# A LEADER IN LIFE SCIENCE

Merck KGaA, Darmstadt, Germany

Morgan Stanley Global Healthcare Conference, New York

Udit Batra, CEO Life Science

12 September 2016



# Disclaimer

Publication of Merck KGaA, Darmstadt, Germany. In the United States and Canada the group of companies affiliated with Merck KGaA, Darmstadt, Germany operates under individual business names (EMD Serono, Millipore Sigma, EMD Performance Materials). To reflect such fact and to avoid any misconceptions of the reader of the publication certain logos, terms and business descriptions of the publication have been substituted or additional descriptions have been added. This version of the publication, therefore, slightly deviates from the otherwise identical version of the publication provided outside the United States and Canada.



# **Disclaimer**

#### Cautionary Note Regarding Forward-Looking Statements and financial indicators

This communication may include "forward-looking statements." Statements that include words such as "anticipate," "expect," "should," "intend," "plan," "project," "seek," "believe," "will," and other words of similar meaning in connection with future events or future operating or financial performance are often used to identify forward-looking statements. All statements in this communication, other than those relating to historical information or current conditions, are forward-looking statements. We intend these forward-looking statements to be covered by the safe harbor provisions for forward-looking statements in the Private Securities Litigation Reform Act of 1995. These forward-looking statements are subject to a number of risks and uncertainties, many of which are beyond control of Merck KGaA, Darmstadt, Germany, which could cause actual results to differ materially from such statements.

Risks and uncertainties include, but are not limited to: the risks of more restrictive regulatory requirements regarding drug pricing, reimbursement and approval; the risk of stricter regulations for the manufacture, testing and marketing of products; the risk of destabilization of political systems and the establishment of trade barriers; the risk of a changing marketing environment for multiple sclerosis products in the European Union; the risk of greater competitive pressure due to biosimilars; the risks of research and development; the risks of discontinuing development projects and regulatory approval of developed medicines; the risk of a temporary ban on products/production facilities or of non-registration of products due to non-compliance with quality standards; the risk of an import ban on products to the United States due to an FDA warning letter; the risks of dependency on suppliers; risks due to product-related crime and espionage; risks in relation to the use of financial instruments; liquidity risks; counterparty risks; market risks; risks of impairment on balance sheet items; risks from pension obligations; risks from e-crime and cyber attacks; risks due to failure of business-critical information technology applications or to failure of data center capacity; environmental and safety risks; unanticipated contract or regulatory issues; a potential downgrade in the rating of the indebtedness of Merck KGaA, Darmstadt, Germany; downward pressure on the common stock price of Merck KGaA, Darmstadt, Germany and its impact on goodwill impairment evaluations; the impact of future regulatory or legislative actions; and the risks and uncertainties detailed by Sigma-Aldrich Corporation ("Sigma-Aldrich") with respect to its business as described in its reports and documents filed with the U.S. Securities and Exchange Commission (the "SEC").

The foregoing review of important factors should not be construed as exhaustive and should be read in conjunction with the other cautionary statements that are included elsewhere, including the Report on Risks and Opportunities Section of the most recent annual report and quarterly report of Merck KGaA, Darmstadt, Germany, and the Risk Factors section of Sigma-Aldrich's most recent reports on Form 10-K and Form 10-Q. Any forward-looking statements made in this communication are qualified in their entirety by these cautionary statements, and there can be no assurance that the actual results or developments anticipated by us will be realized or, even if substantially realized, that they will have the expected consequences to, or effects on, us or our business or operations. Except to the extent required by applicable law, we undertake no obligation to update publicly or revise any forward-looking statement, whether as a result of new information, future developments or otherwise.

This quarterly presentation contains certain financial indicators such as EBITDA pre exceptionals, net financial debt and earnings per share pre exceptionals, which are not defined by International Financial Reporting Standards (IFRS). These financial indicators should not be taken into account in order to assess the performance of Merck KGaA, Darmstadt, Germany in isolation or used as an alternative to the financial indicators presented in the consolidated financial statements and determined in accordance with IFRS. The figures presented in this quarterly statement have been rounded. This may lead to individual values not adding up to the totals presented.



# Agenda

# **D** Business introduction

**02** Strategic review Life Science









# Group Portfolio of three high-tech businesses



# Leading in specialty pharma markets

- Biologics and small-molecules
- Research focus: Oncology, Immunology & Immuno-Oncology
- Over-the-counter medicine



### Leading life science company

- Tools and services for biotech
   research & production
- Tools and laboratory supply for the academic research and industrial testing



# Market leader in display materials

- Innovative display materials
- Effect pigments and functional materials
- High-tech materials for electronics



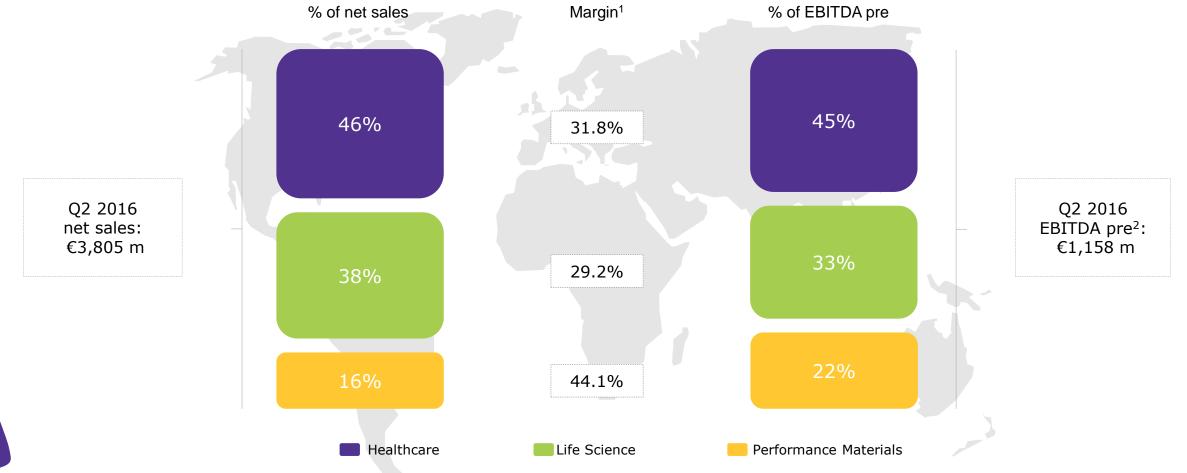
# Group Profitability improved fundamentally





<sup>1</sup>Included since 2 May 2014; <sup>2</sup>2007 and 2014 EBITDA pre margin adjusted for comparability

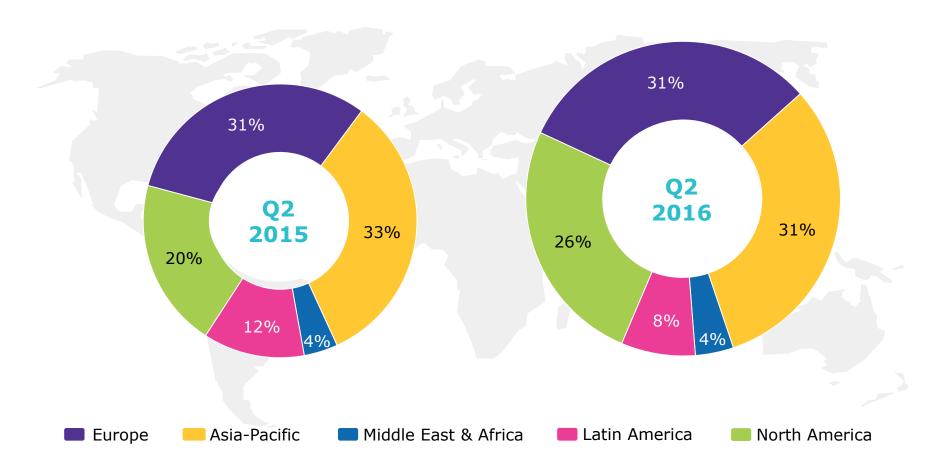
# Group Strong businesses with attractive margins





# Group Balanced geographic footprint

Group Q2 2015 and Q2 2016 net sales by region [in %]





# Group We are well set for profitable and sustainable growth



Maximize growth of existing franchises Deliver on pipeline

Focus on seamless integration and deliver cost synergies Leverage strategic capabilities for value creation

Drive innovation and technology leadership across all businesses Innovate in applications beyond displays



# Group Upgrade of full-year 2016 guidance

Group guidance for 2016













# MilliporeSigma: Serving customers across Life Science



- Academic and government institutions
- Biopharma R&D
- Industry R&D



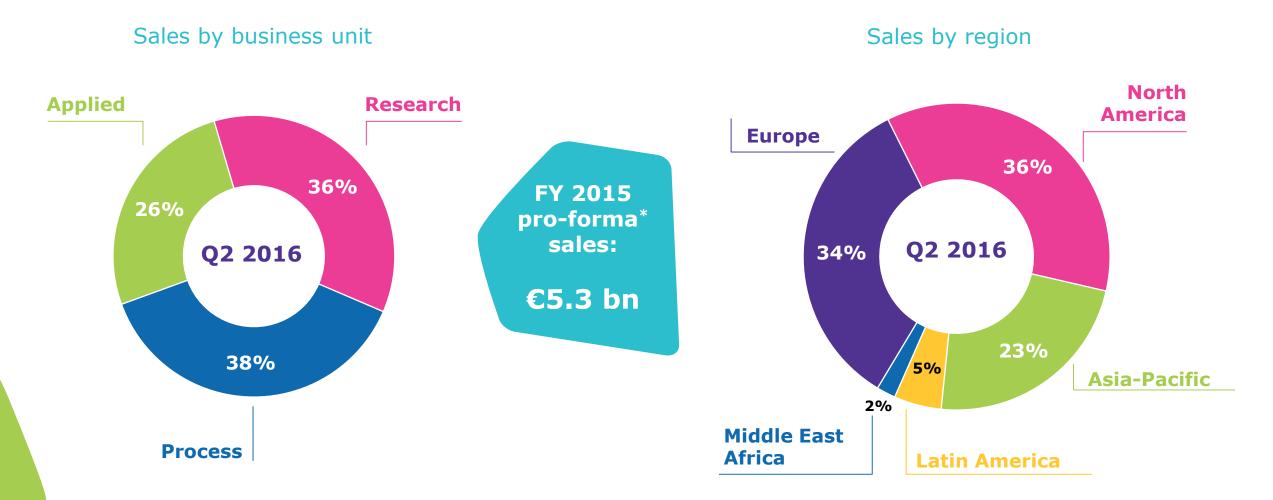
- Pharmaceutical companies
- Small biotech
- Contract manufacturing organizations



