GENERAL 2018 MEETING 2018

May 2, 2018







Introduction

Serge Weinberg Chairman of the Board of Directors

Forward Looking Statements

This presentation contains forward-looking statements as defined in the Private Securities Litigation Reform Act of 1995, as amended. Forwardlooking 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 absence of guarantee that the product candidates if approved will be commercially successful, the future approval and commercial success of therapeutic alternatives, Sanofi's ability to benefit from external growth opportunities 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 conditions, the impact of cost containment initiatives and subsequent changes thereto, the average number of shares outstanding as well as those 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, 2017.

Other than as required by applicable law, Sanofi does not undertake any obligation to update or revise any forward-looking information or statements.



Agenda

Introduction & Governance

Serge Weinberg, Chairman of the Board of Directors

Compensation Policy

Patrick Kron, Chairman of the Compensation Committee

Strategic Outlook

Olivier Brandicourt, Chief Executive Officer

Progress in Research & Development

Elias Zerhouni, President, Global R&D

2017 Financial Performance

Jérôme Contamine, Executive Vice-President, Chief Financial Officer

Reports by Statutory Auditors at the Combined General Meeting

PricewaterhouseCoopers Audit ERNST & YOUNG et Autres

Questions & Answers

Vote on the Resolutions







Governance

Serge Weinberg Chairman of the Board of Directors

An Independent, Experienced and Diversified Board of Directors

Board of Directors composed of 16 directors

- Widely independent (11 out of 14⁽¹⁾) and gender sensitive (43%⁽²⁾)
- International (6 non-French directors, i.e. 38%)
- 2 directors representing employees

Further implementation of roadmap with a controlled and progressive renewal

- More scientific and pharmaceutical expertise
- Development of competences in the digital area
- Preservation of key competences
- Restricted number of directorships allowing an enhanced participation and dedication



Renewal of the terms of:

- Olivier Brandicourt
- Christian Mulliez
- Patrick Kron

Appointment of:

Emmanuel Babeau



¹⁾ Numbers not taking into account directors representing employees, pursuant to the recommendations of the AFEP-MEDEF Corporate Governance Code
2) Percentage not taking into account directors representing employees, pursuant to the relevant regulation

A Dedicated and Active Board of Directors

In 2017

- A sustained level of activity: 9 meetings
- Activities related to:
 - financial statements and financial matters
 - compensation matters
 - appointment and governance matters
- Review of significant proposed alliances, acquisitions and strategic opportunities
- Review of key businesses
- Evaluation of the activities of the Board and its Committees
- 3 executive sessions

High directors' attendance rate

> 95 %



Four Specialized Committees

1 Audit Committee

- Chaired by Robert Castaigne⁽¹⁾
- 3 financial experts
- 3 independent members out of 4
- 7 meetings in 2017
 - Review of the annual, half-year and quarterly financial results
 - Review of the main risks that may affect the results
 - Review of the draft financial resolutions
 - Update on the implementation of IFRS 15 and IFRS 9 accounting standards

Attendance rate: 93%

2 Compensation Committee

- Chaired by Patrick Kron
- 3 independent members out of 4
- 3 meetings in 2017
 - Review of the fixed and variable compensation of the CEO, the Chairman of the Board, and the members of the Executive Committee
 - Determination of the amount of directors' attendance fees
 - Update on changes in « say on pay » requirements
 - Launch of a new employee share-ownership plan

Attendance rate: 100%



Four Specialized Committees (cont'd)

3 Appointments and Governance Committee

- Chaired by Serge Weinberg
- 3 independent members out of 3
- 3 meetings in 2017
 - Succession planning
 - Follow-up of the discussions held with main shareholders and proxy advisors on governance matters
 - Evaluation of the activities of the Board and its Committees
 - Changes in the Board's composition

Attendance rate: 100%

4 Strategic Committee

- Chaired by Serge Weinberg
- 2 independent members out of 4
- 10 meetings in 2017
 - Review of external growth opportunities
 - Review of the strategy
 - Global environment
 - · Research and development
 - Long range financials

Attendance rate: 100%



A Fifth Specialized Committee in 2018



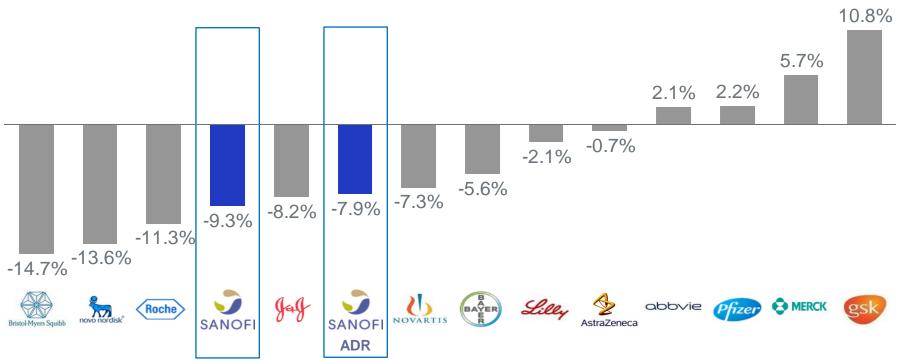
5 Scientific Committee

- Decision of the Board of Directors on March 6, 2018, upon recommendation of the Appointments and Governance Committee
- Scientific Committee in charge of assisting the Board on the strategic orientations in the field of R&D
- Thomas Südhof appointed Chairman of the Committee



Sanofi Share Performance vs. Pharma Company Peers since January 2018

Stock Performance in local currencies





Dividend: A Crucial Element of the Return to Shareholders



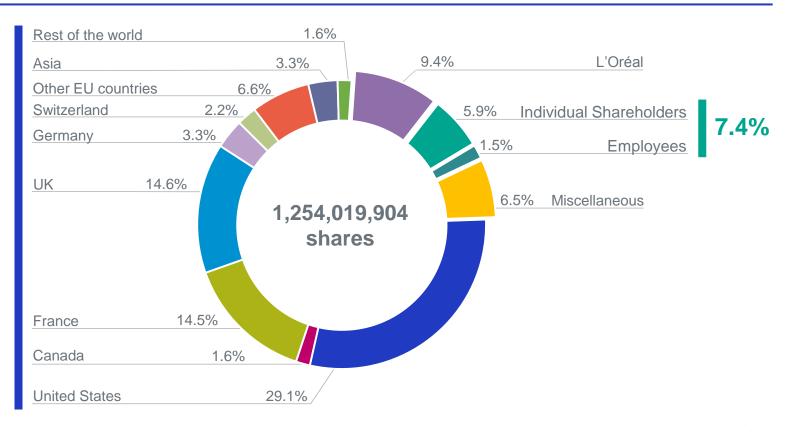


- 2017 dividend proposed by the Board of Directors: €3.03 per share⁽¹⁾
- 24th consecutive year of dividend increase

