to be supported by targeted investments, improved awareness and availability of guidelines.

## Other main medicines

Net sales (€ million)	Q1 2024	Change at CER
Lantus	360	-15.4%
Toujeo	321	+18.0%
Lovenox	262	-13.9%
Fabrazyme	253	+7.7%
Plavix	238	+6.8%
Cerezyme	214	+23.0%
Myozyme/Lumizyme	191	-13.6%
Alprolix	130	+6.4%
Praluent	121	+25.5%
Thymoglobulin	117	+12.8%
Aubagio	102	-74.7%
Eloctate	86	-24.6%
Cerdelga	83	+15.1%

**Lantus** sales were €360 million, a decrease of 15.4%. In the US, sales decreased by 13.6% to €112 million, reflecting a high comparison in Q1 2023 ahead of an adjustment to reflect increasing sales to government customers. These developments more than offset an increase in volume due to a competitor medicine withdrawing from the US market. In China, sales decreased by 52.5% million to €27 million as patients are shifting to Toujeo.

**Toujeo** sales increased by 18.0% to €321 million driven by China where the market share now exceeds that of Lantus. In the US, sales decreased mainly due to a shift in sales channel mix, resulting in a lower average net price, more than offsetting an increase in volume and market share.

**Lovenox** sales decreased by 13.9% to €262 million, reflecting an impact from volume-based procurement in China as well as biosimilar competition, which was largely offset by growth in some other countries.

**Fabrazyme** sales increased by 7.7% to €253 million, driven by the Rest of World.

**Plavix** sales increased by 6.8% to €238 million driven by use in China.

**Cerezyme** sales increased by 23.0% to €214 million, reflecting growth in high-inflation countries (Argentina and Turkey), part of the Rest of World. Gaucher disease franchise (Cerezyme + Cerdelga) sales were €297 million, an increase of 20.8%.

**Myozyme/Lumizyme** sales decreased by 13.6% to €191 million reflecting the conversion to Nexviazyme/Nexviadyne mentioned above.

**Alprolix** sales were €130 million, an increase of 6.4%, driven by the Rest of World.

**Praluent** sales were €121 million, an increase of 25.5%, driven by Europe and China.

**Thymoglobulin** sales were €117 million, an increase of 12.8%, driven by China.

**Aubagio** sales decreased by 74.7% to €102 million, reflecting the loss of exclusivity in the US in March 2023 and the competition from generics across all regions, including Europe where generics entered the market at the end of September 2023. The negative impact is anticipated to continue during 2024 as the losses of exclusivity annualize.

**Eloctate** sales were €86 million, a decrease of 24.6%, reflecting patients converting to **ALTUVIIIO**.

**Cerdelga** sales were €83 million, an increase of 15.1%, reflecting continued growth in the US, Europe and favorable shipment pattern.

## Vaccines

Net sales (€ million)	Q1 2024	Change at CER
Polio/Pertussis/Hib vaccines incl. Boosters	636	-0.5%
Meningitis, Travel and endemic vaccines	286	+7.7%
RSV (Beyfortus)	182	— %
Influenza vaccines	73	+38.1%
Others	—	-100.0%

**Vaccines** sales increased by 5.6% to €1,177 million driven by strong Beyfortus uptake partly offset by the absence of COVID-19 vaccines sales this quarter compared to €167 million in Q1 2023.

**Polio/Pertussis/Hib** (PPH) vaccines sales, including Boosters, decreased by 0.5% to €636 million. Positive Rest of World sales driven by performance and favorable phasing were offset by declining sales in the US, where Vaxelis became market leader in the three-dose primary series market at the end of 2023. Vaxelis in-market sales are not consolidated by Sanofi but profits are shared equally between Sanofi and Merck & Co.

**Meningitis, Travel and endemic** vaccines sales increased by 7.7% to €286 million reflecting a favorable US Centers for Disease Control and Prevention buying pattern and increased MenQuadfi penetration in Europe.

**Beyfortus** sales reached €182 million, reflecting late deliveries in the US and new implementation of "All Infant Protection" programs in Chile and Australia. No Beyfortus sales are expected in Q2 2024 due to the early delivery in Q1 and the specific seasonality.

**Influenza vaccines** sales reached €73 million, benefiting from positive phasing with earlier deliveries to Southern Hemisphere countries compared to Q1 2023.

## Biopharma business operating income

BOI of Biopharma decreased by 15.2% to €2,365 million. At CER, Biopharma BOI decreased by 5.7% reflecting lower gross profit ratio, higher R&D expenses, and lower capital gains from divestments of medicines/portfolio streamlining as compared to Q1 2023. The ratio of BOI to net sales decreased by 5.5pp to 26.5% (28.3% at CER).

## Pipeline update

Sanofi has 77 projects in a pipeline across four main therapeutic areas (Immunology, Rare diseases, Neurology, and Oncology) and Vaccines, including 39 potential new medicines (NMEs) and vaccines. The following section highlights significant developments in the late- and mid-stage pipeline in the quarter and recently:

### Highlights of the quarter

Regulatory approvals	Dupixent – pediatric eosinophilic esophagitis (EoE) (US) Dupixent – chronic spontaneous urticaria (CSU) (JP) Beyfortus – RSV prevention (JP)
Regulatory submission acceptance	Dupixent – COPD (US, EU, CN) (US priority review, PDUFA June 27)
Phase 3 data readouts	rilzabrutinib - immune thrombocytopenia (ITP) (primary endpoint met)
Major pipeline advancements	SP0125 – toddler RSV vaccine (phase 3)

## Immunology

### Dupixent (dupilumab)

- The US Food and Drug Administration (FDA) updated the label for *Dupixent* in **atopic dermatitis** (AD), adding efficacy and safety data for patients aged 12 years and older with AD with uncontrolled moderate-to-severe hand and/or foot involvement.
- The FDA approved *Dupixent* for the treatment of pediatric patients aged 1 to 11 years, weighing at least 15 kg, with **eosinophilic esophagitis** (EoE). Dupixent is now the first and only medicine approved in the US specifically indicated to treat these patients.
- The Ministry of Health, Labour and Welfare in Japan granted regulatory approval for *Dupixent* for the treatment of **chronic spontaneous urticaria** (CSU) in people aged 12 years and older whose disease is not adequately controlled with existing therapy. Japan is the first country to approve Dupixent for CSU, emphasizing the value of Dupixent as a novel treatment option to manage this disease in patients with unmet medical needs. A regulatory submission is also under review in the European Union based on data from the LIBERTY-CUPID clinical study program.
- The FDA accepted for Priority Review the supplemental Biologics License Application for *Dupixent* in a sixth potential indication as an add-on maintenance treatment in certain adult patients with uncontrolled **chronic obstructive pulmonary disease** (COPD). The target action date for the FDA decision is June 27, 2024. Regulatory submissions are also under review in the European Union and China.