- Diagnostic manufacturers
- Clinical testing labs
- Food & Beverage manufacturers



# A balanced portfolio and geographic presence



14

# Life Science is an attractive market



- Growth in volume of experiments
- Mild growth in academic funding
- Investment in industry R&D



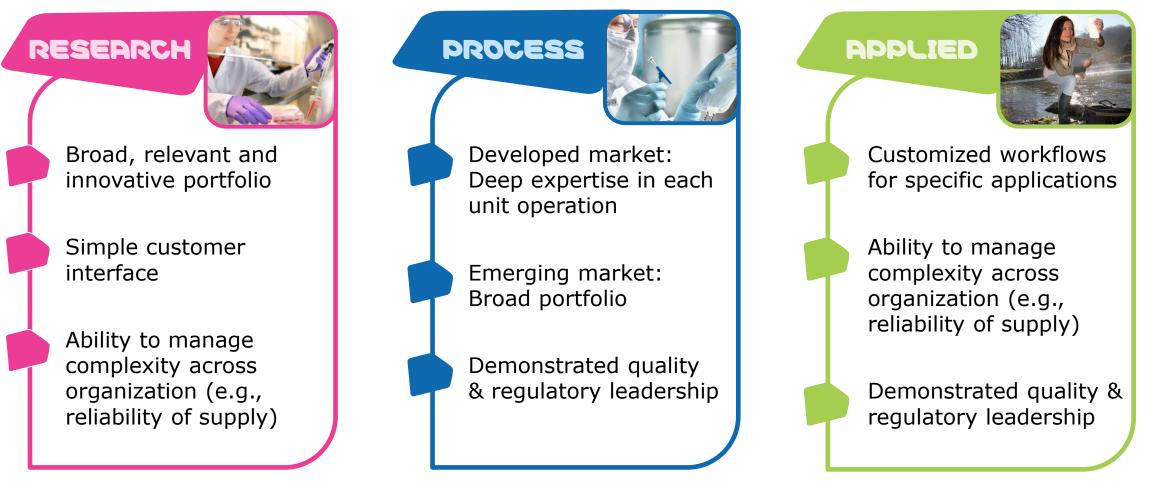
- Drug volume growth
  - from biologics
  - from emerging modalities
- Continued shift to single-use



- Volume growth from
  - Population growth
  - Increased testing needs

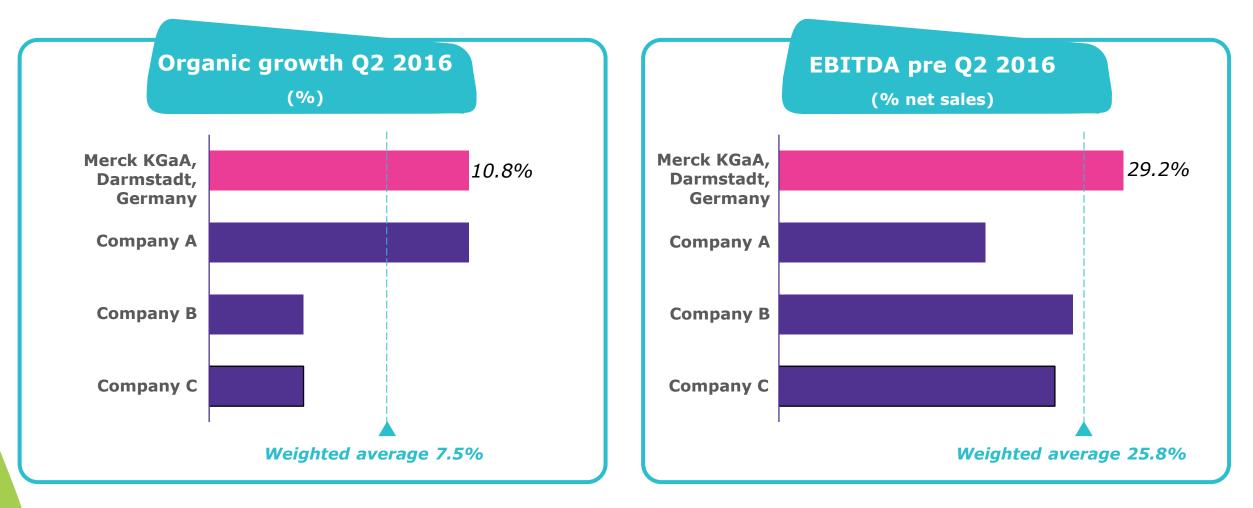


# Success driven by portfolio breadth and differentiation, a customer-centric approach and world-class capabilities





# **Continued momentum and outperformance of peers in Q2 2016**



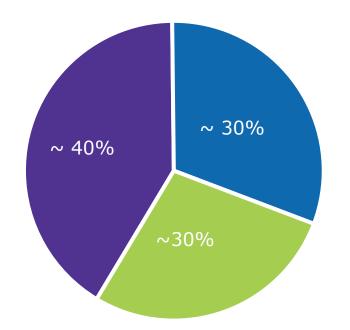


# **Integration on track**





# **Expected synergies identified and fully confirmed**

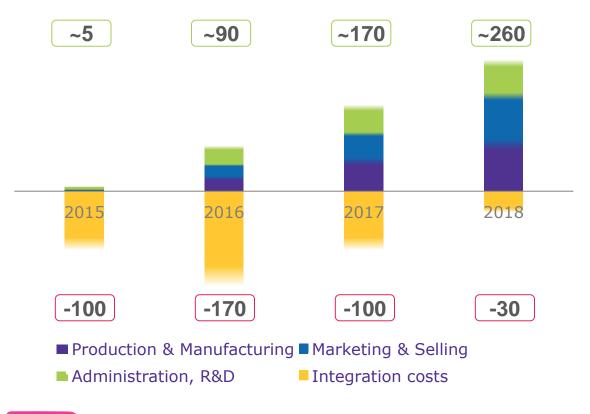


Sources of synergies (3<sup>rd</sup> full year 2018)

Production & Manufacturing Marketing & Selling
 Administration, R&D

Three major areas for delivering the synergies

### Timing of expected synergies and related costs [€m]







# **Three levers for value creation**



- Over 300,000 products
- End-to-end in biologics manufacturing
- Short- and long-term innovation, e.g. antibodies and bioreactors



- Over 60 countries
- Leader in North America, and critical mass in emerging markets, e.g. Latin America, APAC
- Global manufacturing and distribution footprint



- World-class ecommerce platform
- Deep technical expertise
- Focus on quality and service



# Process Solutions **Our end-to-end portfolio for manufacturing mAbs**



MAKE Produce antibodies



to enhance cell

EX-CELL® Advanced™ CHO Fed-batch Medium **Cell culture media** 200



2000L CellReady bioreactor **Tank for** 

Clarisolve <sup>®</sup> clarification filters **Removing ce** debris **PURIFY** Remove cell debris, virus, etc.



FlexReady <sup>®</sup> chromatography **Purifying mAbs** 



Viresolve<sup>®</sup> Pro solution *Removing viruses from protein solutions*  Pellicon<sup>®</sup> cassette filters **Washing and** removing cells lipids, particle



**FORMULATE** Final drug product

Opticap<sup>®</sup> capsules Sterile filtration

2000L CellReady bioreactor Tank for cultivating cells Provantage ® BioReliance ® EMPROVE®



growth

# **#1** website in research life science industry

76 m Visits 270 m Page views

Industry leading e-commerce platform and supply chain capability





and peer reviewed

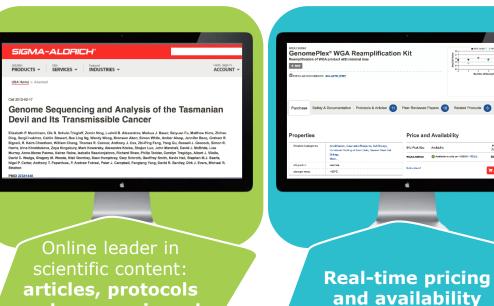
papers



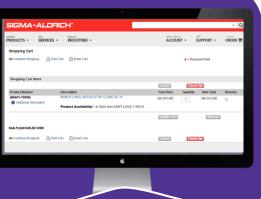
## 2.9 m Orders



Hundreds of thousands of products at your fingertips



Price and Availability Price Ocentity SKU Puck Star 100.50 0 🖸 🕤 🕤



Convenient and simple customer interface: no more than 2 clicks from shopping cart



# **Executive summary: Life Science**

Solid growth drivers for a ~125bn€ market growing at mid-single digit

Strong momentum, performance growing at least with market



# Integration

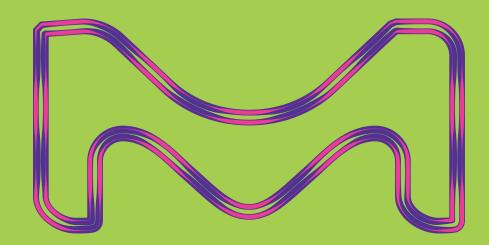
Attractive

Market

strong

**Progress on value delivery, organization** setup, and process integration





# Appendix



# **O2** Sigma acquisition

# **B** Healthcare



Performance Materials











## 2016 business sector guidance





# Additional financial guidance 2016

### Further financial details

Corporate & Other EBITDA pre		~ -€370 – -400 m	
Interest result		~ -€270 – -300 m	
Intangibles amortization from Sigma PPA		~ €250 – 300 m p.a.	
Underlying tax rate		~23% to 25%	
Capex on PPE		~€750 – 800 m	
Hedging/USD assumption	2016 & 20	17 hedge ratio ~40%-45% at EUR/USD ~1.10 to 1.15	
2016 Ø EUR/USD assumption		~1.07 - 1.12	











# Sigma Aldrich acquisition – A compelling transaction rationale



Increasing scale – expanding position in attractive life science industry

Enhancing value for our customers

- Broadens product range and ease of doing business for Laboratories & Academia
- Complements Process Solutions product offering
- Closing the gap in U.S. adequate presence in all geographies
- •Leveraging existing platforms for global innovation rollout