An International and Diversified Shareholder Base

Institutional Investors

76.7%





Strenghten Employee Share Ownership

3 initiatives since 2013

Action 2013

>1.6 million shares subscribed

Action 2016

>1.7 million shares subscribed

Action 2017

>1.5 million shares subscribed

Sanofi wants to strenghen its employee share ownership

- 22nd resolution voted by the AGM of May 10, 2017
- For an implementation in June 2018
- In more than 70 countries







Compensation Policy

Patrick Kron
Chairman of the Compensation Committee

Compensation Policy for the Chairman of the Board

Stable annual fixed gross compensation

- No attendance fees
- No annual variable compensation
- No equity compensation
- No compensation payable upon termination of office
- No pension entitlement
- No exceptional compensation



Compensation Elements of Serge Weinberg in 2017

	Amounts	Comments
Fixed compensation	€700,000	Fixed compensation unchanged since his appointment on May 17, 2010
	€8,353	Company car
Total	€708,353	



Compensation Policy for the Chief Executive Officer

- Annual fixed compensation
- Annual variable compensation
- Equity compensation
 - Options to subscribe for shares
 - Performance shares

- No Director's attendance fees
- No exceptional compensation

- Compensation on leaving office
 - Termination benefit
 - Top-up pension benefit subject to fulfillment of a performance condition (voluntary implementation of the Macron law as of January 2017)
 - Non-compete undertaking



Compensation Elements of Olivier Brandicourt in 2017

	Amounts	Comments
Fixed compensation	€1,200,000	Same compensation since his appointment in 2015
Variable compensation	€1,792,800 ⁽¹⁾	From 0% to 250% of the target fixed compensation to 150% of the fixed compensation Achieved rate: 149.4% of fixed compensation
Options	220,000 options to subscribe for shares	Exercise price: €88.97
Performance shares	50,000 performance shares	
Benefits in kind	€318	Social benefit in relation to social contribution payment made by Sanofi on CEO's behalf



Variable Compensation of Olivier Brandicourt in 2017

40% based on financial indicators

- 60% based on specific individual objectives
 - External growth (14%)
 - Excellence of product launches (10%)
 - Operational transformation (12%)
 - Organization and staff relations (12%)
 - New product pipeline (12%)

 Quantitative criteria account for 76% of the overall annual gross variable compensation objectives



99.6% of the 150% target, *i.e.* 149.4% of the fixed compensation



Equity Compensation of Olivier Brandicourt in 2017

2017 Grant

- 220,000 options to subscribe for shares
- 50,000 performance shares

2015 Plan

- Allocation rate of 81.12%
- Definitive acquisition in June 2019



Plan Conditions

- Performance conditions measured over 3 years
- 3 performance criteria
 - Business Net Income (50%)
 - Return On Assets or ROA (30%)
 - Total Shareholder Return or TSR (20%)

Plan Conditions

- Performance conditions measured over 3 years (2015-2017)
- 3 performance criteria
 - Business Net Income (50%) 102.2%
 - Return On Assets or ROA (30%) 100%
 - Total Shareholder Return or TSR (20%) 0%









Strategic Outlook

Olivier Brandicourt Chief Executive Officer

Sanofi, a Health Journey Partner







For 3 Years: Strategic Transformation for Better Healthcare and Continued Value Creation

- An unprecedented effort in Research & Development
- 2 Refocusing on our strenghs
- 3 A new operational model
- 4 Impact of digital opportunities





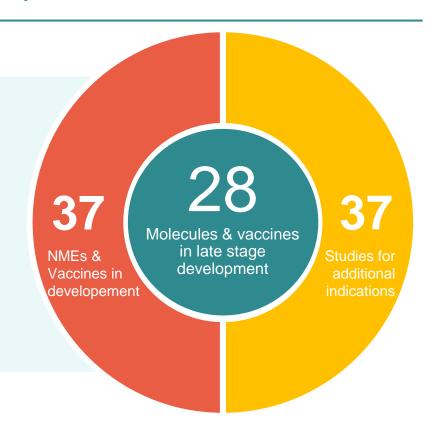
A Robust R&D Pipeline in 6 Therapeutic Areas

• €5.5bn invested in R&D in 2017

15.6% of company sales

74 projects in development

- Immunology
- Multiple Sclerosis & Neurology
- Oncology
- Rare Diseases & Rare Blood Disorders
- Diabetes & Cardiovascular
- Vaccines





Dupixent® Launched in Atopic Dermatitis A Product with Multiple Potential Indications

Moderate-to-Severe Atopic Dermatitis

- Strong launch in the U.S.
- Ongoing or planned launches in 20 countries in the coming months
- Moderate-to-Severe Asthma
 - Regulatory submissions in the U.S. and Europe
- Significant commercial potential in multiple diseases⁽¹⁾
 - Nasal Polyposis, Eosinophilic Esophagitis, Chronic Obstructive Pulmonary Disease, Allergies...





