

Forward looking statements

This document 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, 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" 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 related transactions and/or obtain regulatory clearances, 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 market conditions, cost containment initiatives and subsequent changes thereto, and the impact that COVID-19 will 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. Any material effect of COVID-19 on any of the foregoing could also adversely impact us. This situation is changing rapidly and additional impacts may arise of which we are not currently aware and may exacerbate other previously identified risks. 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, 2019. Other than as required by applicable law, Sanofi does not undertake any obligation to update or revise any forward-looking information or statements.



Agenda CMD21 - part 1

Driving growth with strategic choices	Paul Hudson	CEO
Focus on transformative therapies	Dietmar Berger	Head of Development, CMO
Leveraging our DNA to build our future	Olivier Charmeil Alexandre de Germay	Head of General Medicines Head of Global CV & EP Franchise
Unlocking value	Julie van Ongevalle Josephine Fubara	Head of Consumer Healthcare Chief Scientific Officer, CHC
Our people ambition	Natalie Bickford	Chief People Officer
Q&A (also joining)	Jean-Baptiste de Chatillon John Reed Arnaud Robert Cyril Grandchamp-Desraux Grace Gu Paul Martingell	CFO Head of R&D Chief Digital Officer Head of Diabetes Franchise CFO China General Medicines Head of CHC Global Growth Hub

Part 2 of Capital Markets Day to start at 3:45 CET / 9:45 EST





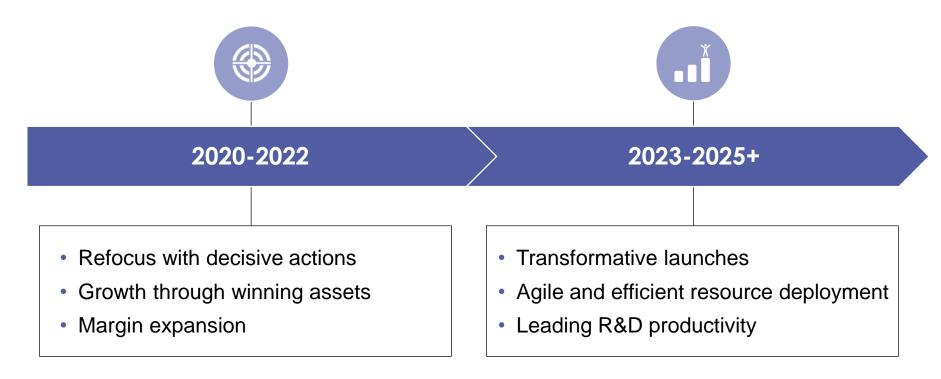
Driving growth with strategic choices

Paul Hudson

Chief Executive Officer



Play to Win: Our six-year plan – ahead of schedule





Our key growth drivers are delivering



Dupixent®

Maximize patient benefits with ambition to achieve >€10 billion peak sales across type 2 inflammatory diseases

€3.5bn sales in 2020, 3 years

after launch



Vaccines

Expected mid-to-high single-digit growth⁽¹⁾, through differentiated products, market expansion, launches

8.8% growth in 2020



Pipeline

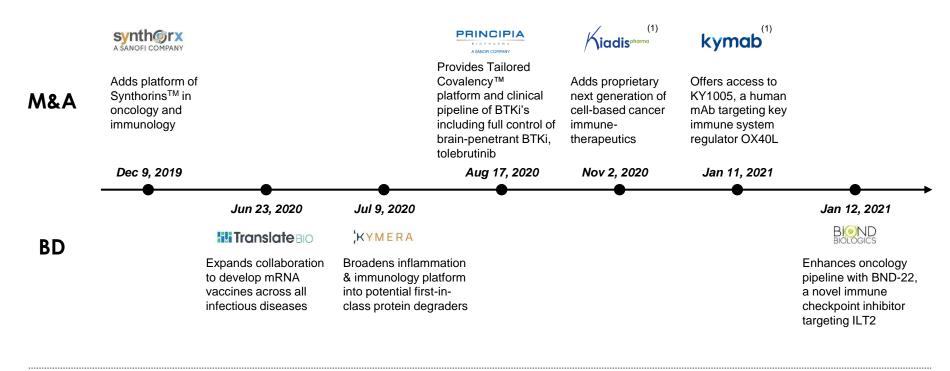
Prioritize and accelerate portfolio of potentially transformative therapies