### amlitelimab (OX40L mAb)

Positive results from part 2 of *amlitelimab* STREAM-AD phase 2b study showed sustained improvement of signs and symptoms for 28 weeks in adults with moderate-to-severe **AD** who previously responded to amlitelimab and continued treatment. High responder rates were also observed in participants who were taken off amlitelimab. The safety profile was consistent with part 1 of the study with amlitelimab being well-tolerated and no new safety concerns identified. These results were presented as part of a late-breaking session at the American Academy of Dermatology 2024 Conference in San Diego and support the dosing of once every 12 weeks now being assessed in a larger phase 3 clinical program (OCEANA). Enrollment of the four main AD studies is on track and designed to evaluate the on- and off-treatment efficacy and safety in adults and adolescents, with a regulatory submission anticipated in 2027.

### itepekimab (IL33 mAb)

The first participants were treated in the phase 2 study evaluating *itepekimab* in patients with **bronchiectasis** (clinical study identifier: NCT06280391).

### rilzabrutinib (BTK inhibitor)

Positive results from the RILECSU phase 2 study showed that *rilzabrutinib* significantly improved itch, hives and urticaria in adults with **moderate-to-severe CSU** whose symptoms are not adequately controlled by H1-antihistamines. The data were presented in a late-breaking poster at the 2024 American

Academy of Allergy, Asthma and Immunology Annual Meeting in Washington, DC, US and form the basis for the phase 3 program which is on track to start in H2 2024.

Additionally, results from the second (high-dose) cohort of the ACT17208 phase 2 study of *rilzabrutinib* showed a higher relative reduction magnitude in loss of asthma control and confirmed meaningful improvement in symptoms with overall good safety in patients with **moderate-to-severe asthma**. The data are anticipated to be presented at the upcoming American Thoracic Society International Conference.

#### **frexalimab** (CD40L mAb)

The development of *frexalimab* in **Sjögren's Syndrome** (SjS), an immunological disorder with currently no approved disease-modifying treatments, was discontinued based on the findings of a phase 2 study. The 84-patient study evaluated the therapeutic efficacy of one dose level of frexalimab versus placebo over 12 weeks in adult patients with primary SjS, assessed by the change of the EULAR SjS Disease Activity Index (ESSDAI). The data confirmed pharmacologic activity and well-tolerated safety profile but not the necessary efficacy outcomes to continue to move forward the development in this indication. Development of frexalimab continues in relapsing multiple sclerosis (RMS) and secondary-progressive multiple sclerosis (SPMS) in phase 3, and type 1 diabetes and systemic lupus erythematosus in phase 2.

### **Rare diseases**

#### **Nexviazyme** (avalglucosidase alfa)

New data from the Mini-COMET phase 2 long-term extension study of *Nexviazyme* in pediatric patients with **infantile-onset Pompe disease** (IOPD), suggested that it meaningfully improved ptosis, or drooping eyelid, over nearly three years. Data were featured at the 20th annual WORLDSymposium™, along with positive safety debut data from the Baby-COMET phase 3 study, the first study in never-before-treated patients with IOPD, and the first study in over 20 years of any treatment in naïve IOPD patients. The phase 3 Baby-COMET study is currently ongoing and is expected to read out in 2025, followed by a regulatory submission later in the year.

#### **venglustat** (GCS inhibitor)

The AMETHIST phase 3 study of *venglustat* for the treatment of **GM2 gangliosidosis** was discontinued based on the absence of positive trends on clinical endpoints. The data reinforced the favorable safety profile and do not impact the other indications currently being tested.

#### **rilzabrutinib** (BTK inhibitor)

In April, *rilzabrutinib* met the primary endpoint of durable platelet response in the LUNA 3 phase 3 study in adult patients with persistent or chronic **immune thrombocytopenia** (ITP), showing clinically and statistically significant result in a population of patients with primary ITP that had been refractory to prior therapy. The safety profile of rilzabrutinib was favorable and consistent with that reported in previous studies. Regulatory submission is planned for H2 2024. Previously, rilzabrutinib was granted Fast Track Designation and Orphan Drug Designation by the US FDA.

### **Neurology**

#### **oditrasertib** (RIPK1 inhibitor)

The phase 2 study of *oditrasertib* (also known as SAR443820) in **amyotrophic lateral sclerosis** (ALS) was discontinued due to the primary endpoint of improvement based on ALS Functional Rating Scale-Revised (ALSFRRS-R), the conventional assessment of physical function in activities of daily living for patients, not being met. The parties are committed to sharing the full efficacy and safety data at a forthcoming medical meeting.

Sanofi will continue the K2 phase 2 study of *oditrasertib* in participants with **MS**.

#### **frexalimab** (CD40L mAb)

Phase 2 data for *frexalimab* in MS were published in The New England Journal of Medicine, demonstrating that frexalimab significantly slowed disease activity in people with RMS, corresponding to an 89 percent and 79 percent reduction in new Gd+ T1 brain lesions at 12 weeks in the high- and low-dose treatment arms compared to placebo, meeting the study's primary endpoint. Findings also showed both doses of frexalimab provided a significant reduction in new or enlarging T2 lesions, a secondary endpoint of the

study. Phase 3 studies in RMS and non-relapsing secondary-progressive MS (nrSPMS) are initiated with regulatory submission expected in 2027.

## **Vaccines**

**Beyfortus** (nirsevimab)

*Beyfortus*, a long-acting monoclonal antibody, was approved in Japan for the prophylaxis of lower respiratory tract disease (LRTD) caused by respiratory syncytial virus (RSV) in all neonates, infants and children entering their first RSV season, and the prevention of RSV LRTD in neonates, infants and children at risk of serious RSV infection entering their first or second RSV season.

**SP0125** (live attenuated RSV vaccine for toddlers)

The study evaluating *SP0125* for the prevention of **respiratory syncytial virus (RSV) in toddlers** entered phase 3.