Pictures of a patient from a Phase 3 clinical trial before and after treatment with dupilumab. Results may vary

Two Promising Products in Oncology



cemiplimab

- PD-1 inhibitor monoclonal antibody
- Being evaluated by European and U.S. health authorities in Cutaneous Squamous Cell Carcinoma
- Phase 3 studies ongoing in other cancers

isatuximab

- Anti-CD38 monoclonal antibody
- Pivotal studies in Multiple Myeloma (bone marrow cancer) ongoing; submission planned in 2018

Praluent® Significantly Reduces Risk of Cardiovascular Events in High-Risk Patients





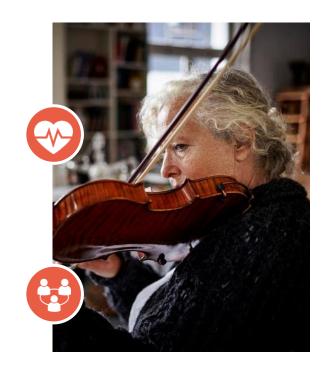


A more pronounced effect observed in patients with baseline LDL-C levels ≥ 100 mg/dL



Approved in more than 60 countries







(2) HR=0.85; IC: 0.73-0.98, nominal p value = 0.026

For 3 Years: Strategic Transformation for Better Healthcare and Continued Value Creation

- An unprecedented effort in Research & Development
- 2 Refocusing on our strenghs
- 3 A new operational model
- 4 Impact of digital opportunities





Consolidating our Leadership in Rare Diseases

Building a center of expertise in rare blood disorders



Three complementary moves



Restructuring of the alliance with Alnylam

 Global rights obtained on fitusiran, currently in Phase 3 in hemophilia





2 Refocusing on our strenghs

Bioverativ: a Leading Hemophilia Portfolio, a New Platform in Other Rare Blood Disorders





U.S. biotechnology company

• 2017 Revenues: \$1,168m, +31.7%

Leadership position in the large hemophilia market

Two marketed products





R&D programs

 Cold agglutinin disease, hemophilia and other rare blood disorders 2 Refocusing on our strenghs

Ablynx: Provides a Leading Technology Platform and Strengthens R&D Pipeline⁽¹⁾





Belgian biotechnology company

 An innovative Nanobody® platform which strengthens Sanofi's multi-targeting R&D strategy

Expanding rare blood disorders franchise(2)

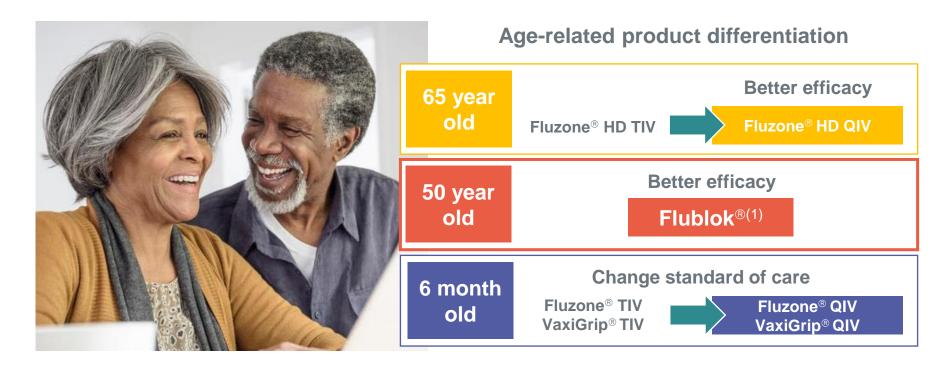
 Caplacizumab: submitted in Europe for the treatment of Acquired Thrombotic Thrombocytopenic Purpura

Complementary R&D programs

Hematology, inflammatory diseases, immuno-oncology, respiratory diseases

2 Refocusing on our strenghs

Flu Vaccines: Protein Sciences Broadens our Portfolio with Flublok®





For 3 Years: Strategic Transformation for Better Healthcare and Continued Value Creation

- An unprecedented effort in Research & Development
- 2 Refocusing on our strenghs
- 3 A new operational model
- 4 Impact of digital opportunities





An Organization Based on Five Global Business Units (GBUs)



For 3 Years: Strategic Transformation for Better Healthcare and Continued Value Creation

- An unprecedented effort in Research & Development
- 2 Refocusing on our strenghs
- 3 A new operational model
- 4 Impact of digital opportunities





Digitalization Offers Enormous Potential

- Accelerating research
- Better diabetes management, beyond medicines
- Better interaction with patients, physicians and payers
- Digitalization of the industrial tool





A diversified and strong model based on three categories of activity



High scientific content Strong added value

Diabetes and cardiovascular disease

Predictable and profitable growth



Sanofi Genzyme – Specialty Care Success of the New Immunology Franchise



Immunology

- Successful launch of Dupixent[®]
- Kevzara[®] launch progressing well

Multiple Sclerosis

Strong, growing franchise

Rare Diseases

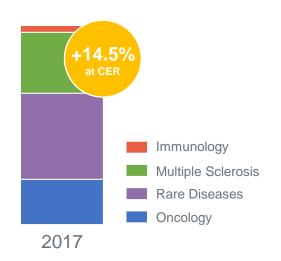
 Bioverativ acquisition and Ablynx planned acquisition strenghten leadership position⁽¹⁾

Oncology

 Significant expansion of development pipeline

Global Specialty Care Franchise Sales €6,678m

19% of company sales





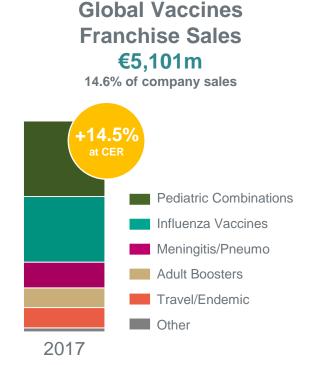
Sanofi Pasteur – Vaccines A Global Leader in a Growing Market



Leading vaccines in 5 areas:

- Influenza
- Pediatric Combinations
- Meningitis
- Adult Boosters
- Travelers and other vaccines for endemic countries

A strenghtened product portfolio with the acquisition of Protein Sciences





CER = Constant Exchange Rates

Diabetes and Cardiovascular Diseases



Diabetes Franchise

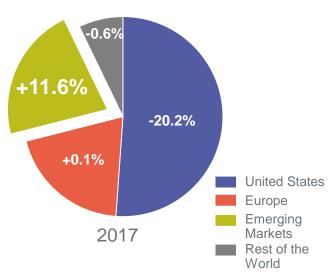
- Global sales down 11.1% at CER in 2017
 - Decrease in U.S. sales partly offset by performance in Emerging Markets
 - Success of Toujeo[®] in 2017: sales up 27% to €816m

Cardiovascular Franchise

 Praluent[®]: positive results of ODYSSEY OUTCOMES

Global DCV Franchise Sales €6,905m

19.7% of company sales





Sanofi is One of the Top 3 Players in Consumer Healthcare⁽¹⁾



Integration of Boehringher Ingelheim CHC business completed



Allergies,

Digestive

Other

Nutritionals

Pain











Doliprane













Global CHC

Franchise Sales

€4,832m 13.8% of company sales

at CER



General Medicines & Emerging Markets: Leading Position in Emerging Countries Due to an Adapted Product Portfolio