12

projects entered Phase 3 in 2020



Clear capital allocation priorities to strengthen R&D



Sale of equity investment in Regeneron to support execution of 'Play to Win' strategy



EUROAPI to become a leading European company providing active pharmaceutical ingredients

New industry champion with six European manufacturing sites

- Expected sales of ~€1 bn by 2022, world #2
- ~3,200 employees headquartered in France
- CEO appointed Karl Rotthier, with ~30 years international experience
- IPO on Euronext Paris by 2022⁽¹⁾
- Debt-free at inception
- Sanofi to hold minority stake of ~30%

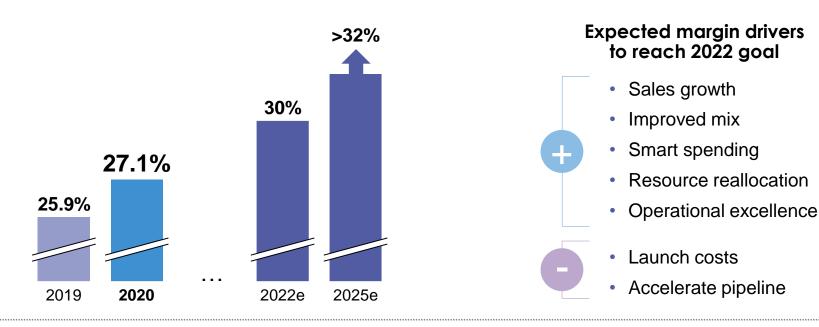


Carve-out activities on track



BOI margin up 120bps in 2020 tracking toward 2022 target

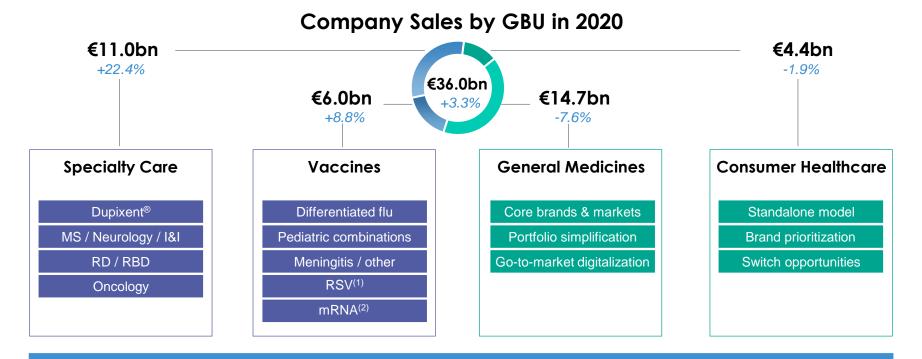
Sanofi expected BOI margin evolution



Dupixent® to become accretive to BOI margin by end of 2022



Driving growth with strategic choices



Reallocating to fund core drivers



⁾ In collaboration with AstraZeneca



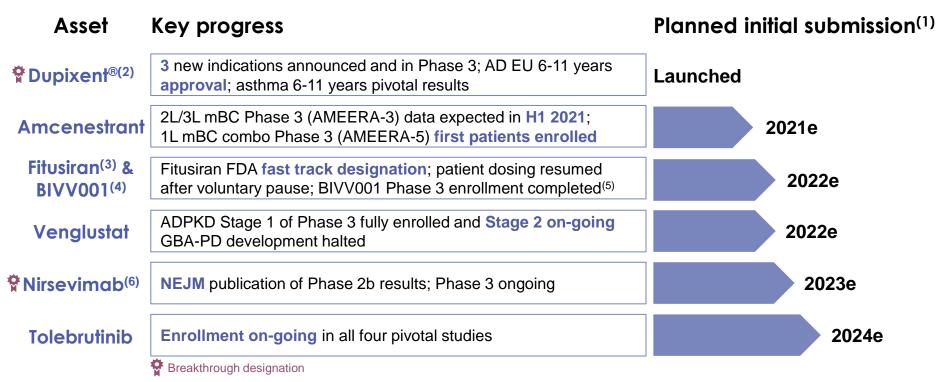
Focus on transformative therapies – status update

Dietmar Berger

Chief Medical Officer



Status update of our late-stage priority assets



Investigational uses of priority assets have not been approved by regulators for the uses being investigated.