## Anticipated major upcoming pipeline milestones

	Medicine/vaccine	Indication	Description
<b>H1 2024</b>	Dupixent	COPD	Regulatory decision (US)
		CSU	Regulatory submission (EU)
	Kevzara	Polyarticular juvenile idiopathic arthritis	Regulatory decision (US)
	Sarclisa	Multiple myeloma (MM), front line (1L) (IMROZ study)	Regulatory submission (US)
	MenQuadfi	Meningitis six weeks+	Phase 3 data
<b>H2 2024</b>	Dupixent	COPD	Regulatory decision (EU)
		CSU	Phase 3 data
		Bullous pemphigoid (BP)	Phase 3 data
	rilzabrutinib	ITP	Regulatory submission
		IgG4-related disease	Phase 2 data
		Warm autoimmune hemolytic anemia	Phase 2 data
	amlitelimab	Asthma	Phase 2 data
	Anti-TL1A	Inflammatory bowel disease	Phase 2 data (interim analysis)
	fitusiran	Hemophilia A/B	Regulatory submission (US)
	tolebrutinib	RMS	Phase 3 data Regulatory submission (US)
		nrSPMS	Phase 3 data Regulatory submission (US)
	Sarclisa	MM, relapsed/refractory (R/R) (IRAKLIA study), subcutaneous	Phase 3 data
MM, newly diagnosed (GMMG study)		Phase 3 data	
MenQuadfi	Meningitis six weeks+	Regulatory submission (US)	
<b>2025</b>	Dupixent	COPD	Regulatory decision (CN)
		CSU	Regulatory submission (US)
		BP	Regulatory submission (US)
	itepekimab	COPD	Phase 3 data Regulatory submission (US)
	amlitelimab	Hidradenitis suppurativa (HS)	Phase 2 data
		Alopecia areata	Phase 2 data
	eclitasertib	Ulcerative colitis	Phase 2 data
	frexalimab	Systemic lupus erythematosus	Phase 2 data
	IRAK4 degrader	AD	Phase 2 data
		HS	Phase 2 data
	Oral TNFR1si	Psoriasis	Phase 2 data
		Rheumatoid arthritis	Phase 2 data
	lunsekimig	Asthma	Phase 2 data
	TNFa/OX40L	HS	Phase 2 data
	Nezviazyme	IOPD	Phase 3 data
			Regulatory submission (US)
	venglustat	Fabry disease	Phase 3 data
			Regulatory submission (US)
	tolebrutinib	Primary progressive MS	Phase 3 data
			Regulatory submission (US)
Sarclisa	MM, R/R (IRAKLIA study), subcutaneous	Regulatory submission (US)	
	MM, newly diagnosed (GMMG study)	Regulatory submission (US)	
SP0256	RSV older adults	Phase 2 data	
SP0087	Rabies	Phase 3 data	
		Regulatory submission (US)	

An update of the Sanofi pipeline as of March 31, 2024, is available at: <https://www.sanofi.com/en/our-science/our-pipeline>.

## Consumer Healthcare

Net sales (€ million)	Q1 2024	Change at CER
Seasonal symptoms & pain relief	693	-1.0%
Wellness brands	660	+26.3%
Others	172	-5.8%

Consumer Healthcare (CHC) sales increased by 9.0% to €1,525 million supported by growth in the Rest of World region and the US which included the acquisition of Qunol. The divestments of non-core products had a negative impact of -1.6pp, mainly reflected in the Others category in the first quarter. Excluding divestments and the Qunol acquisition, CHC sales growth was 5% in the quarter.

### CHC business operating income

BOI of CHC decreased by 11.6% to €472 million. At CER, the BOI of CHC was up 3.0% mainly reflecting the consolidation of Qunol. The ratio of BOI to net sales decreased by 4.7pp to 31.0% (33.8% at CER) compared to Q1 2023.

During the quarter, Sanofi reached an agreement in principle to resolve the vast majority of Zantac personal injury state court cases in US state courts outside of Delaware. Assuming the agreement concludes in due course, Sanofi is settling these cases, not because Sanofi believes the claims have any merit, but rather to avoid expenses and ongoing distraction of the litigation, and with no concession of liability. This resolution is not anticipated to have any material financial impact.

## Corporate Social Responsibility update at the end of the Q1 2024

### Access to healthcare

#### Sanofi Global Health Unit: making a difference for our patients in low- and middle-income countries

Sanofi's Global Health Unit (GHU) works to address today's many growing healthcare challenges – with a focus on countries with the highest unmet medical needs – through a self-sustained not-for-profit social business model.

Sanofi's GHU aims to provide access to a broad portfolio of medicines in 40 countries with the highest unmet medical needs. To that end the GHU created Impact, a unique not-for-profit brand with 30 standard-of-care medicines produced by Sanofi, some of which are considered essential by the World Health Organization (WHO). The Impact medicines cover a wide range of therapeutic areas including diabetes, cardiovascular disease, tuberculosis, malaria and cancer.

Sanofi's GHU aims to reach two million people with non-communicable disease (NCD) care in its 40 countries in scope by 2030. Since its creation in 2021, the GHU has made significant progress towards its objective, having already treated 506,130 NCD patients in 31 countries as of the end of March 2024.

To support the set up and development of sustainable healthcare systems, the GHU is also working closely with local communities, authorities and non-governmental organizations to develop disease awareness programs and establish partnerships to drive better care through:

- strengthening supply chains;
- conducting medical training;
- providing services to patients.

Sanofi's GHU has engaged with Ministries of Health and other partners in several countries, including Rwanda, Uganda, Tanzania and Cambodia. As of March 2024, the GHU pilots 44 active partnerships in 21 countries. Selected examples of projects supported are described below:

Name	Therapeutic Area	Country(s)	Activity pillar(s)	Overview and progress in numbers
PharmAccess	Cardio Diabetes	Zanzibar	Patient Care model	The project is an integrated patient-centered model of care aiming at improving diagnosis and disease management for patients with cardio-metabolic diseases through a care bundle consisting of access to patient group meetings, digital self-management support, remote care and medications.
CHAZ FBO Zambia	Cardio Diabetes	Zambia	Scaling Patient Care services with faith-based organizations	The primary goal is to institutionalize NCD Prevention WHO Best Buys as a standard of care within the church health institutions participating in the project. It includes building the capacity of health workers and community educators in church health institutions in diabetes and hypertension prevention and management, raising awareness of common NCD risk factors, and providing diabetes and hypertension diagnostic and treatment services in the selected church health institutions.
WCEA	Cardio Diabetes	Malawi, Tanzania, Sierra Leone, Zimbabwe, Uganda	Online HCP Training	Online NCD training of healthcare professionals across multiple countries.
CNSS	Cardio Diabetes	Djibouti	Empowering HCPs and supply chain actors	The specific objectives of this partnership are focused on strengthening advocacy and knowledge about NCDs, increasing the capacity of healthcare professionals for better management of NCDs and of supply chain actors, while building a sustainable procurement mechanism for affordable access to treatment.
Touch Foundation	Cardio Diabetes	Tanzania	Strengthen Supply Chain	The primary goal is to improve supply chain management for NCD medicines and patient tracking at each facility to ensure patients are adhering to treatment.