2017 GEM Business Unit Sales €14,048m

40% of company sales

Established Products





 Account for nearly one third of company sales



Emerging Markets⁽¹⁾

- Sanofi ranks #1⁽²⁾
- 2017 sales: €10,258m, +6.0%⁽³⁾
- Supported by strong sales growth in China:
 €2.2bn, +15.1%⁽³⁾



- (1) World excluding U.S., Canada, Europe, Japan, South Korea, Australia, New Zealand and Puerto Rico
- (2) Market share (without Vaccines), IMS MIDAS | FY 2017
- (3) At Constant Exchange Rates and Constant Structure

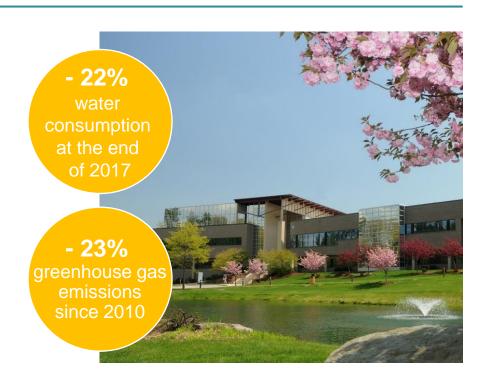
Progress in Protecting the Environment

Water saving

 Objective: reduce water consumption between 2010 and 2020 by 25 %

Carbon footprint

 Objective: reduce greenhouse gas emissions by 50 % in 2025





Access to Healthcare for Those Most in Need in Emerging Countries



Commitment against infectious diseases

- Malaria: >450 million treatments provided in 10 years
- Tuberculosis
- Poliomyelitis: world leader in polio vaccine

Fight against tropical neglected diseases

- Sleeping sickness
 - In 15 years: reduction from 30,000 to 2,000 cases
 - Fexinidazole, developed with DNDi, would be the first all-oral treatment



Take Action for Disadvantaged Children and Educate them About Health



Sanofi Espoir Foundation – My Child Matters

 For 12 years, enabling children with cancer in low-income countries to benefit from better care

Kids

- Preventing diabetes in children at school and encourage healthy lifestyles
- Program launched in Brazil, the United Arab Emirates, Egypt, Japan, India, Pakistan and Poland



In France: Promoting Access to Healthcare for Those in Need



Women in vulnerable situations

 Pregnancy monitoring at the Maison des femmes de St Denis

Those who live on the street

• Partnerships with: Samu social, Croix Rouge Française, Emmaüs, Médecins du Monde

Migrants

 A new program prepared by Sanofi Espoir Foundation







Progress in R&D Elias Zerhouni

Président, Global R&D

Key Achievements in R&D in 2017



Product approvals

- Dupixent^{®(1)} in Atopic Dermatitis in adult patients
- **Kevzara**®(1) in Rheumatoid Arthritis
- Suliqua® in Diabetes (Europe)



Regulatory submissions

- **Dupixent**®(1) in Asthma in adult patients (U.S.)
- VaxiGrip® QIV IM in 6-35 month-olds (EU)



Pivotal study starts

- Dupixent^{®(1)} in Asthma in patients aged 6-11 year-old
- Dupixent^{®(1)} in Atopic Dermatitis in patients 6-11 years-old, 12-17 years-old, 6 month-5 years-old
- **Dupilumab**⁽¹⁾ in Nasal Polyposis
- **Efpeglenatide**⁽²⁾ in type 2 Diabetes
- Sotagliflozin⁽³⁾ in combination therapies in type 2 Diabetes
- Isatuximab in Multiple Myeloma and other cancers
- Cemiplimab⁽¹⁾ in Non-small Cell Lung Cancer, Basal Cell Carcinoma and Cervical Cancer
- Fluzone ® QIV HD



⁽²⁾ In collaboration with Hanmi

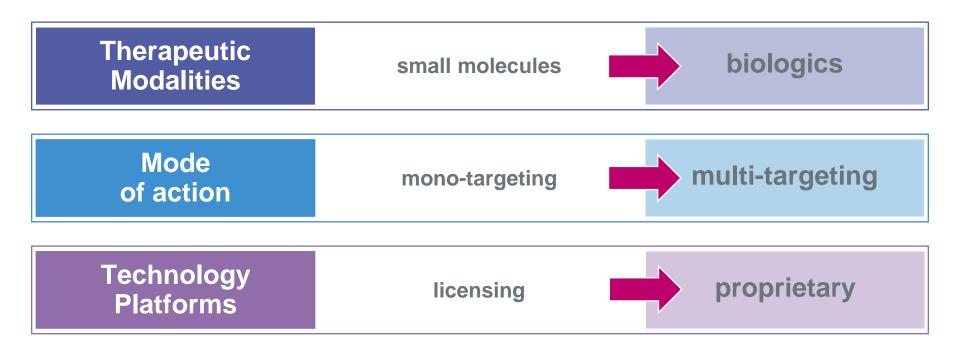
Key R&D Milestones in 2018

Potential submissions	Cemiplimab ⁽¹⁾ in Cutaneous Squamous Cell Carcinoma Sotagliflozin ⁽²⁾ in Type 1 Diabetes Dupilumab ⁽¹⁾ in Asthma in adolescents/adults (EU) Praluent ^{®(1)} ODYSSEY OUTCOMES label update Dupilumab ⁽¹⁾ in Atopic Dermatitis in adolescents Isatuximab in Multiple Myeloma in combination with PomDex (U.S.)	Q1	Q2	Q3	Q4
Expected pivotal trial read-outs	Praluent®(1) ODYSSEY OUTCOMES Cemiplimab(1) in Basal Cell Carcinoma Dupilumab(1) in Nasal Polyps	✓			
Expected start of pivotal studies	Mavacamten ⁽³⁾ in Obstructive Hypertrophic Cardiomyopathy Fitusiran in Hemophilia A & B as prophylactic treatment Isatuximab in 1 st line Multiple Myeloma in Stem Cell Transplant eligible patients Venglustat in Autosomal Dominant Polycystic Kidney Disease Sotagliflozin ⁽²⁾ in Worsening Heart Failure in Diabetes patients Dupilumab ⁽¹⁾ in Eosinophilic Esophagitis Dupilumab ⁽¹⁾ in Chronic Obstructive Pulmonary Disease Alemtuzumab in Primary Progressive Multiple Sclerosis	✓			