AD: moderate to severe atopic dermatitis; mBC: metastatic breast cancer; ADPKD: autosomal dominant polycystic kidney disease; GBA-PD:

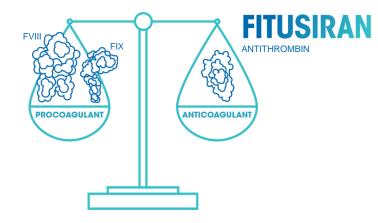


(1) First submission for assets with multiple potential indications; (2) Breakthrough designation for AD 6-11 years. Dupixent® in collaboration with Regeneron; (3) Fitusiran 2022 submission subject to future discussion with regulators (4) BIVV001 in collaboration with Sobi, recommended INN: efanesoctocog alfa; (5) Enrollment completed to meet the end of study criterion: (6) In collaboration with AstraZeneca

Fitusiran pivotal trial dosing resumed – new dose and regimen potentially strengthening target profile

- Fitusiran dosing resumed⁽¹⁾ in January, less than 3 months following voluntary pause
- Amendments to protocol to maintain antithrombin levels in target corridor
 - Starting dose lowered (~40%)
 - Starting dosing interval extended (every other month)

Fitusiran has the potential to transform the treatment of Hemophilia



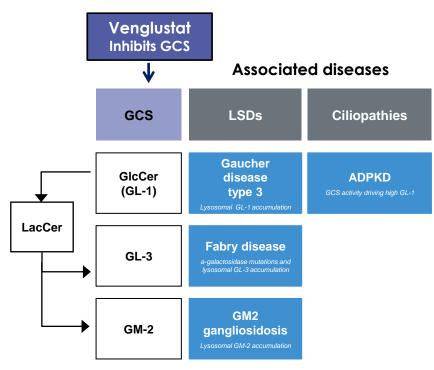
Potential first-in-class siRNA therapy

Next steps of regulatory path to be discussed with health authorities in H1 2021



Venglustat – a central regulator of GSL pathway, potentially transformative across multiple rare diseases

- Next-generation oral, brain-penetrating GCS inhibitor
 - PoC achieved in Gaucher Type 3 and Fabry Diseases
- Development in Parkinson's Disease halted following MOVES-PD Ph2 readout in GBA-PD end January
 - Did not meet primary or secondary efficacy endpoints; one-year trend favors placebo
 - Biomarkers indicated consistent, predictable GL-1 reductions in both plasma and cerebrospinal fluid
 - Continued favorable safety profile
- Venglustat development continues in ADPKD and LSDs
 - Regulates GSL metabolism linked to cystogenesis in ADPKD
 - SRT with GCS inhibitors established in GD Type 1



Pivotal data in ADPKD expected in H1 2022







Leveraging our DNA to build our future General Medicines

Olivier Charmeil

Head of General Medicines

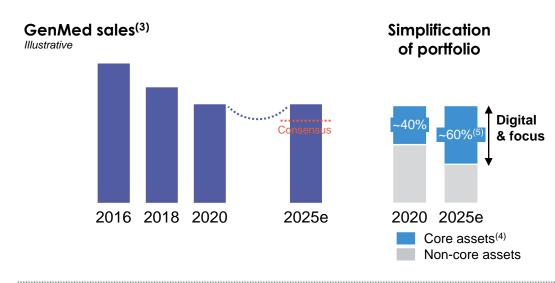
Alexandre de Germay

Head of Global CV & EP Franchise



Actively managing an accretive⁽¹⁾ and resilient business

Stabilize sales and maintain current BOI ratio⁽²⁾



Volume & price

- 2018-2020 decline mostly driven by price vs. stable volumes
- Portfolio remains critical for chronic disease management

Drivers to maintain BOI margin

- Core assets expected to grow to ~60%⁽⁵⁾ of sales by 2025
- Focus on key markets
- Continued divestments
- COGS improvement

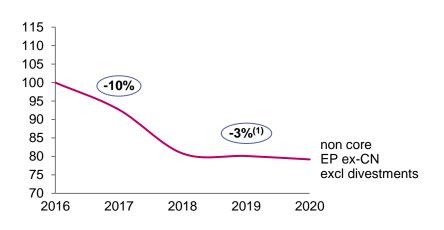
Funding the Specialty Care pipeline



- (1) GenMed BOI ratio expected to be accretive to Sanofi BOI ratio over the period
- (2) General Medicine BOI margin in 2020-2025
- (3) Excluding Industrial Affairs third party sales of which EUROAPI is expected to be deconsolidated in 2022
- (4) Core assets include Toujeo, Soliqua, Praluent, Multaq, Lovenox, Plavix and others for a total of €5.6bn in 2020
- (5) At CER