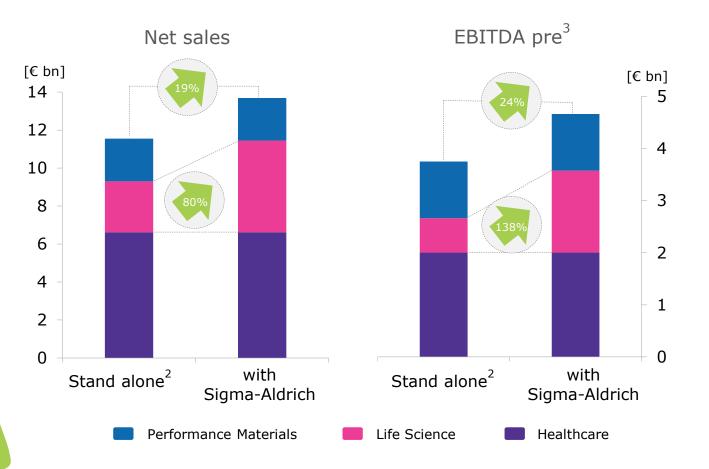
### Financial fit

- Further diversification of revenue stream
- •Substantial synergy potential
- •Immediately accretive to EPS pre<sup>\*</sup> and EBITDA margin
- Solid investment grade rating will be maintained



# Sigma-Aldrich acquisition enhances our financial profile

### Pro-forma 2014<sup>1</sup>



### Pro-forma financial impacts

- •Group sales<sup>1</sup> increase by  $\sim 19\%$
- •Group EBITDA pre<sup>3</sup> rises by ~24% with margin<sup>4</sup> expansion from ~30% to ~33%
- Synergies: ~€260m p.a. fully implemented in 3<sup>rd</sup> full year after closing
- Expected PPA impact: Mid triple-digit €m p.a.
- •Immediately EPS pre accretive

<sup>1</sup>Pro-forma calculation based on published sales for FY 2014 for Merck KGaA, Darmstadt, Germany (including pro-forma AZ Electronic Materials) and Sigma-Aldrich; <sup>2</sup>Pro-forma calculation based on published sales for FY 2014 for Merck KGaA, Darmstadt, Germany (including pro-forma AZ Electronic Materials); <sup>3</sup>Pro-forma calculation based on 100% expected synergies; excluding Corporate & Other; <sup>4</sup>Including Corporate & Other



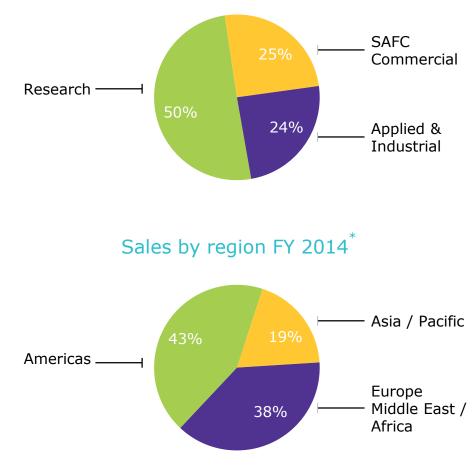
# Sigma-Aldrich – A leading life science consumables supplier

### **Business**

- •Total revenues of \$2.8 billion in 2014
- •~9,000 employees including ~3,000 scientists and engineers
- •Headquartered in St. Louis, MO
- •Chemical and biochemical products, kits and services provider to laboratories and pharma production
- •No. 1 eCommerce platform in the industry; ~1,600 sales people

### Footprint

- •Balanced regional exposure; strength in North America
- •Operations in  $\sim$ 40 countries; products available in  $\sim$ 160 countries

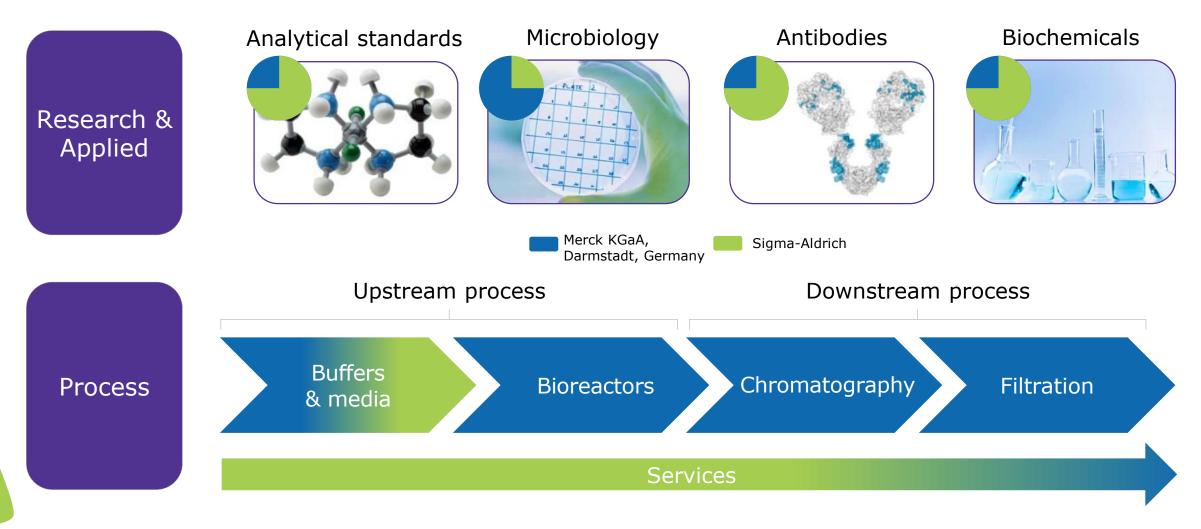


Sales by division FY 2014



32

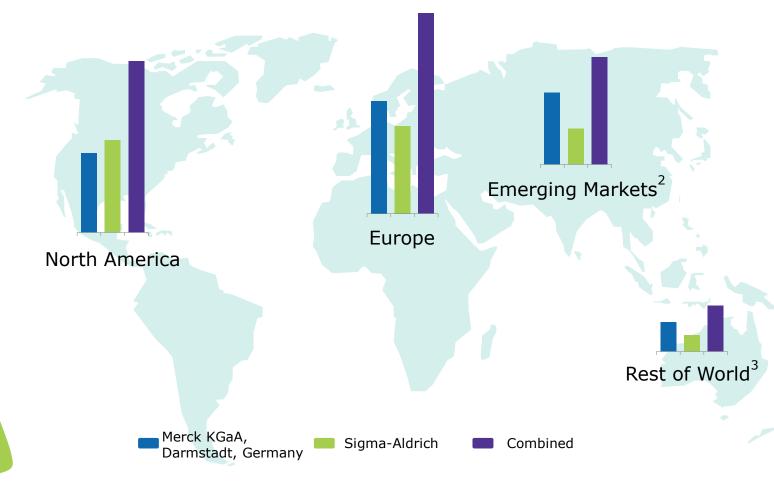
### **Broad and complementary product fit in attractive segments**





# **Expanding global reach and scale**

Global sales<sup>1</sup> footprint of both businesses



### Increased presence in North America

• Benefiting from a leading position in U.S. Laboratory sector

Increased access to U.S. academia

### Exposure to fast-growing Asia

Accelerating growth momentum

•Opportunity to leverage eCommerce platform



## Leveraging operational excellence to deliver superior value to customers

Product innovation

Process innovation

- Delivering innovative workflow solutions to increase customers' efficiency
- Broad technology and platforms
- Recurring winners of renowned innovation awards



Mobius FlexReady





Amnis

- Efficient supply chain for >300,000 products
- •Best in class customer experience; e.g. 24 hour delivery in major markets
- •Top-notch customer interface supported by eCommerce platform



eCommerce platform



Supply chain

Efficient work flow solutions and unique customer experience



# Sigma-Aldrich – Business and transaction financials

### Overview of financial data<sup>1</sup>

US\$ m	2012	2013	2014 <sup>4</sup>
Revenue	2,623	2,704	2,785
% YoY at constant FX	+3%	+3%	+4%
EBITDA (adjusted)	809	821	847
% of sales	31%	30%	30%
D&A	136	138	132
% of sales	5%	5%	5%
Net financial debt (period end)	-41	-357	-513
No. of shares (diluted, m)	122	121	120

#### Proposed transaction details<sup>2</sup>

- •Equity value ~US\$17 bn (€13.1 bn)
- Enterprise value (EV) ~€12.7 bn including net cash ~€360 m<sup>5</sup>
- Financing through cash and debt; no equity
- Assumed synergies: ~€260m
- In line with core acquisition criteria
  - Immediately accretive to EPS pre
  - Solid investment grade rating will be maintained

### Implied forward transaction multiples<sup>3</sup>

	2013	2014
EV/Sales	6.1x	5.9x
EV/EBITDA	20.1x	19.4x
EV/EBITDA pro-forma incl. synergies <sup>3</sup>	14.3x	13.9x



# Support from meaningful synergies

#### Our experience



- Significant restructuring and integration experience
- •Deep knowledge and understanding of the life science industry

#### Source of synergies



- Consolidate manufacturing footprint
- Increase conversion to eCommerce channels
- Optimize sales & marketing
- Streamline admin functions and infrastructure
- •Save U.S. public company costs
- Optimize R&D portfolio

#### Planned delivery

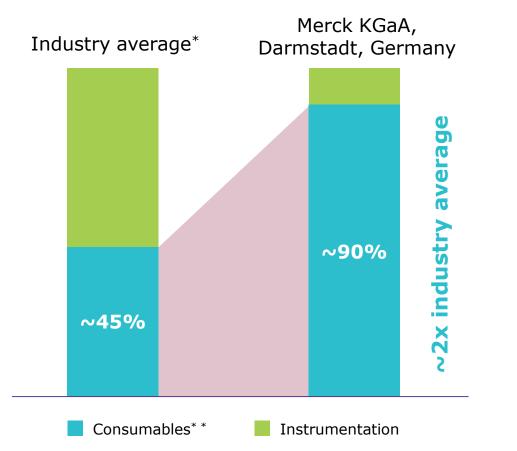


- •Synergies: ~€260 m, i.e. ~12% of Sigma-Aldrich sales
- Fully implemented in third full year after closing
- Expected integration costs:
   ~€400 m; spread over
   2015-2018