⁽¹⁾ In collaboration with Regeneron(2) In collaboration with Lexicon

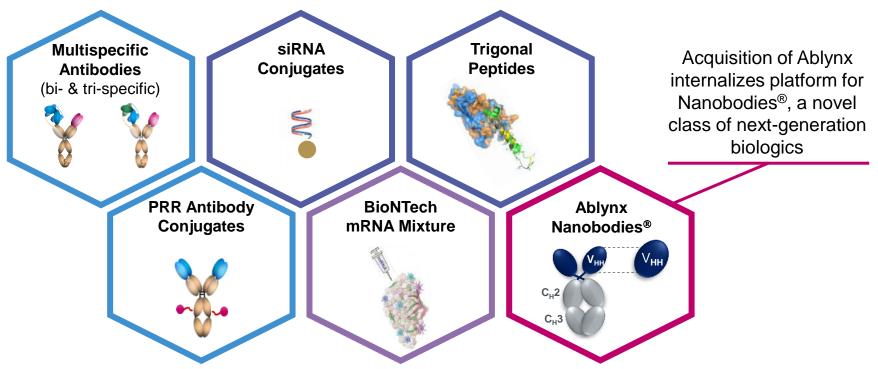
New R&D Model Based on Three Strategical Transformations



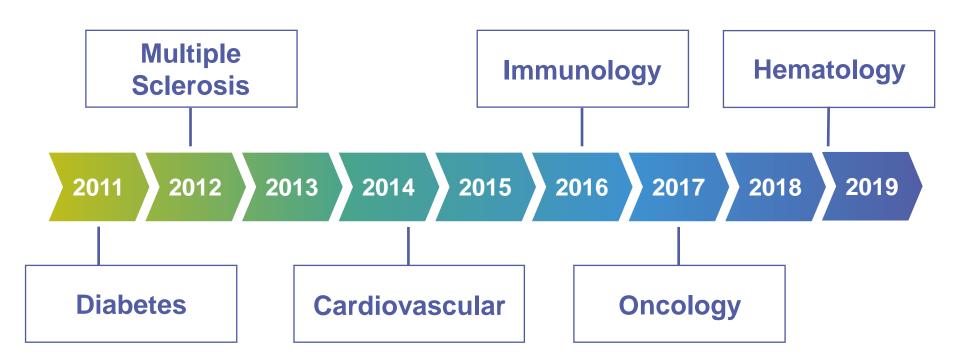


Sanofi Focuses on Leading Technology Platforms

Addressing Multiple Disease Targets with Single Complex Molecules



Build New Foundations for Growth





Dupilumab – A Pipeline in a Product Being Developed in Multiple Indications

Atopic Dermatitis



- Breakthrough therapy in moderate-to-severe AD
- First-in-class biologic treatment

Asthma



- Efficacy in 3 pivotal trials
- Largest Phase 3
 program of a biologic therapy in asthma

Nasal Polyposis



- Positive Proof of Concept data
- No currently approved biologic
- Phase 3 fully enrolled

Eosinophilic Esophagitis



- Positive Proof of Concept data
- No currently approved biologic
- Phase 3 expected to start in Q4 2018

dupilumab



ODYSSEY OUTCOMES Provides Strong Clinical Evidence of Patient Benefit from Long-Term Therapy with Praluent®(1)





Long-term trial ~19,000 patients



Positive Results

- Significant reduction in major adverse cardiac events of 15%⁽²⁾
- Associated with 15% reduction in all-cause mortality⁽³⁾
- Safety profile consistent with previous findings

Praluent® is developed in collaboration with Regeneron

ACS = acute coronary syndrome; LDL-C = low-density lipoprotein cholesterol



Patient Impact

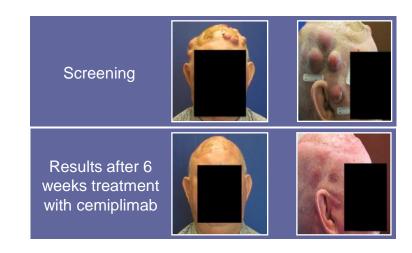
- Greatest clinical benefit shown in high risk post-ACS patients
- In highest risk patients with LDL-C ≥100mg/dL, Praluent[®] was associated with a 24% reduction of major adverse cardiovascular events and 29% reduction in all-cause death⁽⁴⁾



Cemiplimab: Promising Results in Treatment of Cutaneous Squamous Cell Carcinoma

Cutaneous Squamous Cell Carcinoma - Results from Phase 2 Pivotal Study

- One of the most common cancers globally with significant unmet needs
 - 200K to 400K new cases/year in the U.S.⁽¹⁾
 - Severe morbidity and mortality with recurrence
 - \sim 3,900 8,800 deaths/year in the U.S.⁽¹⁾
- Pivotal studies in other indications
 - Non-Small Cell Lung Cancer
 - Basal Cell Carcinoma
 - Cervical Cancer

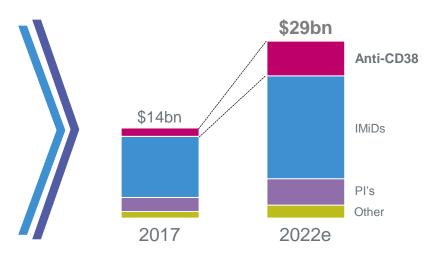




Isatuximab: a Significant Opportunity in Large and Growing Multiple Myeloma Market

Worldwide Multiple Myeloma market expected to reach \$29bn in 2022⁽¹⁾

- Double/triple branded combination use
- New options with prolonged progression free survival benefit
- Globally ~114k new cases diagnosed annually
- Anti-CD38 class rapidly becoming standard of care
 - Combinability without increased toxicity
 - Unprecedented progression free survival prolongation