Overall, volumes of off-patent EP stabilizing with core assets still benefiting from strong demand in key EMs

Sanofi EP non-core volume stabilization at low price level



Lovenox® & Plavix®: Capturing class market growth opportunities in Emerging Markets and China





Source: Sanofi internal data based on ex-factory sales, normalized at 100 from 2016;

- (1) After correction from estimated COVID effect in 2020, and excl. divestments
- (2) IQVIA volume data MAT Oct 2020, excluding Turkey
- (3) IQVIA volume data: syringes per capita consumption

- (5) IQVIA treatment days MAT Oct 2020
- (6) United Nations, Department of Economic and Social Affairs, Population Division (2019)

Cardiometabolic diseases: Differentiated growth drivers to address patient needs









Patent Differentiation Geographic expansion

- Only PCSK9i associated with reduction in all-cause mortality(12)
- 1st line & new class 1A rec. in auidelines(4)

Only AAD to reduce CV

hospitalization

- Anticipated further differentiation vs. Degludec insulin with InRange clinical study(9)
- Anticipated differentiation vs. Premix insulins with Solimix clinical study(10)

- Rapid China uptake leveraging ACS leadership
- Germany to re-launch, largest potential in EU
- · China launch planned late 2021 / early 2022
- Sanofi leading BI market⁽⁶⁾ and Toujeo next generation BI market⁽⁷⁾
- · Toujeo MAX launch and broadest commercial coverage in U.S.(11)
- Double-digit growth EU⁽⁸⁾

- Expansion in countries with high premix use
- ~70% premix volume in China & emerging markets(6)

Market expansion

- PCSK9i market acceleration +52%(1)
- Favorable EU guidelines⁽²⁾
- · New evidence for earlier rhvthm control⁽⁵⁾
- Stabilized market share U.S.(3)
- Launched in China in 2020.
- Further expansion planned in key emerging markets: India, Algeria, Turkey in 2021
- >50% insulin initiation with premix in China
- Focus in U.S. on post-OAD HbA1c >9%

Focus on 13 key markets and digitally-enabled go-to-market model

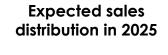
- (1) IQVIA treatment days data MAT Oct 2020, excluding US, Germany and Japan
- (2) 2019 ESC/EAS Guidelines for the management of dyslipidemias
- (3) U.S. TRx Nov 2020
- (4) 2020 ESC Guidelines for the diagnosis and management of atrial fibrillation
- Kirchhof P et al. N Engl J Med. 2020 Oct 1;383(14):1305-1316
- IQVIA sales data MAT Oct 2020
- (7) IQVIA sales data MAT Oct 2020

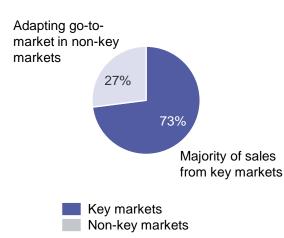
- (8) IQVIA sales data MAT Oct 2020 vs 2019
- (9) https://clinicaltrials.gov/ct2/show/NCT04075513; results expected H2 2021
- (10) https://www.clinicaltrialsregister.eu/ctr-search/search?query=LPS15017: results expected mid 2021
- (11) Toujeo® returning to coverage with CVS in 2021 Sanofi estimate of total lives covered
- (12) with only nominal statistical significance by hierarchical testing (HR 0.85, 95% CI 0.73,



Critical enablers to transform GenMed

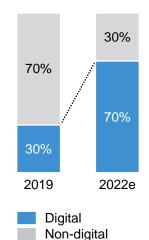
Focus on 13 key markets





Interacting digitally with our customers

HCP interactions



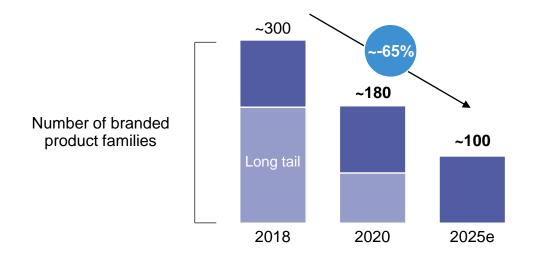
2020 status

- Digital transformation (accelerated by COVID)
 - ePermission for ~60% of our targeted physician universe
 - Digital content centralized production increased 400% in 2020 with increased standardization and acceptance⁽¹⁾
- OPEX reduction 2019-2020 of ~20%