Action 4 Diabetes (A4D)	Diabetes (type 1)	Cambodia, Laos, Myanmar	Care for Type 1 Diabetes Patients	Action 4 Diabetes focuses on type 1 diabetes patients and includes healthcare professional training, patient services, support in monitoring blood glucose levels and access to insulins, to increase efficiency in the management of type 1 diabetes patients. A4D also holds diabetes camps for patients and their families to build awareness and understanding.
City Cancer Challenge	Oncology	Cambodia Rwanda	Health System Strengthening	Working with City Cancer, the objectives are to create city-wide oncology stakeholder leadership groups and complete situational analysis and needs assessments of oncology services (including digital oncology services), forming the basis for a successful approach to empower and strengthen the health system.

### Cancer and work: Sanofi supporting health and wellbeing in the workplace

Sanofi has launched 'Cancer & Work: Acting Together', a program which covers all Sanofi employees in the world if they are diagnosed with cancer or critical illnesses<sup>1</sup>. It provides social, emotional and financial support and secures the job, salary and benefits of any employee for up to twelve months, no matter the role or geographical location.

It will allow employees to incorporate further flexible work arrangements to better navigate cancer and work and will have access to a network of volunteer colleagues trained to help them navigate from initial diagnosis through the treatment journey and return to work. The program is also designed to better equip managers to support members of their team who are affected by cancer. Throughout 2024, Sanofi also intends to implement coverage of miscellaneous non-medical expenses. Moreover, Sanofi permanent employees will become eligible for an unpaid caregiver leave which allows them to carry out caregiving duties for their close family member suffering from a critical illness<sup>1</sup>.

In 2017, several volunteer employees in France, with complementary expert skills and experience as patients, caregivers or managers, started the initiative. The program has since grown to a network of 27 partner teams with one team at each Sanofi site in France, with 150 members who share feedback and best practice. More than 350 employees have benefited (42% sick employees, 30% caregivers, 28% managers).

The program "Cancer & Work" has started to roll out globally in early 2024 and is part of our programs supporting health and wellbeing in the workplace. This complements other initiatives already launched for employees such as the gender-neutral parental leave, allowing all new parents 14 weeks of paid leave to welcome a new child into their lives.

1. Specific criteria identifying the conditions and circumstances that are eligible for coverage under this program might be governed by the terms and conditions of country-specific policies or legal requirements.

## ESG ratings

Latest Sanofi ESG rankings:

### Sanofi ESG ratings

#### Rating agencies

<b>SCORE</b>	<b>21.2</b> Medium risk	<b>79/100</b>	<b>87/100</b>	<b>Climate Change: A-</b> <b>Water: A-</b>	<b>B</b>	<b>4.5/5</b>	<b>3.47/5</b>	<b>65/100</b>
A Score stable since 2021	21.5 21st among 447 pharmaceutical companies	78/100 Percentile of 99 within 348 scored companies in the industry	New Disclosure score of 87/100 vs a 67/100 average for the healthcare sector 2023 WDI Awards Special mention for Workforce Action	A/A-	B 1st decile of the 476 companies in the industry	4.3/5 With very high rating across the 3 pillars ESG	3.47/5 Top 10 company	64/100 1st pharmaceutical company out of 57 Score improving since 2018

vs. previous rating

Scores assigned by the rating agencies are not equivalent.

## Q1 2024 financial results

### Business net income<sup>2</sup>

Sanofi generated **net sales** of €10,464 million, an increase of 2.4% (up 6.7% at CER).

**Other revenues** increased by 2.0% (up 9.8% at CER) to €654 million, including higher VaxServe sales of non-Sanofi products of €407 million (up 5.6% at CER).

**Gross Profit** decreased by 1.2% (up 4.2% at CER) to €7,694 million. The gross margin ratio decreased by 2.6pp to 73.5% (74.4% at CER) compared with Q1 2023. This decrease mainly reflected a deterioration of the Biopharma gross margin ratio (from 77.7% to 75.1%) due to the product mix evolution (e.g. sales decrease of Aubagio, COVID sales) and unfavorable currency evolution. CHC gross margin ratio decreased from 67.0% to 63.9% due to product and country mix, as well as unfavorable currency evolution.

**Research and Development** (R&D) expenses increased by 10.0% to €1,719 million. At CER, R&D expenses increased by 11.8%, reflecting increased expenses in Vaccines (mRNA) and Pharma (pipeline acceleration).

**Selling, general and administrative expenses** (SG&A) decreased by 0.1% to €2,605 million. At CER, SG&A expenses increased by 2.9%, reflecting increased commercial investments and launch costs in Biopharma and Vaccines as well as higher CHC commercial expenses. The ratio of SG&A to sales decreased by 0.6pp to 24.9% compared to Q1 2023.

**Operating expenses** were €4,324 million (up 3.7% and up 6.2% at CER).

**Other current operating income net of expenses** were -€562 million compared to -€304 million in Q1 2023. Other current operating income net of expenses included an expense of €825 million (compared to an expense of €674 million in Q1 2023) corresponding to the share of profit to Regeneron from the monoclonal antibodies alliance, the share of profit paid by Regeneron towards development costs and the reimbursement of commercialization-related expenses incurred by Regeneron. This line also included €199 million of capital gains from divestments of medicines/portfolio streamlining, compared to €307 million in Q1 2023. Sanofi expects the amount of capital gains from divestments of medicines/portfolio streamlining to exceed €500 million in 2024.

**Share of profit from associates** was €44 million compared to €33 million in Q1 2023 and included the share of US profit related to Vaxelis.

**Business operating income<sup>5</sup>** (BOI) decreased by 14.7% to €2,843 million. At CER, BOI decreased by 4.2%. The ratio of BOI to net sales decreased by 5.4pp to 27.2% (down 3.3pp at CER).

**Net financial expenses** were €43 million compared to €7 million in Q1 2023.

**The effective tax rate** increased to 21.0% from 19.0% in Q1 2023. Sanofi expects its effective tax rate to be around 21% in 2024.

**Business net income<sup>5</sup>** decreased by 17.8% to €2,219 million and decreased by 7.4% at CER. The ratio of business net income to net sales decreased by 5.2pp to 21.2% compared to Q1 2023 (down 3.5pp at CER).

**Business earnings per share<sup>5</sup>** (EPS) was €1.78, down 17.6% on a reported basis (down 7.4% at CER). The average number of shares outstanding was 1,248.8 million compared to 1,249.3 million in Q1 2023.