 Proof of concept studies to evaluate combination use of isatuximab in 9 other cancers



			avalglucosidase alfa Neo GAA Pompe Disease - U.S./EU			GZ389988 TRKA antagonist Osteoarthritis - U.S./EU	SAR422459 ABCA4 gene therapy Stargardt Disease U.S./EU	Combination ferroquine / OZ439(**) Antimalarial - U.S./EU
isatuximab anti-CD38 mAb 3L RRMM (ICARIA) - U.S.			olipudase alfa rhASM ASD ⁽⁶⁾ - U.S./EU			SAR156597 IL4/IL13 bi-specific mAb Systemic Scleroderma - u.s./EU	venglustat Oral GCS inhibitor GrPD ⁽⁹⁾ - U.S./EU	SP0232 mAbs(10)(**) Respiratory syncytial virus U.S.
cemiplimab ^{(2)(**)} PD-1 inhibitor mAb Advanced CSCC - u.s./Eu			fitusiran (7) siRNA inhibitor Hemophilia A/B - U.S./EU			SAR440340(**) Anti-IL33 mAb Asthma - U.S./EU	SAR425899 GLP-1/GCG dual agonist Obesity/Overweight in T2D U.S./EU	Tuberculosis Recombinant subunit vaccine
sotagliflozin(**) Oral SGLT-1&2 inhibitor Type 1 Diabetes - U.S./EU	SAR341402 Rapid acting insulin Type 1/2 Diabetes - EU ⁽⁴⁾		BIVV009 ⁽⁸⁾ Anti Complement C1s mAb Cold Agglutinin Disease		efpeglenatide(**) Long acting GLP1-R agonist Type 2 Diabetes - U.S./EU	SAR566658 Maytansin-loaded anti-CA6 mAb Triple Negative Breast Cancer	SAR407899 rho kinase Microvascular Angina - U.S./EU	HIV Viral vector prime & rgp120 boost vaccine
2018	20	19	20	20	2021	202	2 and beyo	nd
dupilumab ^{(3)(**)}			sarilumab(**)		(2)(**)	044		
Anti-IL4Rα mAb Asthma adults & adolesc EU	Dupixent®(3)(**) Anti-IL4Rα mAb AD 6 - 11 years old - U.S./EU	sotagliflozin(**) Oral SGLT-1&2 inhibitor Type 2 Diabetes – EU ⁽⁵⁾	Anti-IL6R mAb Polyarticular Juvenile Idiopathic Arthritis - U.S./EU	Shan 6 DTP-HepB-Polio-Hib Pediatric hexavalent vaccine	dupilumab^{(3)(**)} Anti-IL4Rα mAb Asthma 6 - 11 years old _{U.S./EU}	Dupixent [®] (**) Anti-IL4Rα mAb AD 6 months - 5 years old U.S/EU	venglustat Oral GCS inhibitor Fabry Disease - U.S./EU	Adacel+ Tdap booster
Anti-IL4Rα mAb	Anti-IL4Rα mAb	Oral SGLT-1&2 inhibitor	Anti-IL6R mAb Polyarticular Juvenile	DTP-HepB-Polio-Hib	Anti-IL4Rα mAb Asthma 6 - 11 years old	Anti-IL4Rα mAb AD 6 months - 5 years old	Oral GCS inhibitor	
Anti-IL4Rα mAb Asthma adults & adolesc EU Dupixent ^{®(3)(**)} Anti-IL4Rα mAb	Anti-IL4Ra mAb AD 6 - 11 years old - u.s./EU dupilumab(*)(**) Anti-IL4Ra mAb Nasal Polyposis Adult - u.s./EU cemiplimab(*2)(**)	Oral SGLT-1&2 inhibitor Type 2 Diabetes – EU ⁽⁵⁾ Fluzone [®] QIV HD Quadrivalent inactivated	Anti-IL6R mAb Polyarticular Juvenile Idiopathic Arthritis - U.S./EU cemiplimab ^{(2)(**)} PD-1 inhibitor mAb	DTP-HepB-Polio-Hib Pediatric hexavalent vaccine Pediatric pentavalent vaccine	Anti-IL4Ra mAb Asthma 6 - 11 years old U.S./EU isatuximab Anti-CD38 mAb (IMROZ) 1L Newly Diagnosed MM	Anti-IL4Ra mAb AD 6 months - 5 years old U.S/EU sarilumab(") Anti-IL6R mAb Systemic Juvenile Arthritis	Oral GČS inhibitor Fabry Disease - u.s./EU venglustat Oral GCS inhibitor Gaucher Disease Type 3	Tdap booster Rabies VRVg

- (1) Excluding Phase 1 Data related to all studies published on clinicaltrials.gov
- (2) Also known as SAR439684 and REGN2810
- (3) Also known as SAR231893
- (4) Submission strategy for the U.S. under evaluation
- (5) Submission for the U.S. expected in 2020
- (6) Acid Sphingomyelinase Deficiency
 (7) Following the Alnylam/Sanofi strategic restructuring of the RNAi therapeutics rare disease alliance announced in January 2018, Sanofi now has global rights on fitusiran
- (8) Currently operating as separate entities. Reported dates are based on prior Bioverativ disclosure of study completion date
- (9) Gaucher Related Parkinson's Disease
- (10)Also known as MEDI8897
- (**) Partnered and/or in collaboration Sanofi may have limited or shared rights on some of these products
- Immuno-inflammation Oncology Cardiovascular & metabolism Infectious Diseases Rare Diseases Vaccines Rare Blood Disorders MS, Neuro, Gene therapy