Executing on streamlining the tail ahead of target

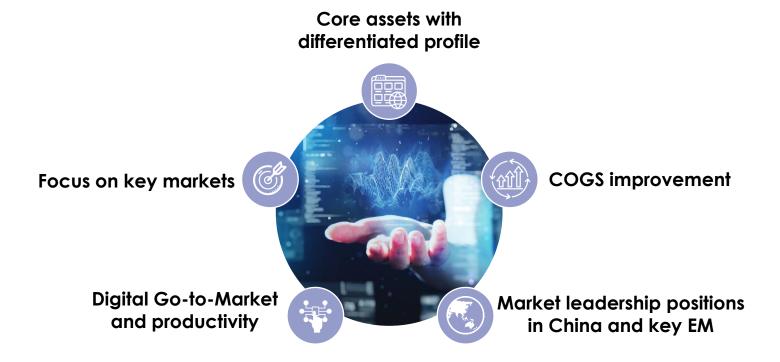
Simplification drives COGS improvement



~€1.5bn cash proceeds expected in 2019-2025



General Medicines: Accretive and resilient contributor to group BOI margin









Unlocking value Consumer Healthcare

Julie Van Ongevalle

Head of Consumer Healthcare

Josephine Fubara
Chief Science Officer, CHC



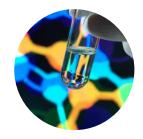
Sanofi's Consumer Healthcare 'Play to Win' strategy



Focus on growth

Brand prioritization

Acceleration in U.S. and China



Lead with innovation

Reignite innovation engine

Deliver switches



Accelerate efficiency

Simplify portfolio

Optimize go-tomarket model



Reinvent how we work

Become a fast-moving consumer healthcare company

Build digital and data edge

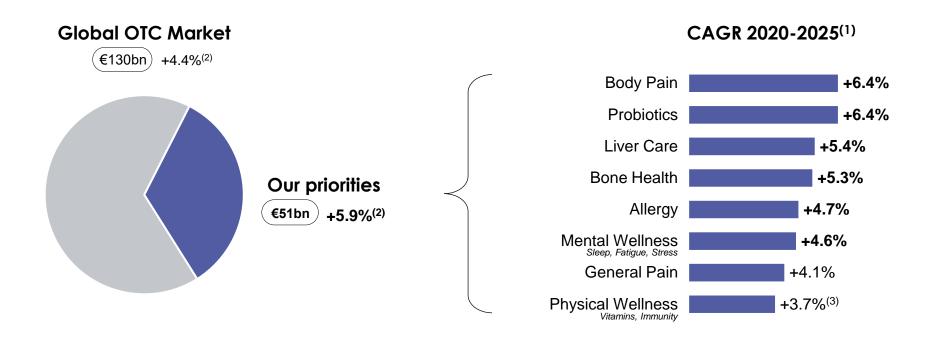
Sustainability at the core

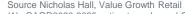
Ambition to be the fastest growing Global CHC company⁽¹⁾



(1) From 2024, including switches

Global OTC market is growing at 4.5% CAGR 2020-2025⁽¹⁾





⁽¹⁾ CAGR2020-2025 estimates released Q4 2020

SANOFI 👣

⁽²⁾ Actuals MAT Q3 2020

^{(3) 3.7%} growth in Physical Wellness (below market) represents only the retail growth. VMS (Vitamins, Minerals and Supplements) represents over 60% of e-commerce (Source NH) and grew +25% in 2020 Vs PY

Growth driven by local, regional and global love brands



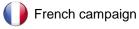




Strengthening our presence in Mental Wellness

- Mental Wellness expected to grow 4.6% over next 5 years
- Sleep represents ~40% of Mental Wellness
 - Double-digit growth in 2020 enhanced by COVID-19
 - Expected to grow 5.3% over the next 5 years
- Established local positions in:
 - U.S. Unisom
 - Europe NOVANUIT
 - Japan FUIII





'No matter what. Sleep at night.'