# High exposure to consumables makes us unique compared to industry

#### Consumables exposure



#### Pros and cons

+

Consumables paid out of operating versus capex budgets

- Products are often not discretionary and must be used to conduct research or manufacture drugs
- Risk of getting locked out by equipment manufacturer









# Healthcare Healthcare is set to deliver on promising pipeline candidates

**Deliver** on organic growth

Focus on pipeline



Stable existing business to fuel slight organic growth



Solid pipeline of oncology, immuno-oncology and immunology molecules



Transformation of R&D operating model ongoing



Competitive R&D funding in our focus areas



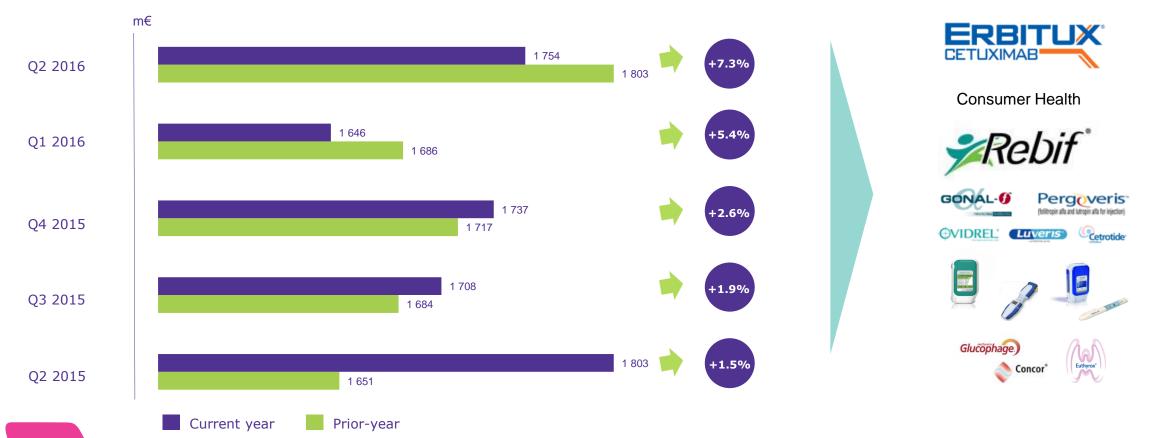
Cost discipline and efficient execution





# Healthcare **Delivered organic sales growth, committed to future performance**

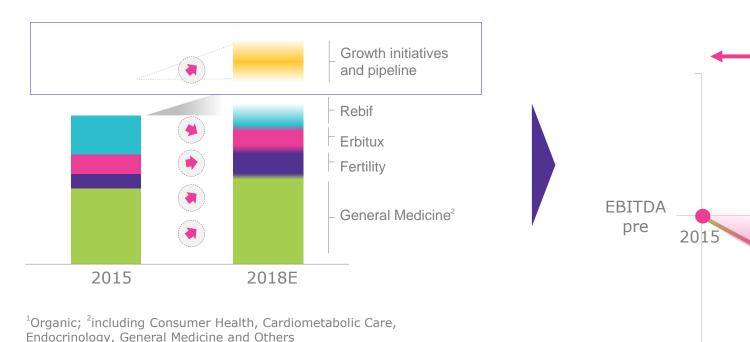
Stable to slight organic growth



Defending the existing product portfolio is a key strength of Healthcare



# Healthcare Pipeline opportunities will lead to rising investments



Stable to slightly growing<sup>1</sup> sales until 2018 confirmed

Should pipeline catalysts materialize, investments will lead to considerable payback as of 2018+

Investment

phase

2017

2018

2016

Long-term

EBITDA pre

upside

2019

2020

Illustration

# Rising investments until 2017 to accelerate sales and earnings growth as of 2018



# Healthcare Investments in future growth

 Immuno-Oncology: avelumab and ramp-up of earlier pipeline projects +€150-200m cost increase in 2016

- Oncology/Immunology, e.g. tepotinib, BTK inhibitor: mid to high double-digit €m cost increase in 2016
- Launch readiness to be ensured for avelumab and cladribine
- Costs for launch preparation in the mid to high double-digit €m range in 2016



- Stringent pipeline assessments continue
- Investments based on sound business cases and robust clinical data

#### Long-term growth investments partly mitigated by strict cost management

\*For scenario that pipeline catalysts materialize



R&D\*

Marketing & Selling<sup>\*</sup>

# Portfolio management: Differentiating across diverse business models

#### General Medicine portfolio



- •Limited risk with high cash generation
- Sustainable steady growth fueled by Emerging Markets

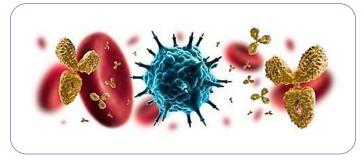
#### Biologicals portfolio



- Moderate risk and reward profile
- Economies of scale due to stateof-the-art production capabilities
- Emerging Markets gain importance



#### Oncology & Immunology innovation portfolio



- High reward at high risk
- Innovation key success factor high R&D spend
- Promising pipeline projects



Mid-term, all parts of the portfolio need to earn their cost of capital



# The road to maximizing existing franchises is clear



Continue to drive front-line mCRC share by increasing patient testing and expanding head and neck coverage



Capitalize on strong efficacy and new smart devices to maximize differentiation and defend franchise



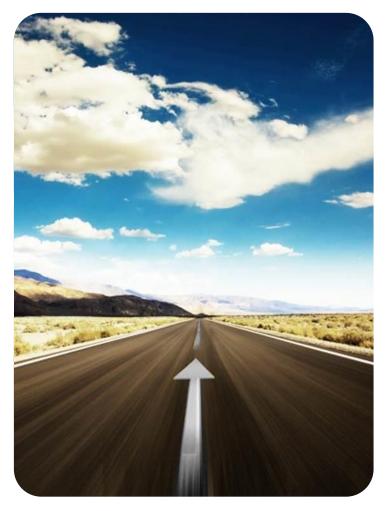
Build on No.1 position and ART<sup>1</sup> channel access with embryo diagnostics and other innovative technologies



Harness strengths of existing business and build a new focus area driven by innovative devices and services for patients



Build on existing track record in Emerging Markets, drive brand and life-cycle management and expand business including asset repatriation





# **Clinical pipeline**

#### Phase I

Tepotinib c-Met kinase inhibitor Solid tumors M2698 p70S6K & Akt inhibitor Solid tumors M3814 DNA-PK inhibitor Solid tumors Beigene-283 BRAF inhibitor Solid tumors

Avelumab Anti-PD-L1 mAb Solid tumors

Avelumab Anti-PD-L1 mAb Hematological malignancies

M9241 (NHS-IL12)<sup>1</sup> Cancer immunotherapy Solid tumors

M7824 Bifunctional immunotherapy Solid tumors

M1095 (ALX-0761) Anti-IL-17 A/F nanobody Psoriasis M2951 BTK inhibitor Systemic lupus erythematosus

#### Phase II

M2736 (ATX-MS-1467) Immune tolerizing agent Multiple sclerosis

#### Tepotinib c-Met kinase inhibitor Non-small cell lung cancer Tepotinib c-Met kinase inhibitor Hepatocellular cancer

Avelumab Anti-PD-L1 mAb Merkel cell carcinoma

Sprifermin Fibroblast growth factor 18 Osteoarthritis Atacicept Anti-Blys/anti-APRIL fusion protein Systemic lupus erythematosus M2951 BTK inhibitor Rheumatoid arthritis

#### Phase III

Avelumab – Anti-PD-L1 mAb Non-small cell lung cancer 1L<sup>2</sup> Avelumab – Anti-PD-L1 mAb Non-small cell lung cancer 2L<sup>3</sup> Avelumab – Anti-PD-L1 mAb Gastric cancer 1L<sup>2</sup> Avelumab – Anti-PD-L1 mAb Gastric cancer 3L<sup>4</sup> Avelumab – Anti-PD-L1 mAb Bladder cancer 1L<sup>2</sup> Avelumab – Anti-PD-L1 mAb Ovarian cancer platinum resistant/refractory Avelumab – Anti-PD-L1 mAb Ovarian cancer 11<sup>2</sup> Avelumab - Anti-PD-L1 mAb Renal cell cancer 1L<sup>2</sup>

MSB11022 Proposed biosimilar of Adalimumab Chronic plaque psoriasis

Pipeline as of July 28th, 2016

Pipeline products are under clinical investigation and have not been proven to be safe and effective. There is no guarantee any product will be approved in the sought-after indication.

#### Registration

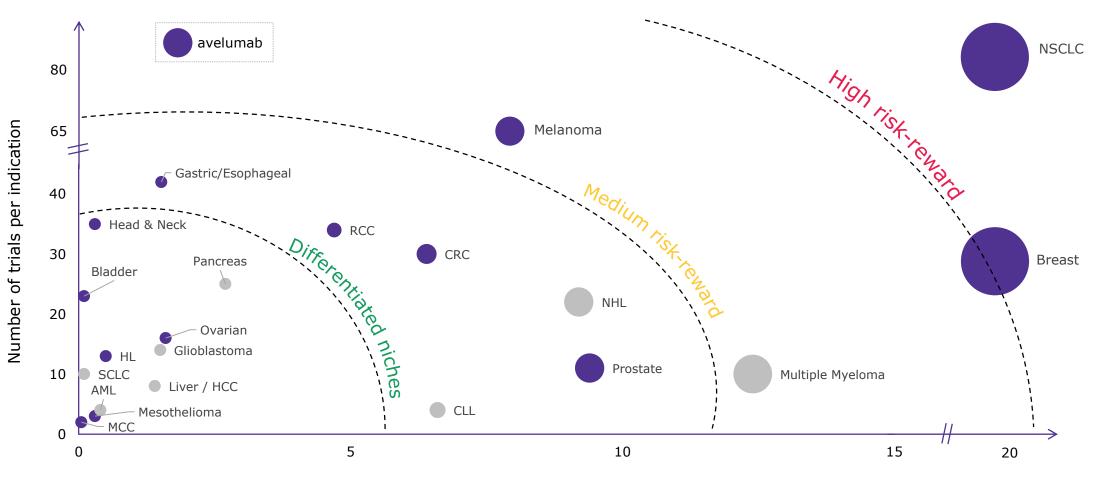
**Cladribine Tablets**<sup>5</sup> – **Lymphocyte targeting agent** Relapsing-remitting multiple sclerosis