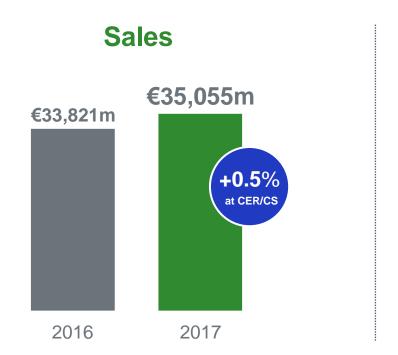




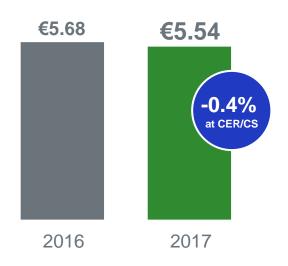
2017 Financial Performance

Jérôme Contamine Executive Vice-President, Chief Financial Officer

2017 Business EPS in-Line with Expectations



Business EPS

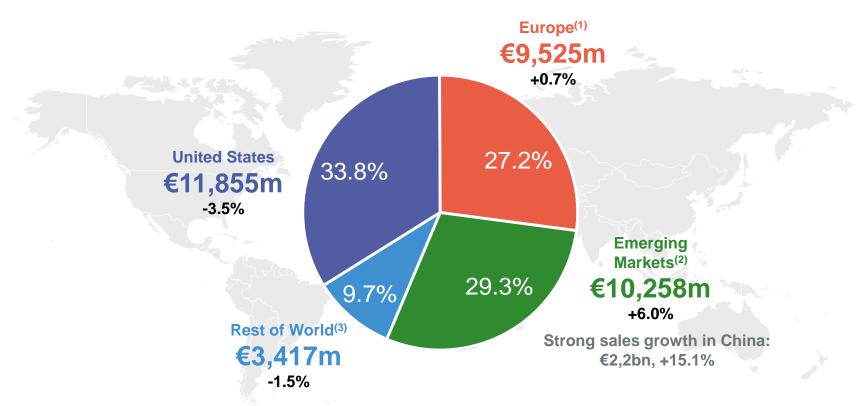


2017: A Year of Transition Supported by Solid Growth Drivers

	of Sales				Growth at CER	Growth at CER/CS
7	1.6%	Pharmaceutica	ls	€25,122M	-1.2%	-1.3%
	19.7%	Diabetes &	Cardiovascular	€6,905M	-9.6%	-9.6%
	19.1%	Specialty N	ledicines	€6,678M	+14.5%	+14.6%
	27.7%	Established	d Rx Products	€9,761M	-3.4%	-3.8%
	5.1%	Others		€1,778M	-3.3%	-3.1%
9 000	3.8%	Consumer Heal	thcare	€4,832M	+46.3%	+2.1%
1	4.6%	Vaccines		€5,101M	+14.5%	+8.3%



A Balanced Geographic Breakdown of Sales





All growth at CER (constant exchange rates) / CS (constant structure), adjusted for BI CHC business, termination of SPMSD and others

(1) Western Europe & Eastern Europe (excluding Eurasia)

- (2) World excluding U.S., Canada, Europe, Japan, South Korea, Australia, New Zealand and Puerto Rico
- (3) Japan, South Korea, Canada, Australia, New Zealand and Puerto Rico

Sanofi Met 2017 Financial Performance Objectives

	Objectives	FY 2017 Results
✓ Gross margin	70-71% at CER	70.6%
✓ OpEx growth rate at CER	At similar rate as in 2016 ⁽¹⁾	+2.0%
✓ Tax rate	24-25%	23.5%
Business EPS guidance at CER	Broadly stable	-0.4%
✓ Dividend growth	Progressive	+2.4%

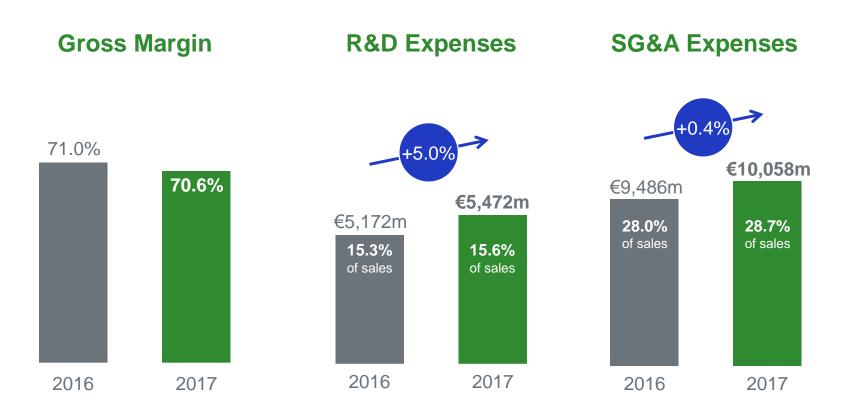


Business Operating Income Up 3% at CER in 2017

€m	2017	2016	%Change (reported €)	% Change (CER)	% Change (CER/CS)
Sales	35,055	33,821	+3.6%	+5.6%	+0.5%
Other revenues	1,149	887	+29.5%	+32.9%	+36.6%
Gross profit	24,759	24,006	+3.1%	+5.3%	+0.4%
R&D expenses	(5,472)	(5,172)	+5.8%	+7.0%	+5.0%
SG&A expenses	(10,058)	(9,486)	+6.0%	+7.8%	+0.4%
Business operating income	9,343	9,285	+0.6%	+3.0%	-1.1%
Effective tax rate	23.5%	23.3%	-	-	-
Business net income excluding Animal Health	6,964	6,832	+1.9%	+4.2%	-
Business net income of Animal Health	-	476	-100.0%	-100.0%	-
Business net income	6,964	7,308	-4.7%	-2.6%	



Continued Investment in R&D and Strict Cost Management





Net Income Benefited From Gain on the Disposal of the Animal Health Business

€m	2017	2016	% Change (reported €)
Business net income	6,964	7,308	(4.7%)
Amortization of intangible assets	(1,866)	(1,692)	
Impairment of intangible assets	(293)	(192)	
Fair value remeasurement of contingent consideration liabilities	(159)	(135)	
Expenses arising from the impact of acquisitions on inventories	(166)	-	
Restructuring costs and similar items	(731)	(879)	
Other gains and losses, and litigation	(215)	211	
Tax effect of items listed above	1,126	841	
Other tax items	(742)	(113)	
Associates and non-controlling interests	(127)	31	
Animal Health items	4,643	(162)	
Others ⁽¹⁾	-	(509)	
Net income attributable to equity holders of Sanofi	8,434	4,709	79.1%



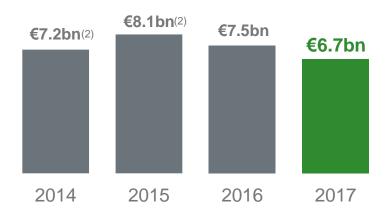
Strong Balance Sheet on December 31, 2017

		December 31, 201 (€bn)	7 Change vs. December 31, 2016
	Intangible assets	53.4	+2.2
ASSETS	Other non-current assets	20.1	-0.3
ASSETS	WCR ⁽¹⁾	1.8	+0.2
	Net cash (B)	10.3	+0.0
_	Equity attributable	58.3	+0.6
& EQUITY	Provisions and other non-current liabilities	11.8	-0.7
	Financial debt (A) ⁽²⁾	15.5	-3.0
	Net Debt (A-B)	5.2	-3.0