Standalone CHC on track to be in place end of 2022

Implementation ongoing



Design and overall planning complete



Relevant social processes on schedule



Operationalize majority of standalone CHC legal entities by end of 2021

Benefits to growth



Consumer and data at the core of every business decision



Reignite our innovation engine and reduce time-to-market by around 20%



Fully integrated R&D, manufacturing and supply network with dedicated support functions



First-in-class U.S. switches with €1bn U.S. sales potential

Cialis[®]

- >30m⁽¹⁾ U.S. men suffer from erectile dysfunction (ED)
- U.S market leader as Rx product (Lilly)
- Anticipate being the first OTC treatment for ED (PDE5i)
- Works up to 36 hours

Label comprehension, selfselection, human factors

Program underway

First patient enters Actual Use Trial this year

Expected launch

2024

Tamiflu®

- **30m**⁽²⁾ people suffer from flu in the U.S. annually
- U.S. market leader as Rx product (Roche)
- Anticipate being the first OTC flu antiviral
- Significantly reduces duration & severity of illness

Label comprehension, self-selection, human factors

Program underway

First patient enters Actual Use Trial in 2022

Expected launch

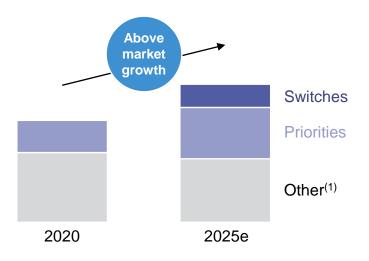
2023/

2024



⁽¹⁾ Burnett AL, Nehra A, Breau RH, Culkin DJ, Faraday MM, et al. Erectile Dysfunction. American Urological Association. Available at https://www.auanet.org/guidelines/male-sexual-dvsfunction-erectile-dvsfunction-(2018)

Invest in brands and geographies of focus for best-in-market growth in 2024/2025



- Ambition for best-in-market growth with switches from 2024/2025
- Grow priority brands above market growth as early as 2022 in key geographies
- Supported by consumer insights, digital and e-commerce channels and a standalone model with dedicated support functions



(1) Also includes brands to be divested 29



Our people ambitions

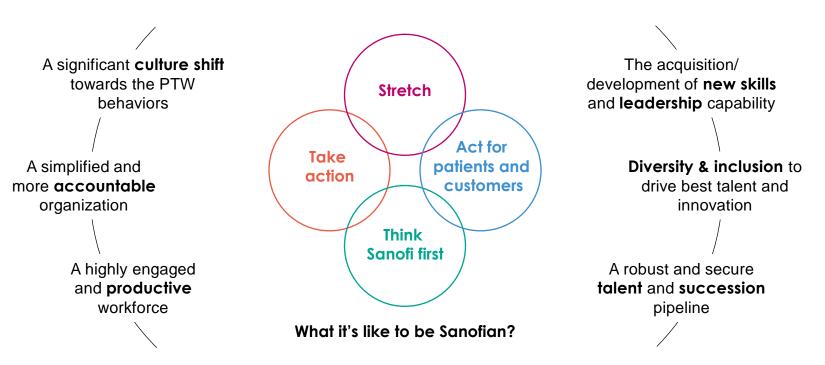
Natalie Bickford

Chief People Officer



Our business imperative

What we need to successfully deliver the 'Play to Win' strategy





Our 'Play to Win' 2025 people ambitions



Healthy organization

Sanofi is an agile and competitive organization meeting patient and market needs, with a robust talent pool and the right capabilities



Purposeful experience

Employees own their career journey, stretching themselves through compelling work experiences



Winning culture

Our culture allows our people to thrive and enables business success



Diversity edge

Our business outperforms through our ability to fully leverage the diversity of our people and our partners



Q&A session



Paul Hudson CEO



Olivier Charmeil
Head of General Medicines



Julie van Ongevalle Head of Consumer Healthcare



Arnaud RobertChief Digital Officer



Josephine Fubara Chief Scientific Officer, CHC



Cyril Grandchamp-Desraux Head of Diabetes Franchise



Paul Martingell
Head of CHC Global Growth Hub



Natalie Bickford Chief People Officer



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John Reed Head of R&D



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Grace GuCFO China General Medicines