- Neurodegenerative Diseases
   Oncology
   Immunology
   Immuno-Oncology
- Biosimilars

<sup>1</sup>Sponsored by the National Cancer Institute (USA); <sup>2</sup>1st line treatment; <sup>3</sup>2nd line treatment; <sup>4</sup>3rd line treatment <sup>5</sup>As announced on July 18<sup>th</sup>, 2016 the European Medicines Agency accepted Merck KGaA, Darmstadt, Germany, Marketing Authorization Application



## Avelumab plays predominantly in attractive and differentiated niches



Market size in 2020 per indication [€bn]

Sources: Trialtrove and Cortellis as of September 2015, Boston Consulting Group, Evaluate Pharma forecast 2020 Acronyms: SCLC = Small Cell Lung Cancer; HL = Hodgkins Lymphoma; NHL = Non Hodgkins Lymphoma; AML = Acute Myeloid Leukaemia



# **Avelumab – Differentiation strategy varies according to chosen target indication and market**

#### Ambition: Smart leader

- Indications (Merkel cell) or markets (Asia for gastric)
- Quick to market strategy, e.g. BTD for MCC in November 2015
- Small, but less crowded markets and sales potential with notable impact for us
- Strategic strength of Healthcare in niche markets

#### Ambition: Smart follower

Saturated and / or major indications

Unsaturated

and / or niche

indications

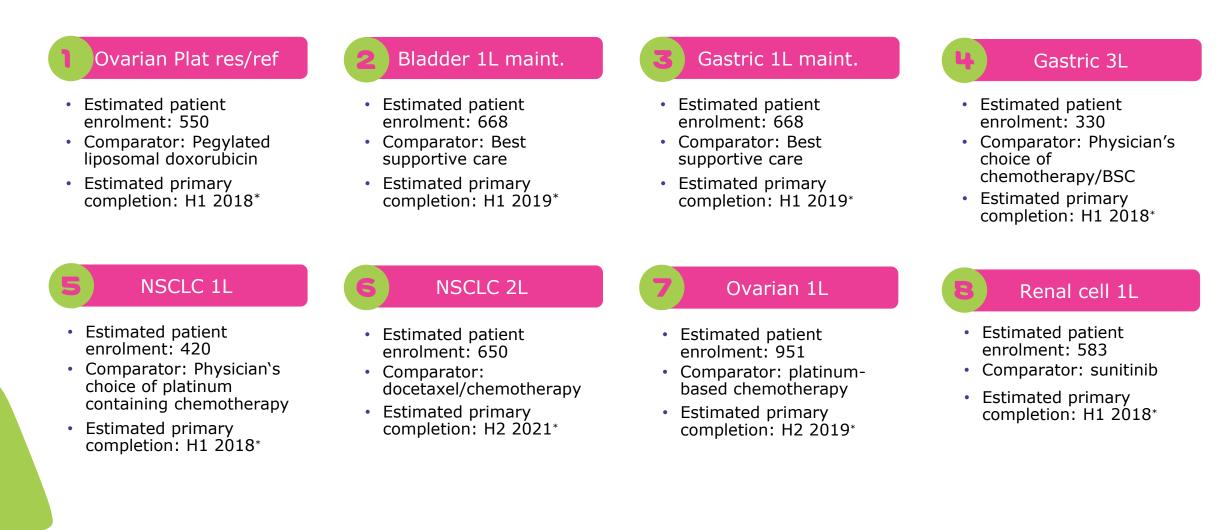
- Indications such as NSCLC or Bladder
- Learn from experience of incumbents/early movers
- Potential for combinations given breadth of combined development pipelines, e.g. lung
- Differentiate in trial design and explore application of further biomarkers





48

# The alliance initiated 8 Phase III studies





49

# MCC 2L: Clinical results support avelumab as potential therapeutic option – planned to apply for marketing authorization in H2 2016

#### Encouraging response rates<sup>1</sup>

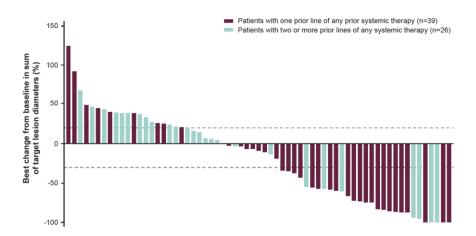
- ORR: 31.8%
  - 9.1% complete response
  - 22.7% partial response
  - Rapid (78.6% responding within 7 weeks of treatment)
  - Durable (82.1% still responding at time of analysis)
- 6-mo OS: 69% (median OS: 11.3 months)
- 6-mo PFS rate: 40%
- Manageable safety profile; no unexpected safety signals





#### Potential for differentiation

- Largest international multicenter, open-label study of anti-PD-L1/PD-1 reported in this patient population (88 patients) – Responses observed in large number of patients
- Improved response rates observed when used earlier, i.e. fewer lines of prior chemotherapy appeared to be associated with better response to avelumab in MCC 2L and beyond
  - ORR of 40.4% for patients with one prior systematic treatment
  - ORR of 19.4% for patients with two and more prior treatments



Note: timelines are event-driven and may change Note: avelumab is the proposed nonproprietary name for the anti-PD-L1 monoclonal antibody (MSB0010718C) <sup>1</sup>Avelumab (MSB0010718C; anti-PD-L1) in patients with metastatic Merkel cell carcinoma previously treated with chemotherapy: results of the phase 2 JAVELIN Merkel 200 trial\* / Oral Presentation at the 52nd ASCO Annual Meeting, June 3-7, 2016; Chicago, Illinois. Abstract No. 9508; Howard Kaufman et al.



# Going forward, avelumab combinations will drive differentiation strategy



- Phase II 2L MCC (BTD, ODD and FTD)
- Phase III 1L and 2L Plat res/ref ovarian
- Phase III 1L MN and 3L gastric
- Phase III 1L and 2L NSCLC
- Phase III 1L MN bladder
- Phase I Hodgkins Lymphoma
- Multiple other tumor types



- Phase III, RCC 1L
- Phase Ib/II, NSCLC 1L ALK+
- Phase I/II
- Phase Ib/II, ovarian
- Phase I/Ib, ovarian

- (avelumab + Inlyta) (avelumab + Xalkori/lorlatinib) (avelumab + 4-1BB) (avelumab + Entinostat; Syndax collaboration) (avelumab + VS-6063; Verastem collaboration)
- Further combination trials under consideration



# **Cladribine tablets – MAA submission accepted by EMA in July 2016**

#### Background

- Targets lymphocytes (both B and T cells), integral to MS pathogenesis
- Two Phase III and one Phase IIIb extension studies conducted in RRMS and early MS<sup>1,2,3</sup>; Phase II study in patients failing IFN beta therapy<sup>5</sup>
- Substantial new efficacy & safety characterization including data from long-term follow up (>10,000 patient-years)
- Most recent analyses provide relevant information on benefit/risk profile of cladribine tablets in RRMS:
  - ARR reduction (58%)
  - Risk of disability progression (33% reduction)
  - Relative reduction in mean number of lesion (86% reduction in T1 gadolinium-enhanced lesions)
  - 47% of patients experience NEDA over 2 years<sup>4</sup>

#### Potential for differentiation

- We aim to address significant unmet needs for agents delivering high efficacy with favorable safety profile in a convenient dosing regimen
- Administered orally (tablet formulation)
- Extremely short treatment courses (8–10 days per year) leading to long-term efficacy<sup>1</sup>

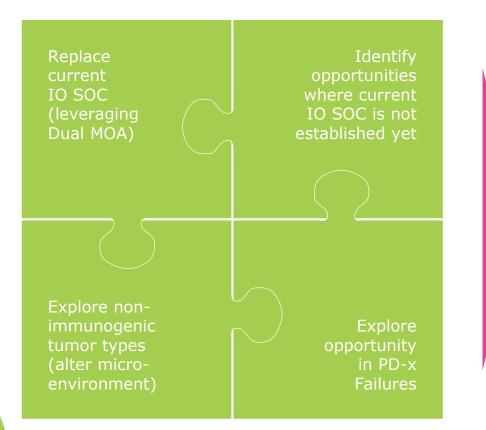
#### Note: timelines are event-driven and may change

EMA = European Medicines Agency; ARR = Annualized Relapse Rate; MAA = Marketing Authorization Application; MS = multiple sclerosis; NEDA = no evidence of disease activity; RRMS = relapsing-remitting multiple sclerosis. <sup>1</sup> Giovannoni G et al. New Engl J Med 2010;362:416–26; <sup>2</sup> Giovannoni G et al. 65th annual meeting of the American Academy of Neurology 2013. P07.119. <sup>3</sup> Leist TP et al. Lancet Neurol 2014;13:257–67. <sup>4</sup> Giovannoni G et al. Lancet Neurol. 2011;10:329–37. <sup>5</sup> Montalban X et al. 65th annual meeting of the American Academy of Neurology 2013. P07.099.