High and Strong Cash-Flow Allows to Finance our Acquisitions at Excellent Conditions

Free Cash Flow⁽¹⁾

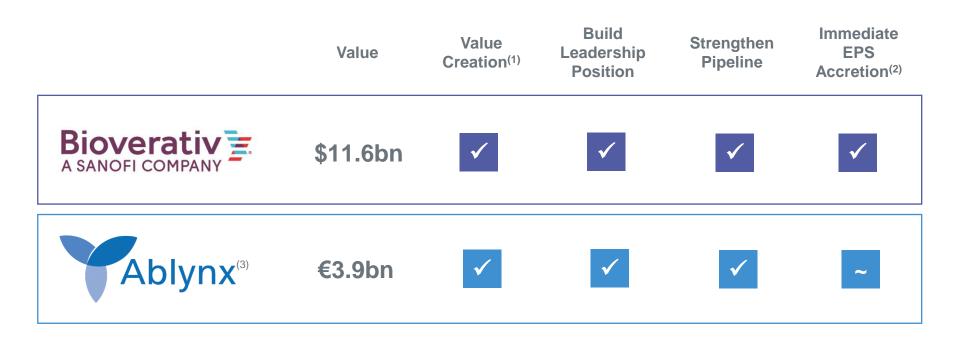


Net Debt €7.1bn €7.3bn €5.2bn March 2014 2015 2016 2017 March 2018

- Strong long-term credit ratings
 - Moody's A1; S&P AA

- Successful bond issues in March 2018
 - €8bn; average cost of 0.96%

Strategically and Financially Compelling Acquisitions to Enhance Sanofi's Growth Profile and Create Value





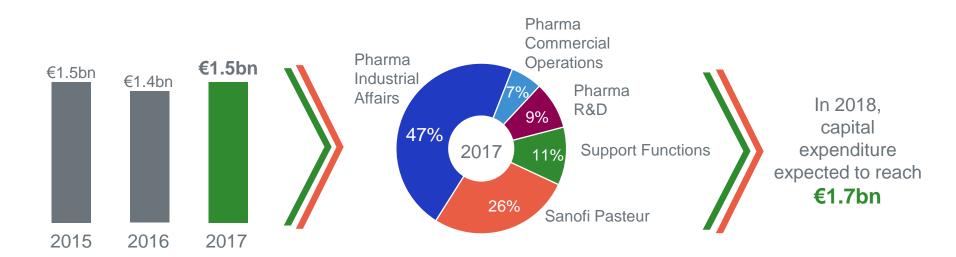
⁽¹⁾ Bioverativ: projected to achieve ROIC in excess of cost of capital within three years

(3) Subject to completion of the acquisition

⁽²⁾ Business EPS is a non-GAAP financial measure (see appendix to Sanofi quarterly financial release definitions) Ablynx: including R&D expenses, the acquisition is expected to be neutral to 2018 and 2019 Business EPS

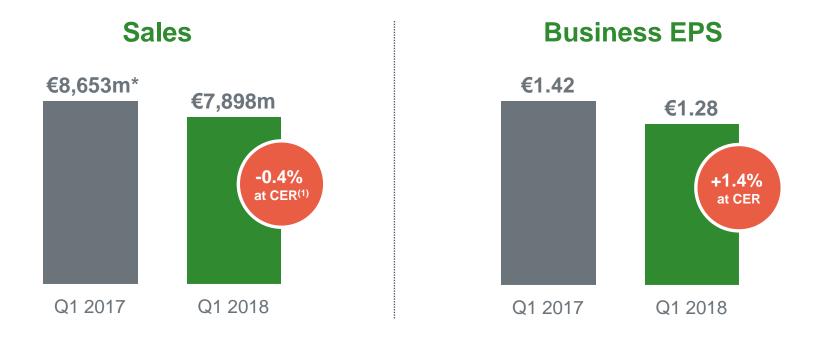
€1.5bn Capital Investments in 2017: Further Expansion in Biologics Production Capacity

Capital Expenditure Evolution and Breakdown in 2017





Q1 2018: Performance of Specialty Care and Emerging Markets Offsets U.S. Lantus[®] and Sevelamer Loss of Exclusivity





Expected Return to Growth in H2 2018

- 1 First quarter in line with 2018 guidance
- 2 Impact of exchange rates movements
- 3 Progress of new products
- 4 Establishment of a leadership in rare blood disorders
- 5 Strong long-term credit ratings confirmed

Expected evolution of Business EPS in 2018: +2% to +5% at CER^(1,2)

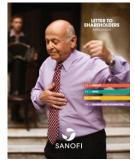


An Active Communication with our Individual Shareholders

Your publications

- Shareholder Handbook
- Letter to Shareholders
- Fact Sheet













Your online information

- www.sanofi.com/shareholders
- Sanofi IR mobile app
- Social Media













Meetings with Shareholders

Meetings planned in 2018



- Salon Actionaria 2018
 - November 22-23: Paris



Shareholders Committee



- 4 events planned in 2018
- Visit of Marcy-l'Etoile Sanofi Pasteur industrial site
- Meetings with management and the Chairman







Reports by Statutory Auditors at the Combined General Meeting

PricewaterhouseCoopers Audit, ERNST & YOUNG et Autres

Reports and Statements Prepared by the Statutory Auditors and Made Available to Shareholders

Financial statements

- Statutory Auditors' report on the statutory financial statements (1st resolution)
- Statutory Auditors' report on the consolidated financial statements (2nd resolution)

Related-party agreements and commitments

 Statutory Auditors' special report on related-party agreements and commitments (4th resolution)

Social, environmental and societal information

 Report by one of the Statutory Auditors, appointed as an independent third party, on the consolidated human resources, environmental and social information included in the management report

Profit forecasts

 Statutory Auditors's report on forecasted business net income per share

Other statements

- Statutory Auditors' statement on compensation
- Statutory Auditors' statement on the information provided in accordance with Article L.225-115-5° of the French Commercial Code with respect to the total amount of payments made pursuant to paragraphs 1 and 4 of Article 238 bis of the French Tax Code







Questions & Answers





Vote on the Resolutions