### Reconciliation of IFRS net income reported to business net income (see Appendix 4)

The IFRS net income was €1,133 million. The main items excluded from the business net income were:

- An amortization charge of €562 million related to fair value remeasurement on intangible assets: 547 million (primarily Bioverativ: €160 million, Genzyme: €94 million, Provention Bio: €53 million, Boehringer Ingelheim CHC business: €45 million, Ablynx: €42 million, Kadmon: €41 million, Beyfortus: €28 million and Qunol: €20 million) and to intangible assets from separate acquisitions - measured initially at acquisition cost (licenses/products): €15 million. These items have no cash impact.
- Restructuring costs and similar items of €740 million mainly related to redundancy plans announced during Q1 2024 and separation costs of the CHC business.
- Other gains and losses, and litigation charge of €79 million related to change in provision for litigation.

<sup>2</sup>See Appendix 3 for Q1 2024 consolidated income statement; see Appendix 7 for definitions of financial indicators, and Appendix 4 for reconciliation of IFRS net income reported to business net income.

- A financial charge of €59 million related to the remeasurement of expected future royalty of Beyfortus US sales.
- A €424 million tax effect arising from the items listed above, mainly comprising €105 million of deferred taxes generated by amortization of intangible assets and €272 million associated with restructuring costs and similar items (see Appendix 4).
- A loss of €81 million corresponding to the equity stake in EUROAPI.

## Capital allocation

Q1 2024 free cash flow before restructuring, acquisitions and disposals amounted to -€78 million, after negative change of net working capital (-€1,764 million), notably including the decrease of US rebates provisions (€980 million) following decision to cut Lantus list price effective January 1, 2024 and capital expenditures (-€512 million). After acquisitions<sup>3</sup> (-€253 million), proceeds from disposals<sup>4</sup> (€478 million) and payments related to restructuring and similar items (-€456 million), Q1 2024 **free cash flow**<sup>4</sup> was -€309 million. The net debt increased from €7,793 million on December 31, 2023 to €8,534 million on March 31, 2024 (amount net of €9,181 million cash and cash equivalents).

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<sup>3</sup> Not exceeding €500 million per transaction (inclusive of all payments related to the transaction).

<sup>4</sup> Non-IFRS financial measure (definition in Appendix 7).

## Forward-Looking Statements

*This press release contains forward-looking statements as defined in the Private Securities Litigation Reform Act of 1995, as amended. Forward-looking statements are statements that are not historical facts. These statements include projections and estimates and their underlying assumptions, statements regarding plans, business transformations, objectives, intentions and expectations with respect to future financial results, events, operations, services, product development and potential, and statements regarding future performance. Forward-looking statements are generally identified by the words "expects", "anticipates", "believes", "intends", "estimates", "plans", "potential", "outlook", "guidance" and similar expressions. Although Sanofi's management believes that the expectations reflected in such forward-looking statements are reasonable, investors are cautioned that forward-looking information and statements are subject to various risks and uncertainties, many of which are difficult to predict and generally beyond the control of Sanofi, that could cause actual results and developments to differ materially from those expressed in, or implied or projected by, the forward-looking information and statements. These risks and uncertainties include among other things, the uncertainties inherent in research and development, future clinical data and analysis, including post marketing, decisions by regulatory authorities, such as the FDA or the EMA, regarding whether and when to approve any drug, device or biological application that may be filed for any such product candidates as well as their decisions regarding labelling and other matters that could affect the availability or commercial potential of such product candidates, the fact that product candidates if approved may not be commercially successful, the future approval and commercial success of therapeutic alternatives, Sanofi's ability to benefit from external growth opportunities, to complete capital markets or other transactions and/or obtain regulatory clearances, risks associated with developing standalone businesses, risks associated with intellectual property and any related pending or future litigation and the ultimate outcome of such litigation, trends in exchange rates and prevailing interest rates, volatile economic and capital market conditions, cost containment initiatives and subsequent changes thereto, and the impact that pandemics, political disruption or armed conflicts or other global crises may have on us, our customers, suppliers, vendors, and other business partners, and the financial condition of any one of them, as well as on our employees and on the global economy as a whole. The risks and uncertainties also include the uncertainties discussed or identified in the public filings with the SEC and the AMF made by Sanofi, including those listed under "Risk Factors" and "Cautionary Statement Regarding Forward-Looking Statements" in Sanofi's annual report on Form 20-F for the year ended December 31, 2023. Other than as required by applicable law, Sanofi does not undertake any obligation to update or revise any forward-looking information or statements. All trademarks mentioned in this document are protected.*

## Appendices

- Appendix 1: Q1 2024 sales by medicine/vaccine/business and geographic region
- Appendix 2: Q1 2024 business net income statement
- Appendix 3: Q1 2024 consolidated income statement
- Appendix 4: Reconciliation of IFRS net income reported to business net income
- Appendix 5: Change in net debt
- Appendix 6: Currency sensitivity
- Appendix 7: Definitions of non-IFRS financial indicators
- Appendix 8: CSR Dashboards

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Appendix 1: Q1 2024 net sales by medicine/vaccine/business and geographic region