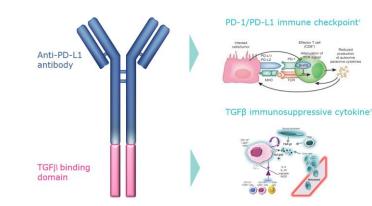
# **PD-L1–TGF-beta indicates potential to move beyond checkpoint inhibitors**

#### Four focus areas for exploration



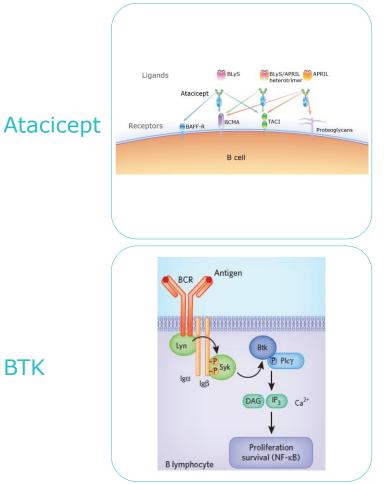
#### Status and next steps

- Novel, first-in-class bifunctional immunotherapy
- Bifunctional mode should result in broader application vs. respective mono-functional agents
- Great potential when combined with Standard of Care, immunotherapy and internal pipeline drug candidates
- Dose level finding of Phase I completed
- Expansion into Ib cohorts expected for Q3 2016





# Update on selected assets (1/2)



- Binds to receptors of two cytokines regulating maturation, function, and survival of B cells (B-lymphocyte stimulator (BLyS) & a proliferation-inducing ligand (APRIL))
- ADDRESS II (Ph IIb) in SLE patients aiming to show reduction in disease activity – 279 patients enrolled
- 24-week, randomized, double-blind, placebo-controlled Subcutaneous injection, once-a-week dosing
- Primary outcome: Percentage of patients with SLE responder index (SRI) response at week 24 compared to screening
- Suppress autoantibody-producing cells
- Preclinical research suggests therapeutic use in certain autoimmune diseases
- High and differentiated efficacy in preclinical models; promising kinase selectivity profile
- Aim to achieve best in class through minimization of off-target effects
- 2<sup>nd</sup> dose level of Phase I completed
- Partnering opportunities under consideration

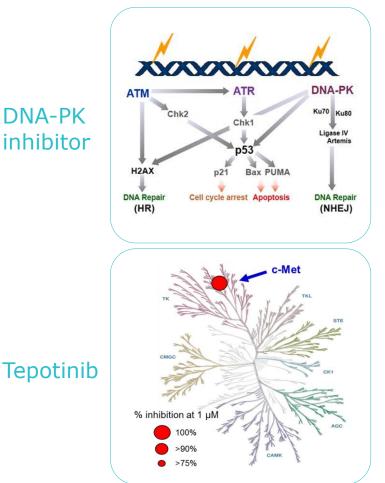
#### Phase III decision expected in H2 2016

Three phase II trials expected to be started until end of 2016 (e.g. RA, SLE)



54

# Update on selected assets (2/2)



- M3814 is a selective and potent inhibitor of DNA-PK, a kinase mediating DNA double strand break repair<sup>1</sup>
- Preclinical PoC showing complete responses and/or increased PFS in combination with radiotherapy in several xenograft models (SCCHN, NSCLC, CRC, PaCa) and strong pre-clinical combination data with SoC chemotherapies
- Two Phase Ia trials ongoing: FIM (monotherapy): 5<sup>th</sup> dose level completed, MTD not yet reached; RT combination: recruitment ongoing

Analysis of Phase I data for RT combination expected in H2 2017

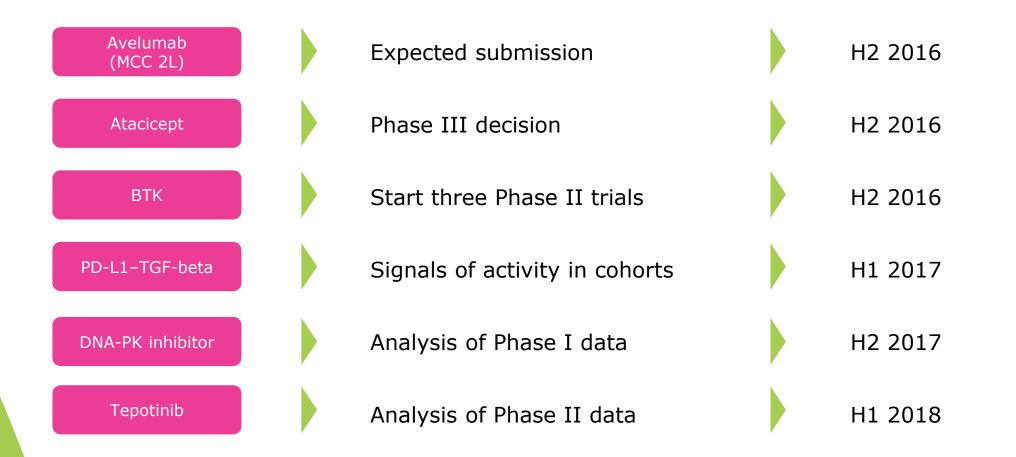
- Highly selective small molecule c-Met inhibitor
- Active in ligand-dependent and ligand-independent tumor models
- Biomarker-driven approach for patient selection
- Preliminary data show encouraging signs of anti-tumor activity in c-Met positive patients in NSCLC and HCC<sup>2,3</sup>
- Phase II trials in progress in NSCLC and HCC

Analysis of Phase II data for HCC and NSCLC expected in H1 2018

Note: timelines are event-driven and may change

<sup>1</sup>Graphics only illustrative; Acronyms: SCCHN = Squamous Cell Carcinoma of the Head and Neck, NSCLC = Non-small Cell Lung Cancer, CRC = Colorectal Cancer, PaCa = Pancreatic Cancer, HCC = Hepatocellular Cancer, PFS = Progression-free Survival, SoC = Standard of Care, FIM = First-in-Man, RT = radiotherapy, CT = chemotherapy, MTD = maximum tolerated dose; <sup>2</sup>Qin, ECC 2015, (3) Kim et al, IASCL-WCLC 2015

# **Outlook – MCC submission planned in H2 2016**





Note: timelines are event-driven and may change Note: avelumab is the proposed nonproprietary name for the anti-PD-L1 monoclonal antibody (MSB0010718C)

# Healthcare is well set for future growth

Stable existing business

Business and market specific initiatives in place to maximize existing business franchises

Strong R&D pipeline Diversified but focused pipeline with high quality assets in the areas Immuno-Oncology, Oncology and Immunology healthily spread across all clinical phases

Successful collaborations

Proven success in partnering through joint investments and collaborations – maximizing potential of assets in competitive space

Promising late stage progress Two expected submissions in 2016 may potentially result in product launches in 2017

Disciplined execution

Systematic pipeline review and timely decision making allow efficient resource and budget allocation











# Performance Materials The four pillars are set for future profitable growth



#### ~50-60% of total sales,

- liquid crystals (LC) and photoresists for TVs, smartphones and tablet computers
- other display and non-display applications (e.g. LC Windows)

#### ~15-20% of total sales,

- effect pigments and functional materials for coatings, plastics, printing and cosmetics
- functional materials for cosmetics and special applications

#### ~15-20% of total sales,

- dielectrics, colloidal silica, lithography materials (photo resists), yield enhancers, edge-bead removers
- polyimide raw materials and printing materials

#### ~5-10% of total sales,

 organic (OLED) and inorganic (LED) light emitting diodes and functional materials for electronics and energy solutions, especially OLED is providing first substantial sales



# Performance Materials Innovation leadership is the backbone of future profitability and growth



Incremental product adjustments - only minor investments

LC	New singles for PSVA technology, new mode SA-VA
pigments	Meoxal luminous metal effect pigments
ICM	low defect CMP <sup>2</sup> slurries
advanced tech.	Organic Photovoltaics



#### Fundamental research - more substantial investments needed

LC	Smart windows, smart antennas
pigments	Counterfeiting prevention applications
ICM	Directed self-assembly (DSA) in lithography
advanced tech.	OLED printing technology



# Performance Materials Sound platform to deliver high earnings

#### Four-pillar platform diversifies earnings stream

- Liquid Crystals remain key earnings contributor
- AZ\* expertise is being leveraged to develop innovative value-added solutions for customers
- OLED is becoming a visible growth driver
- Pigments continue to grow with high-end products



#### **Continuous innovation as key profitability driver**

- New products contribute high growth and profitability
- LC\* technology mode UB-FFS\* launched in 2014 is the most recent example

#### **Balanced sales and consistently high earnings**



# We are the innovation leader IPS\* "Improved picture quality" 1996 VR\* "Large TVs" 2000 PS\* "Display cost reduction & advanced performance" 2008 UB\* "Superior image resolution and bower energy consumption" 2014

#### Diversification of portfolio and ongoing innovation lead to strong profitability

\*Abbreviations: AZ = AZ Electronics, LC = Liquid crystals, UB-FFS = Ultra Brightness Fringe Field Switching, IPS = In-Plane-Switching, VA = Vertical Alignment, PS-VA = Vertical Alignment with additional polymer layer fabricated from reactive mesogenes



First to commercializ

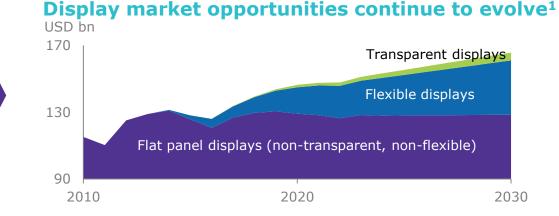
# **Performance Materials**

# Long-term growth and profitability drivers are intact



#### Macroeconomics and electronics remain buoyant

- Global consumer electronics market expected to grow above GDP\*
- Mobile data, Internet of Things and Big Data are key growth drivers for LC and IC
- Display market continues to grow

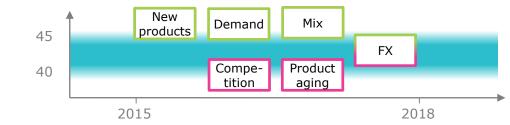


#### High value-added products yield superior profitability

- High market share in liquid crystals expected to prevail
- Strong differentiation by innovation inherent mature of business

#### **Sustainable profitability drivers**





Unique differentiation and market position will continue to lead to strong profitability and maintain low single-digit growth trajectory









# Q2 2016: Overview

#### Key figures

[€m]	Q2 2015	Q2 2016	Δ
Net sales	3,219	3,805	18.2%
EBITDA pre Margin (in % of net sales)	889 <i>27.9%</i>	<b>1,158</b> 30.4%	28.8%
EPS pre	1.30	1.55	19.2%
Operating cash flow	326	311	-4.7%
[€m]	Dec. 31, 2015	June 30, 2016	Δ
Net financial debt	12,654	12,510	-1.1%
Working capital	3,448	3,813	10.6%
Employees	49,613	50,456	1.7%

#### Comments

- EBITDA pre & margin increase driven by Sigma, organic performance and end of Rebif commission expenses
- EPS pre up due to EBITDA pre increase, but higher LTIP\* charges burden financial result
- Healthy operating cash flow due to strong business performance amid higher tax payments
- Net financial debt reflects operating cash flow and dividend payments
- Working capital shows increase in business activity – further room for improvement