Q1 2024 (€ million)	Total Sales	% CER	% reported	United States	% CER	Europe	% CER	Rest of World	% CER
<b>Dupixent</b>	<b>2,835</b>	<b>+24.9%</b>	<b>+22.4%</b>	<b>2,030</b>	<b>+16.8%</b>	<b>371</b>	<b>+33.6%</b>	<b>434</b>	<b>+67.5%</b>
Nexviazyme/Nexviadyne	152	+96.3%	+87.7%	85	+53.6%	43	+147.1%	24	+287.5%
Sarclisa	106	+28.7%	+21.8%	49	+32.4%	31	+10.7%	26	+45.5%
Altuviiio	122	+12300.0%	+12100.0%	115	+11600.0%	—	0.0%	7	0.0%
Rezurock	93	+40.3%	+38.8%	84	+28.8%	5	+400.0%	4	0.0%
Cablivi	59	+3.4%	+1.7%	32	+14.3%	23	-14.8%	4	+66.7%
Xenpozyme	35	+100.0%	+94.4%	18	+80.0%	12	+50.0%	5	0.0%
Enjaymo	29	+87.5%	+81.3%	15	+66.7%	6	+500.0%	8	+50.0%
Tzield	10	0.0%	0.0%	10	0.0%	—	0.0%	—	0.0%
<b>Total Pharma launches</b>	<b>606</b>	<b>+90.5%</b>	<b>+84.8%</b>	<b>408</b>	<b>+99.0%</b>	<b>120</b>	<b>+45.1%</b>	<b>78</b>	<b>+141.0%</b>
Toujeo	321	+18.0%	+11.1%	56	-13.8%	119	+9.2%	146	+44.3%
Lantus	360	-15.4%	-19.5%	112	-13.6%	92	-5.2%	156	-21.0%
Lovenox	262	-13.9%	-18.9%	3	-25.0%	155	-11.5%	104	-16.6%
Plavix	238	+6.8%	+0.8%	2	0.0%	23	-4.2%	213	+8.1%
Fabrazyme	253	+7.7%	+2.8%	126	+3.3%	63	+1.6%	64	+23.0%
Myozyme	191	-13.6%	-16.2%	60	-13.0%	76	-21.1%	55	-3.1%
Alprolix	130	+6.4%	+4.0%	109	+2.8%	—	0.0%	21	+27.8%
Cerezyme	214	+23.0%	+9.2%	48	+4.3%	65	+8.3%	101	+42.7%
Aubagio	102	-74.7%	-75.7%	41	-85.0%	52	-56.7%	9	-50.0%
Praluent	121	+25.5%	+23.5%	—	-100.0%	83	+16.9%	38	+42.9%
Thymoglobulin	117	+12.8%	+7.3%	73	+7.2%	10	0.0%	34	+30.0%
Aprovel	105	-0.9%	-4.5%	—	-100.0%	18	-10.0%	87	+2.2%
Kevzara	87	+23.3%	+19.2%	45	+27.8%	29	+7.4%	13	+50.0%
Eloctate	86	-24.6%	-27.1%	62	-25.9%	—	0.0%	24	-21.2%
Multaq	78	-6.0%	-7.1%	70	-6.6%	3	0.0%	5	0.0%
Jevtana	68	-10.1%	-13.9%	47	-12.7%	2	-50.0%	19	+5.0%
Cerdelga	83	+15.1%	+13.7%	46	+15.0%	32	+10.3%	5	+50.0%
Aldurazyme	83	+12.8%	+6.4%	18	0.0%	23	+9.5%	42	+20.5%
Soliqua/iGlarLixi	58	-4.8%	-7.9%	20	-37.5%	11	+37.5%	27	+26.1%
Fasturtec	42	-6.7%	-6.7%	27	+3.7%	11	-15.4%	4	-40.0%
Mozobil	25	-61.2%	-62.7%	3	-92.9%	16	-5.9%	6	-12.5%
Others	1,140	-6.1%	-10.6%	86	-15.2%	335	-7.5%	719	-4.3%
Industrial Sales	157	+16.3%	+16.3%	1	0.0%	156	+23.6%	—	-114.3%
<b>Total other medicines</b>	<b>4,321</b>	<b>-7.9%</b>	<b>-12.1%</b>	<b>1,055</b>	<b>-24.1%</b>	<b>1,374</b>	<b>-5.5%</b>	<b>1,892</b>	<b>+1.6%</b>
<b>Pharma</b>	<b>7,762</b>	<b>+6.4%</b>	<b>+2.7%</b>	<b>3,493</b>	<b>+4.8%</b>	<b>1,865</b>	<b>+2.8%</b>	<b>2,404</b>	<b>+11.6%</b>
Influenza vaccines	73	+38.1%	+15.9%	5	-16.7%	1	-80.0%	67	+55.8%
Polio/Pertussis/Hib vaccines & Boosters	636	-0.5%	-3.8%	164	-15.3%	109	0.0%	363	+7.6%
RSV (Beyfortus)	182	0.0%	0.0%	114	0.0%	7	0.0%	61	0.0%
Meningitis, Travel and endemic vaccines	286	+7.7%	+5.1%	137	+14.0%	48	+38.2%	101	-7.7%
<b>Vaccines</b>	<b>1,177</b>	<b>+5.6%</b>	<b>+0.9%</b>	<b>420</b>	<b>+30.9%</b>	<b>165</b>	<b>-48.4%</b>	<b>592</b>	<b>+22.7%</b>
<b>Biopharma</b>	<b>8,939</b>	<b>+6.3%</b>	<b>+2.4%</b>	<b>3,913</b>	<b>+7.1%</b>	<b>2,030</b>	<b>-4.9%</b>	<b>2,996</b>	<b>+13.6%</b>
<b>Consumer Healthcare</b>	<b>1,525</b>	<b>+9.0%</b>	<b>+2.0%</b>	<b>403</b>	<b>+10.0%</b>	<b>450</b>	<b>0.0%</b>	<b>672</b>	<b>+14.3%</b>
<b>Company</b>	<b>10,464</b>	<b>+6.7%</b>	<b>+2.4%</b>	<b>4,316</b>	<b>+7.3%</b>	<b>2,480</b>	<b>-4.0%</b>	<b>3,668</b>	<b>+13.8%</b>



## Appendix 2: Business net income statement

Q1 2024	Biopharma			Consumer Healthcare			Other			Total Group		
	Q1 2024	Q1 2023	Change	Q1 2024	Q1 2023	Change	Q1 2024	Q1 2023	Change	Q1 2024	Q1 2023	Change
<b>€ million</b>												
<b>Net sales</b>	<b>8,939</b>	<b>8,727</b>	<b>2.4%</b>	<b>1,525</b>	<b>1,495</b>	<b>2.0%</b>	—	—	—%	<b>10,464</b>	<b>10,222</b>	<b>2.4%</b>
Other revenues	639	626	2.1%	15	15	—%	—	—	—%	654	641	2.0%
Cost of Sales	(2,862)	(2,569)	11.4%	(565)	(508)	11.2%	3	(2)	-250.0%	(3,424)	(3,079)	11.2%
As % of net sales	(32.0%)	(29.4%)		(37.0%)	(34.0%)					(32.7%)	(30.1%)	
<b>Gross Profit</b>	<b>6,716</b>	<b>6,784</b>	<b>-1.0%</b>	<b>975</b>	<b>1,002</b>	<b>-2.7%</b>	<b>3</b>	<b>(2)</b>	<b>-250.0%</b>	<b>7,694</b>	<b>7,784</b>	<b>-1.2%</b>
<b>As % of net sales</b>	<b>75.1%</b>	<b>77.7%</b>		<b>63.9%</b>	<b>67.0%</b>					<b>73.5%</b>	<b>76.1%</b>	
Research and development expenses	(1,675)	(1,510)	10.9%	(44)	(53)	-17.0%	—	—	—%	(1,719)	(1,563)	10.0%
As % of net sales	(18.7%)	(17.3%)		(2.9%)	(3.5%)					(16.4%)	(15.3%)	
Selling and general expenses	(2,093)	(2,124)	-1.5%	(514)	(484)	6.2%	2	1	100.0%	(2,605)	(2,607)	-0.1%
As % of net sales	(23.4%)	(24.3%)		(33.7%)	(32.4%)					(24.9%)	(25.5%)	
Other current operating income/expenses	(617)	(386)		54	71		1	11		(562)	(304)	
Share of profit/loss of associates* and joint ventures	38	30		6	3		—	—		44	33	
Net income attributable to non controlling interests	(4)	(5)		(5)	(5)		—	—		(9)	(10)	
<b>Business operating income</b>	<b>2,365</b>	<b>2,789</b>	<b>-15.2%</b>	<b>472</b>	<b>534</b>	<b>-11.6%</b>	<b>6</b>	<b>10</b>	<b>-40.0%</b>	<b>2,843</b>	<b>3,333</b>	<b>-14.7%</b>
<b>As % of net sales</b>	<b>26.5%</b>	<b>32.0%</b>		<b>31.0%</b>	<b>35.7%</b>					<b>27.2%</b>	<b>32.6%</b>	
										(43)	(7)	
										(581)	(627)	
										21.0%	19.0%	
										<b>2,219</b>	<b>2,699</b>	<b>-17.8%</b>
										<b>21.2%</b>	<b>26.4%</b>	
										<b>1.78</b>	<b>2.16</b>	<b>-17.6%</b>

\* Net of tax.

\*\* Determined on the basis of Business income before tax, associates, and non-controlling interests.

\*\*\* Based on an average number of shares outstanding of 1,248.8 million in the first quarter of 2024 and 1,249.3 million in the first quarter of 2023.

### Appendix 3: Consolidated income statements

€ million	Q1 2024	Q1 2023
<b>Net sales</b>	<b>10,464</b>	<b>10,222</b>
Other revenues	654	641
Cost of sales	(3,439)	(3,079)
<b>Gross profit</b>	<b>7,679</b>	<b>7,784</b>
Research and development expenses	(1,719)	(1,563)
Selling and general expenses	(2,605)	(2,607)
Other operating income	444	436
Other operating expenses	(1,006)	(740)
Amortization of intangible assets	(562)	(489)
Impairment of intangible assets	17	(15)
Fair value remeasurement of contingent consideration	20	(15)
Restructuring costs and similar items	(740)	(240)
Other gains and losses, and litigation	(79)	(88)
<b>Operating income</b>	<b>1,449</b>	<b>2,463</b>
Financial expenses	(253)	(168)
Financial income	151	161
<b>Income before tax and associates and joint ventures</b>	<b>1,347</b>	<b>2,456</b>
Income tax expense	(164)	(459)
Share of profit/(loss) of associates and joint ventures	(37)	12
<b>Net income</b>	<b>1,146</b>	<b>2,009</b>
Net income attributable to non-controlling interests	13	14
<b>Net income attributable to equity holders of Sanofi</b>	<b>1,133</b>	<b>1,995</b>
Average number of shares outstanding (million)	1,248.8	1,249.3
<b>IFRS Earnings per share (in euros)</b>	<b>0.91</b>	<b>1.60</b>

#### Appendix 4: Reconciliation of Net income attributable to equity holders of Sanofi to Business net income

€ million	Q1 2024	Q1 2023
<b>Net income attributable to equity holders of Sanofi</b>	<b>1,133</b>	<b>1,995</b>
Amortization of intangible assets <sup>(1)</sup>	562	489
Impairment of intangible assets	(17)	15
Fair value remeasurement of contingent consideration	(16)	19
Expenses arising from the impact of acquisitions on inventories	15	—
Restructuring costs and similar items	740	240
Other gains and losses, and litigation	79	88
Financial (income) / expense related to liabilities carried at amortized cost other than net indebtedness	59	—
Tax effect of the items listed above:	(424)	(173)
<i>Amortization and impairment of intangible assets</i>	(101)	(94)
<i>Fair value remeasurement of contingent consideration</i>	3	(5)
<i>Expenses arising from the impact of acquisitions on inventories</i>	(3)	—
<i>Restructuring costs and similar items</i>	(272)	(49)
<i>Other items</i>	(51)	(25)
Other tax effects	7	5
Other items	81	21
<b>Business net income</b>	<b>2,219</b>	<b>2,699</b>
<b>IFRS earnings per share <sup>(2)</sup> (in euros)</b>	<b>0.91</b>	<b>1.60</b>

(1) Of which related to amortization expense generated by the intangible assets measured at their acquisition-date fair values: €547 million in the first quarter of 2024 and €468 million in the first quarter of 2023.

(2) Based on an average number of shares outstanding of 1,248.8 million in the first quarter of 2024 and 1,249.3 million in the first quarter of 2023.

## Appendix 5: Change in net debt

€ million	Q1 2024	Q1 2023
<b>Business net income</b>	<b>2,219</b>	<b>2,699</b>
Depreciation & amortization & impairment of property, plant and equipment and software	349	370
Other items	(370)	(323)
<b>Operating cash flow</b>	<b>2,198</b>	<b>2,746</b>
Changes in Working Capital	(1,764)	(627)
Acquisitions of property, plant and equipment and software	(512)	(447)
<b>Free cash flow before restructuring, acquisitions and disposals</b>	<b>(78)</b>	<b>1,672</b>
Acquisitions of intangibles assets, investments and other long-term financial assets <sup>(1)</sup>	(253)	(210)
Restructuring costs and similar items paid	(456)	(333)
Proceeds from disposals of property, plant and equipment, intangible assets and other non-current assets net of taxes <sup>(1)</sup>	478	408
<b>Free cash flow</b>	<b>(309)</b>	<b>1,537</b>
Acquisitions of investments in consolidated undertakings including assumed debt <sup>(2)</sup>	(83)	—
Issuance of Sanofi shares	14	16
Acquisition of treasury shares	(302)	(363)
Other items	(61)	(58)
<b>Change in net debt</b>	<b>(741)</b>	<b>1,132</b>
<b>Beginning of period</b>	<b>7,793</b>	<b>6,437</b>
<b>Closing of net debt</b>	<b>8,534</b>	<b>5,305</b>

(1) Free cash flow includes investments and divestments not exceeding a cap of €500 million per transaction (inclusive of all payments related to the transaction).

(2) Includes transactions that are above a cap of €500 million per transaction (inclusive of all payments related to the transaction).

## Appendix 6: Currency sensitivity

### 2024 business EPS currency sensitivity

Currency	Variation	Business EPS Sensitivity
US Dollar	+0.05 USD/EUR	-EUR 0.17
Japanese Yen	+5 JPY/EUR	-EUR 0.02
Chinese Yuan	+0.2 CNY/EUR	-EUR 0.02
Brazilian Real	+0.4 BRL/EUR	-EUR 0.01
Russian Ruble	+10 RUB/EUR	-EUR 0.01

### Currency exposure on Q1 2024 sales

Currency	Q1 2024
US \$	42.6 %
Euro €	20.6 %
Chinese Yuan	6.6 %
Japanese Yen	3.7 %
Brazilian Real	2.6 %
Mexican pesos	1.8 %
Russian ruble	1.7 %
British Pound	1.4 %
Australian \$	1.3 %
Canadian \$	1.2 %
Others	16.5 %

### Currency average rates

	Q1 2023	Q1 2024	Change
€/\$	1.073	1.085	+1.2%
€/Yen	142.049	161.152	+13.4%
€/Yuan	7.349	7.821	+6.4%
€/Real	5.575	5.375	-3.6%
€/Ruble	78.351	98.637	+25.9%

## Appendix 7: Definitions of non-IFRS financial indicators

### Company sales at constant exchange rates (CER)

When we refer to changes in our net sales “at constant exchange rates” (CER), this means that we exclude the effect of changes in exchange rates.