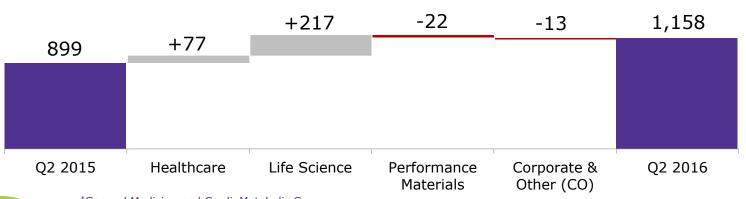


# Life Science and Healthcare drive increase in EBITDA pre

#### Q2 2016 YoY net sales

	Organic	Currency	Portfolio	Total
Healthcare	7.3%	-9.0%	-1.0%	-2.7%
Life Science	8.1%	-2.8%	79.7%	85.0%
Performance Materials	-4.7%	-2.0%	3.1%	-3.5%
Group	5.1%	-6.1%	19.2%	18.2%

#### Q2 YoY EBITDA pre contributors [€ m]



•Growth in Healthcare driven by strong Fertility, GM\* as well as Xalkori commissions

• Process Solutions continues to drive strong organic growth in Life Science

• Organic decline in Performance Materials reflects display supply chain destocking

Portfolio is Sigma and Kuvan

 •HC benefits from strong organic growth, end of Rebif commission expenses and ~€30 m disposal gain

• Life Science driven by Sigma, strong organic growth and synergies

• Performance Materials only slightly lower despite LC sales decline

•CO contains corporate initiatives & hedging

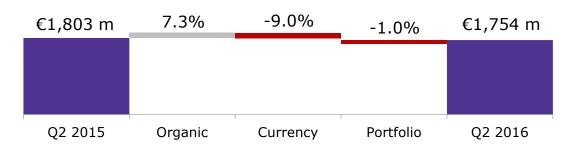


# Healthcare: Strong organic growth, EBITDA pre and profitability

#### Healthcare P&L

[€m]	Q2 2015	Q2 2016
Net sales	1,803	1,754
Marketing and selling	-730	-643
Administration	-69	-66
Research and development	-358	-378
EBIT	267	298
EBITDA	461	558
EBITDA pre	480	557
Margin (in % of net sales)	26.6%	31.8%

#### Net sales bridge

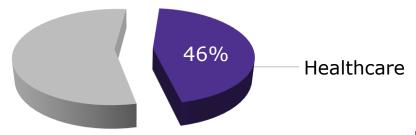


\*Productive Development Partnership Totals may not add up due to rounding

#### Comments

- Rebif organically stable; ramp-up of competition in Europe partially offset by tender in Russia; in U.S. pricing offsets declining volumes
- Solid organic growth of Erbitux across all regions, especially strong volume growth in China, but also low comparables
- Fertility shows strong growth across portfolio especially in China; Gonal-f benefiting from competitive situation in U.S. and low base
- Marketing & selling reflects end of commission expenses for Rebif (U.S.) partially offset by reinvestments in sales force & launch preparations
- R&D spend increases as pipeline development progresses
- Higher EBITDA pre due to strong organic growth, end of Rebif commissions and ~€30 m disposal gain from Venture Fund minority

#### Q2 2016 share of group net sales





# Healthcare organic growth by franchise/product

Q2 2016 organic sales growth [%] by key product [€ m]

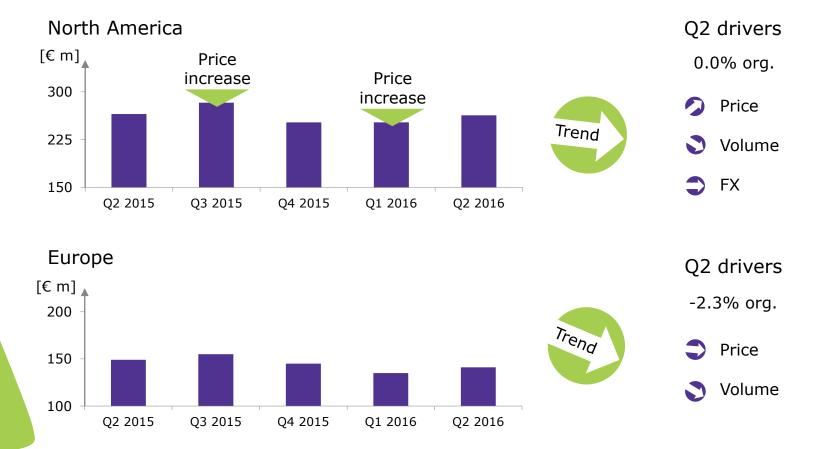
#### 441 **%**Rebif Rebif 863 0% -1% - - -461 891 232 438 **ERBITUX** +7% +5% 233 --438 Consumer Consumer 212 427 0% +3% 248 Health Health 479 209 396 +23% GONAL-GONAL-+20% 177 341 108 214 Concor +2% Concor\* +3% 132 253 104 197 Glucophage +10% Glucophage +5% 114 226 Q2 2016 Q2 2015 H1 2016 H1 2015

H1 2016 organic sales growth [%] by key product [€ m]



# **Rebif: Relief in the U.S. – competitive ramp-up in Europe ongoing**

#### Rebif sales evolution



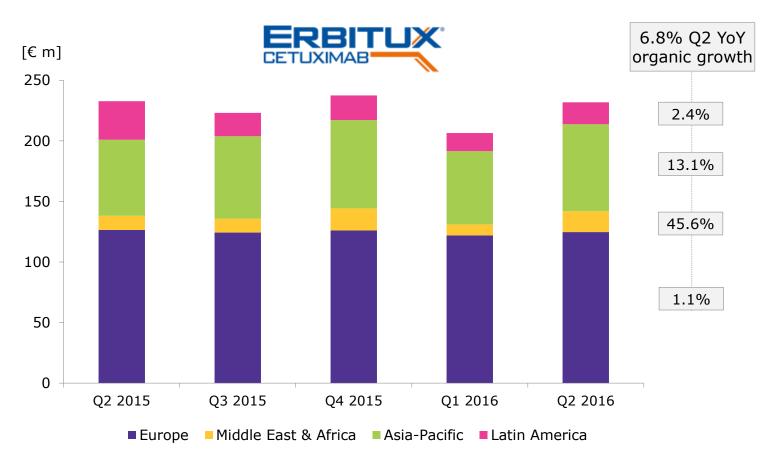
#### Rebif performance

- Rebif sales of €441 m in Q2 2016 reflect flat organic performance amid negative FX effects from LatAm & EU
- Market shares within interferons stable due to high retention rates and longterm safety track record
- •U.S. pricing & market share stabilization offset decline of interferon class
- 5% U.S. price increases in July will support performance going forward
- Phased market entry of orals in Europe causes ongoing volume decline;
   Q2 2016 contains tender in Russia



# **Erbitux: A challenging market environment**

#### Erbitux sales by region

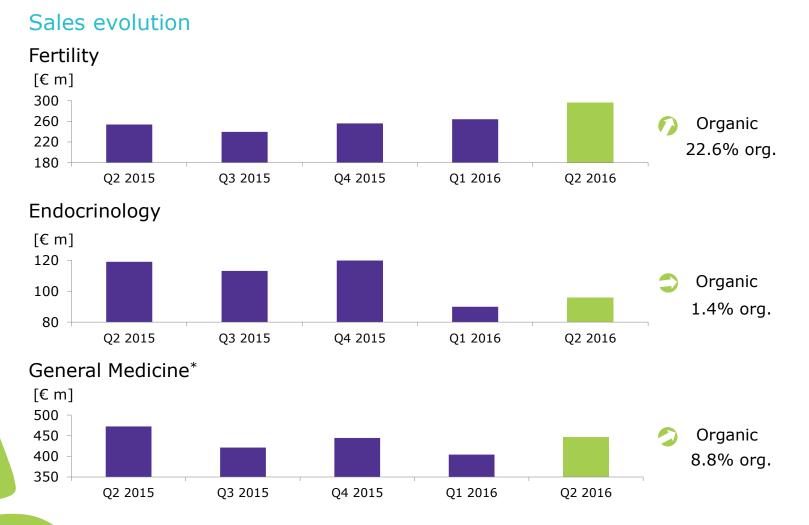


#### Erbitux performance

- Sales increase to €232 m due to solid volume development especially in growth markets
- Europe stable in ongoing tough environment (price and competition)
- Asia-Pacific shows strong growth led by China
- •Organic jump in MEA reflects growing demand but also beneficial tender phasing



# Strong organic growth in Fertility, General Medicine and Endocrinology



#### Q2 drivers

- •Gonal-f continues to benefit from competitive situation in U.S. and strong demand in China
- Sales drop in Endocrinology reflects Kuvan divestment; remaining portfolio (Saizen, Serostim) growing organically
- General Medicine sales burdened by FX headwinds from LatAM, organic performance sustainably healthy
- Euthyrox posts strong growth fueled by all regions, China remains key organic contributor of organic growth
- Glucophage growing in all regions especially benefiting from high demand in Middle East

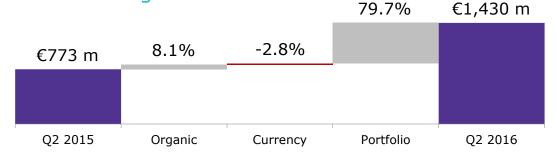


## Life Science: Another strong quarter while integration on track

#### Life Science P&L

[€m]	Q2 2015	Q2 2016
Net sales	773	1,430
Marketing and selling	-244	-413
Administration	-28	-58
Research and development	-49	-65
EBIT	87	166
EBITDA	170	343
EBITDA pre	200	417
Margin (in % of net sales)	25.9%	29.1%

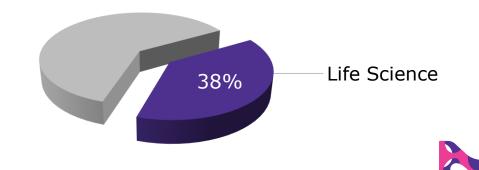
#### Net sales bridge



#### Comments

- Double-digit growth of Process Solutions driven by increasing production of large molecules across global and regional accounts
- Applied Solutions shows moderate organic growth, driven by biomonitoring products for pharma & demand for analytical testing
- Research Solutions benefits from strong demand in Emerging Markets, and for chemical analytics and molecular biology products
- Absolute costs higher due to Sigma, but improve in relation to sales
- Profitability reflects Sigma, business mix and synergies