We eliminate the effect of exchange rates by recalculating net sales for the relevant period at the exchange rates used for

the previous period.

### Reconciliation of net sales to Company sales at constant exchange rates for Q1 2024.

€ million	Q1 2024
<b>Net sales</b>	<b>10,464</b>
Effect of exchange rates	(444)
<b>Company sales at constant exchange rates</b>	<b>10,908</b>

### Business net income

Sanofi publishes a key non-IFRS indicator. Business net income is defined as net income attributable to equity holders of Sanofi excluding:

- amortization of intangible assets,
- impairment of intangible assets,
- fair value remeasurement of contingent consideration related to business combinations or to disposals,
- expenses arising from the impact of acquisitions on inventories
- restructuring costs and similar items<sup>(1)</sup>,
- other gains and losses (including gains and losses on disposals of non-current assets<sup>(1)</sup>),
- costs or provisions associated with litigation<sup>(1)</sup>,
- upfront payments and regulatory milestone payments recognized in the line item Other operating income and arising from transactions outside the scope of Sanofi's ordinary activities,
- financial (income)/expense related to liabilities carried at amortized cost other than net indebtedness,
- tax effects related to the items listed above as well as effects of major tax disputes,
- the share of profits/losses from investments accounted for using the equity method, except for joint ventures and associates with which Sanofi has a strategic alliance,
- net income attributable to non-controlling interests related to the items listed above.

*(1) Reported in the line items **Restructuring costs and similar items** and **Gains and losses on disposals, and litigation**, which are defined in Notes B.16. and B.17. to our consolidated financial statements.*

### Free cash flow

Free cash flow is a non-IFRS financial indicator which is reviewed by our management, and which we believe provides useful information to measure the net cash generated from Sanofi's operations that is available for strategic investments<sup>1</sup> (net of divestments<sup>1</sup>), for debt repayment, and for capital return to shareholders. Free cash flow is determined from the Business net income adjusted for depreciation, amortization and impairment, share of profit/loss in associates and joint ventures net of dividends received, gains & losses on disposals, net change in provisions including pensions and other post-employment benefits, deferred taxes, share-based expense and other non-cash items. It comprises net changes in working capital, capital expenditures and other asset acquisitions<sup>2</sup> net of disposal proceeds<sup>2</sup>, and payments related to restructuring and similar items. Free cash flow is not defined by IFRS and it is not a substitute measure for the IFRS aggregate net cash flows in operating activities.

<sup>1</sup> Amount of the transaction above a cap of €500 million per transaction (inclusive of all payments related to the transaction).

<sup>2</sup> Not exceeding a cap of €500 million per transaction (inclusive of all payments related to the transaction).

## Appendix 8: CSR dashboards

Data are presented in YTD unless stated otherwise.

Topic	Ambition	Progress	
<b>Affordable access</b>			
		Q1 2024	Q1 2023
Sanofi Global Health	Reach 1.5 million NCD patients by 2026 (cumulative since 2022) and 2 million by 2030	57,889 patients treated in 18 countries	54,396 patients treated in 19 countries
		44 active healthcare partnerships in 21 countries	13 active healthcare partnerships in 14 countries
		4 investments signed through the Impact Fund (cumulative since launch)	1 investment signed through the Impact Fund
		Q1 2024	Q1 2023
Viols donations	Donate 100,000 vials a year to treat people with rare diseases	1,112 patients treated 17,287 vials donated	1,065 patients treated 21,542 vials donated
		Q1 2024	Q4 2023
Global Access Plans	Develop a Global Access Plan for all new products to make them available within two years after first launch	10 Global Access Plans initiated or developed covering more than 14 indications	8 Global Access Plans initiated or developed covering more than 12 indications
<b>R&amp;D for unmet needs</b>			
		FY 2023	FY2022
Sleeping sickness	Develop and supply innovative treatments to support the elimination of sleeping sickness by 2030	Data updated annually, next update in Q2 2024	1.5 million patients tested 837 patients treated
		Q1 2024	Q1 2023
Polio	Provide inactivated polio vaccines (IPV) to UNICEF for GAVI countries to support polio eradication efforts	9.4 million IPV doses supplied to UNICEF for GAVI countries	7 million IPV doses supplied to UNICEF for GAVI countries
		Q1 2024	Q4 2023
Pediatric cancer treatment development	Develop innovative treatments to eliminate cancer death in children	3 assets undergoing pre-clinical assessment	3 assets undergoing pre-clinical assessment
		1 asset in clinical study	First pediatric patient dosed with 1 clinical asset in a clinical study (less than 2 years after the 1st adult patient was dosed with this compound)
<b>Planet Care</b>			
		Q1 2024	Q4 2023
Climate change - Carbon footprint (CO <sub>2</sub> emissions)	55% reduction in scope 1&2 greenhouse gas emissions (CO <sub>2</sub> equivalent) by 2030 (cumulative vs 2019 baseline) to contribute to carbon neutrality by 2030 and net zero emissions by 2045 (all scopes)	42% GHG reduction vs 2019	38% GHG reduction vs 2019
Renewable electricity	100% of renewable electricity in all our sites by 2030	84%	79%
Eco-car fleet	100% eco-car fleet in 2030	44% eco-car fleet	43% eco-car fleet
Blister free syringe vaccines	100% blister free syringe vaccines by 2027	Next update at Q4 2024	39% blister free syringe vaccines

Eco-design	All new products to be eco-designed by 2025	13 LCAs completed & 5 in progress (new and marketed products)	13 LCAs completed & 2 in progress (new and marketed products)
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**In and beyond the workplace**

		Q1 2024	Q4 2023
Global Gender balance	Ambition of 50% of women in senior leadership roles by 2025	45%	44%
	Ambition of 40% of women in executive roles by 2025	41%	40%
Engagement with communities	Engage socially and economically with all communities where we operate	Next update in Q2 2024	12,240 volunteers 75,376 hours
From Leaders to Citizens	100% of Sanofi leaders have CSR in their development path	70% of the leaders have completed the eLearning phase	71% of the leaders have completed the eLearning phase
		30% of the leaders have completed the full program	30% of the leaders have completed the full program

*End.*