#### Q2 2016 share of group net sales

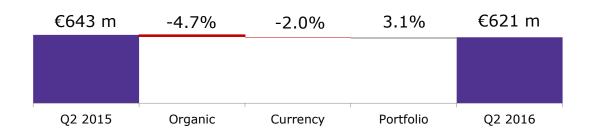


# Performance Materials: Strong profitability amid ongoing destocking

#### Performance Materials P&L

[€m]	Q2 2015	Q2 2016
Net sales	643	621
Marketing and selling	-53	-59
Administration	-14	-14
Research and development	-49	-53
EBIT	238	193
EBITDA	299	267
EBITDA pre	295	273
Margin (in % of net sales)	45.9%	44.1%

#### Net sales bridge

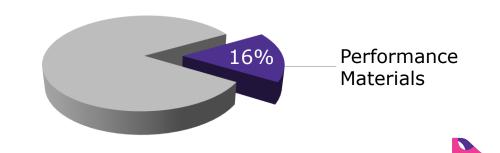


\*Active pharmaceutical ingredient Totals may not add up due to rounding

#### Comments

- LC declining as ongoing supply chain destocking & volume declines of mature TN-TFT outweighs volume increase in PS-VA and IPS
- Destocking is expected to continue into H2 2016
- OLED continues to grow on industry capacity expansion & investments
- Integrated Circuit Materials (ICM) shows above market growth mainly driven by dielectric and lithography materials for chip production
- Strong growth of Pigments & Functionals due to strong Xirallic for automotive coatings and cosmetic functionals, but on low comparables
- High profitability reflects leading market position, positive product mix within ICM and Pigments as well as active cost management

#### Q2 2016 share of group net sales



# **Reported figures reflect strong business performance**

#### Reported results

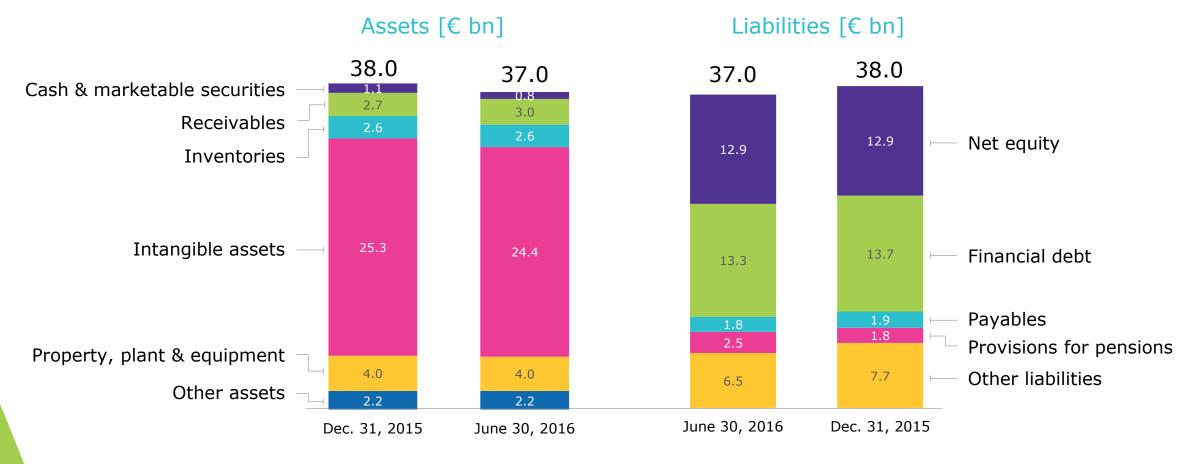
[€m]	Q2 2015	Q2 2016	Δ
EBIT	501	550	9.8%
Financial result	-41	-121	>100%
Profit before tax	461	429	-6.8%
Income tax	-115	-115	0.0%
<i>Effective tax rate (%)</i>	24.9%	26.7%	
Net income	343	312	-9.1%
EPS (€)	0.79	0.72	-8.9%

#### Comments

- •EBIT reflects increased EBITDA pre amid integration costs, Sigma D&A and Xalkori impairment (~€70 m)
- Financial result contains Sigma financing interest expenses and significant adverse effects from LTIP
- Effective tax rate slightly above guided range of ~23% to 25% due to Xalkori impairment



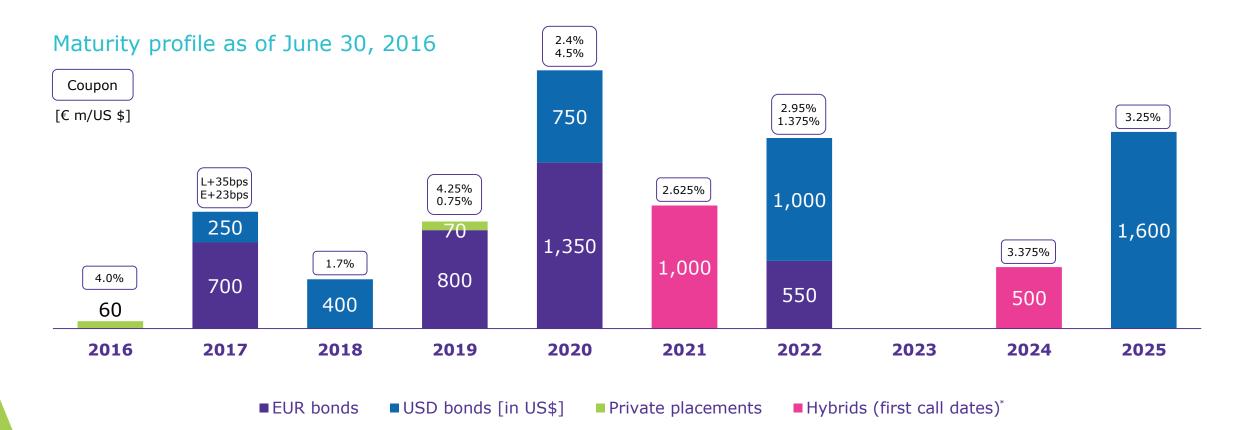
## **Balance sheet – deleveraging in progress after Sigma acquisition**



- Total assets decline by 2.6%, equity ratio grows to 34.7%
- Other liabilities decrease by €1.2 bn mainly due to dividend payment to E. Merck KG, Darmstadt, Germany and lower deferred taxes
- Further decline in interest rates increases pension provisions
- Stable net equity: Profit after tax offset by dividends, FX translation and actuarial losses



# Well-balanced maturity profile reflects capital market transactions related to Sigma-Aldrich



Financing structure enables flexible and swift deleveraging



# Healthy underlying operating cash flow

#### Q2 2016 – cash flow statement

[€m]	Q2 2015	Q2 2016	Δ
Profit after tax	346	314	-32
D&A	343	519	176
Changes in provisions	-70	-67	3
Changes in other assets/liabilities	-270	-397	-127
Other operating activities	3	-28	-31
Changes in working capital	-25	-30	-5
Operating cash flow	326	311	-15
Investing cash flow	1,860	-114	-1,974
thereof Capex on PPE	-93	-125	-32
Financing cash flow	-174	-357	-183

#### Cash flow drivers

- •D&A increases due to Sigma and Xalkori impairment
- Higher tax payments burden changes in other assets/liabilities
- Investing cash flow LY contained cash-in from Sigma hedging
- Capex higher due to HQ & Sigma; 2016 peak expected in Q4
- Financing cash flow reflects repayment of Millipore bond and commercial paper



# **Exceptionals in Q2 2016**

### Exceptionals in EBIT

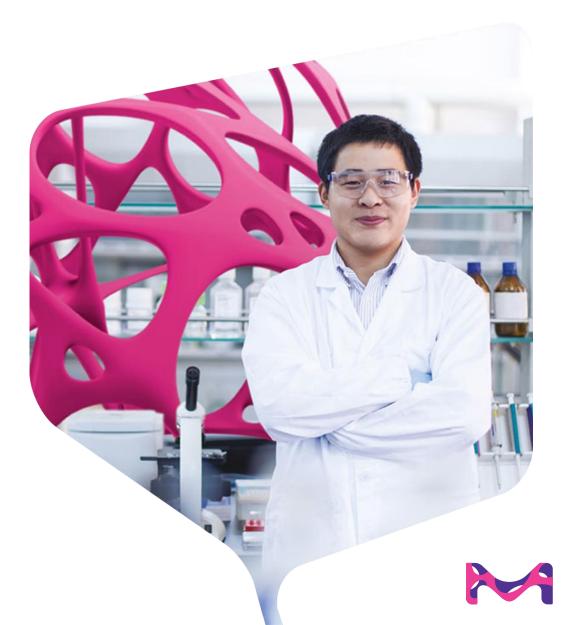
[€m]	Q2 2015		Q2 20	016
	Exceptionals	thereof D&A	Exceptionals	thereof D&A
Healthcare	21	2	70	71
Life Science	30	0	74	0
Performance Materials	-3	0	7	0
Corporate & Other	9	0	10	0
Total	56	2	160	71





# **Financial calendar**

Date	Event
October 13, 2016	Capital Markets Day - 2016
November 15, 2016	Q3 2016 Earnings release
March 9, 2017	Q4 2016 Earnings release
April 28, 2017	Annual General Meeting
May 18, 2017	Q1 2017 Earnings release



#### CONSTANTIN FEST



Head of Investor Relations +49 6151 72-5271 constantin.fest@emdgroup.com

#### ANNETT WEBER



Institutional Investors / Analysts +49 6151 72-63723 annett.weber@emdgroup.com

#### EVA STERZEL



Private Investors / AGM / CMDs / IR Media +49 6151 72-5355 eva.sterzel@emdgroup.com

#### SVENJA BUNDSCHUH



Assistant Investor Relations +49 6151 72-3744 svenja.bundschuh@emdgroup.com

#### NILS KUSTUSCH



Institutional Investors / Analysts +49 6151 72-7434 nils.kustusch@emdgroup.com

#### OLLIVER LETTAU



Institutional Investors / Analysts +49 6151 72-34409 olliver.lettau@emdgroup.com

#### **ALESSANDRA HEINZ**



Assistant Investor Relations +49 6151 72-3321 alessandra.heinz@emdgroup.com

#### **EMAIL:** <u>investor.relations@emdgroup.com</u> **WEB:** <u>www.emdgroup.com/investors</u> **FAX:** +49 6151 72-913321

