GOOD START INTO THE YEAR

Merck Q1 2016 results

Marcus Kuhnert, CFO
Belén Garijo, CEO Healthcare

May 19, 2016

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,” “would,” “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 product-related and patent law disputes; risks from antitrust law proceedings; risks from drug pricing by the divested Generics Group; risks in human resources; 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 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

01 Executive summary

02 Financial overview

03 Guidance

01 EXECUTIVE SUMMARY

MERCK

Highlights

Operations

- Healthcare – good organic growth and first avelumab Phase III combination trial
- Life Science – strong performance continues; Sigma integration on track
- Performance Materials – healthy profitability amid first supply chain destocking

Financials

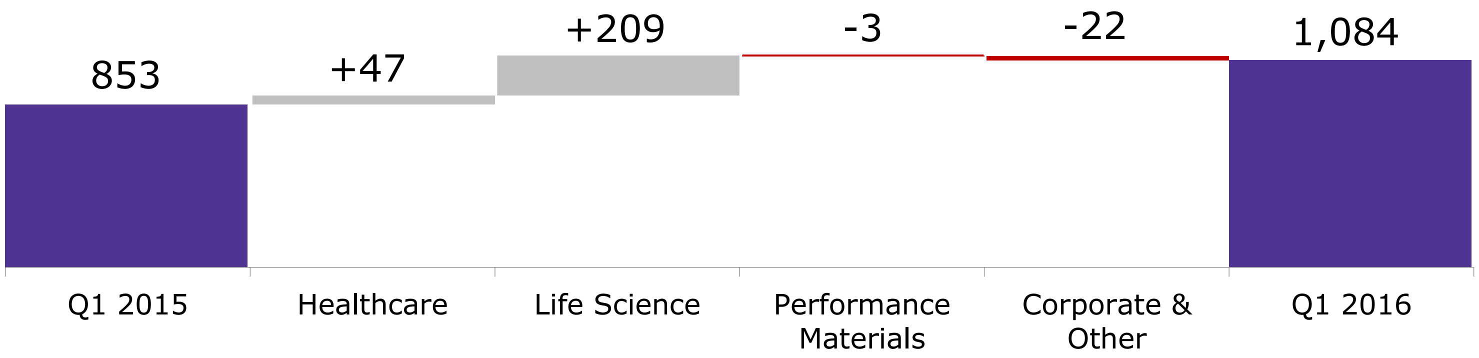
- Acquisition-driven sales growth of 20.5%; EBITDA pre up 27% to €1,084 m
- Deleveraging on track – net financial debt decreases by nearly €600 m
- FY 2016 guidance – net sales: €14.8 – 15.0 bn & EBITDA pre: €4,100 – 4,300 m

Life Science and Healthcare drive increase in EBITDA pre

Q1 2016 YoY net sales

	Organic	Currency	Portfolio	Total
Healthcare	5.4%	-6.8%	-1.0%	**-2.4%**
Life Science	8.9%	-1.3%	81.6%	**89.3%**
Performance Materials	-2.4%	0.5%	2.7%	**0.9%**
Merck Group	4.7%	-4.0%	19.8%	**20.5%**

Q1 YoY EBITDA pre contributors [€ m]

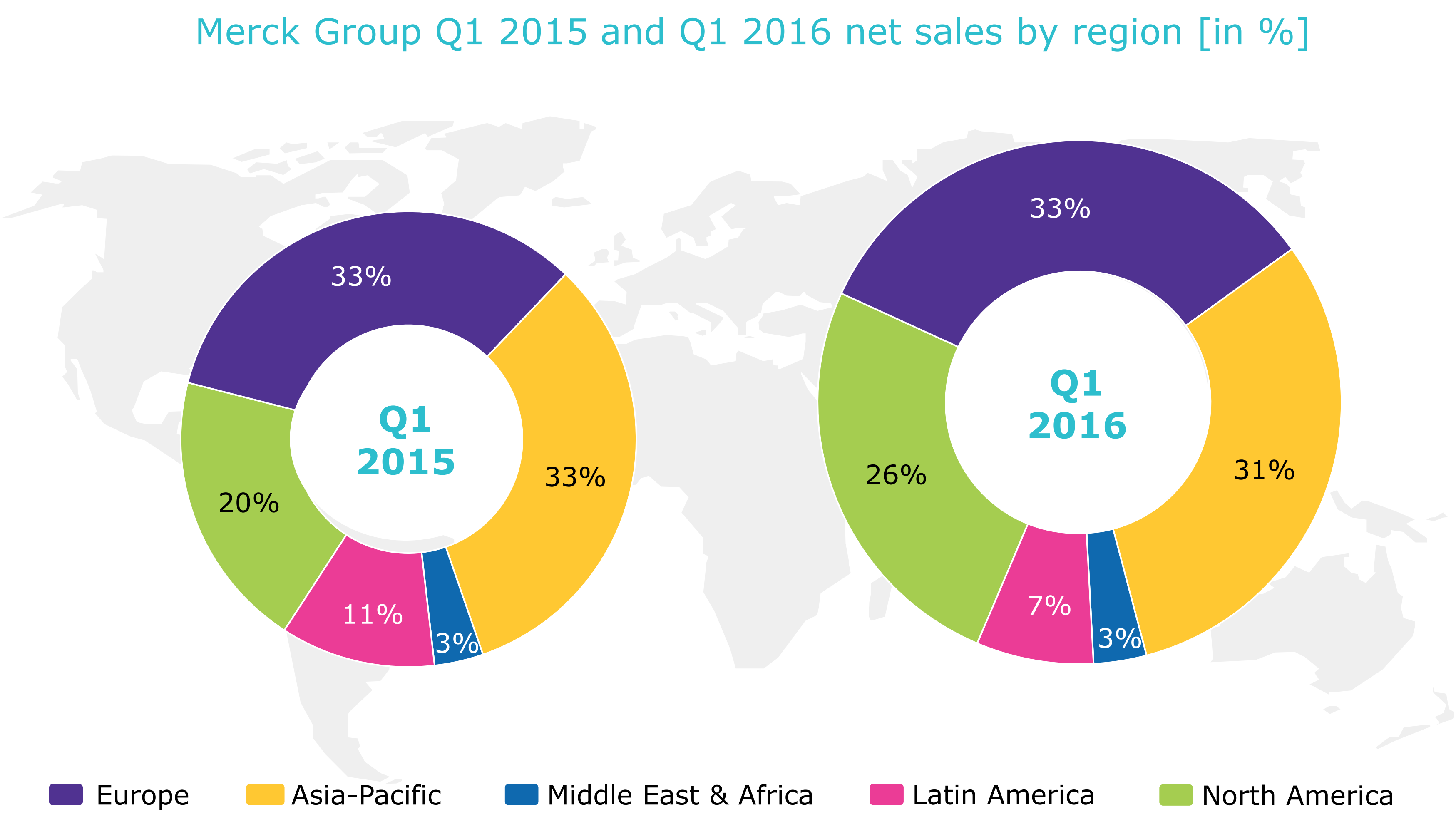


- Healthcare growth reflects strong Fertility, GM* and CH as well as Xalkori commissions
- Strong organic growth in Life Science driven by Process Solutions
- Slight organic decline in Performance Materials confirms expected destocking in display supply chain
- Portfolio reflects Sigma and Kuvan

- HC benefits from solid organic growth, Rebif commission savings and R&D phasing
- Life Science driven by Sigma, strong organic growth and positive product mix
- Performance Materials slightly lower due to product mix
- Corporate EBITDA pre contains hedging and investments in corporate initiatives

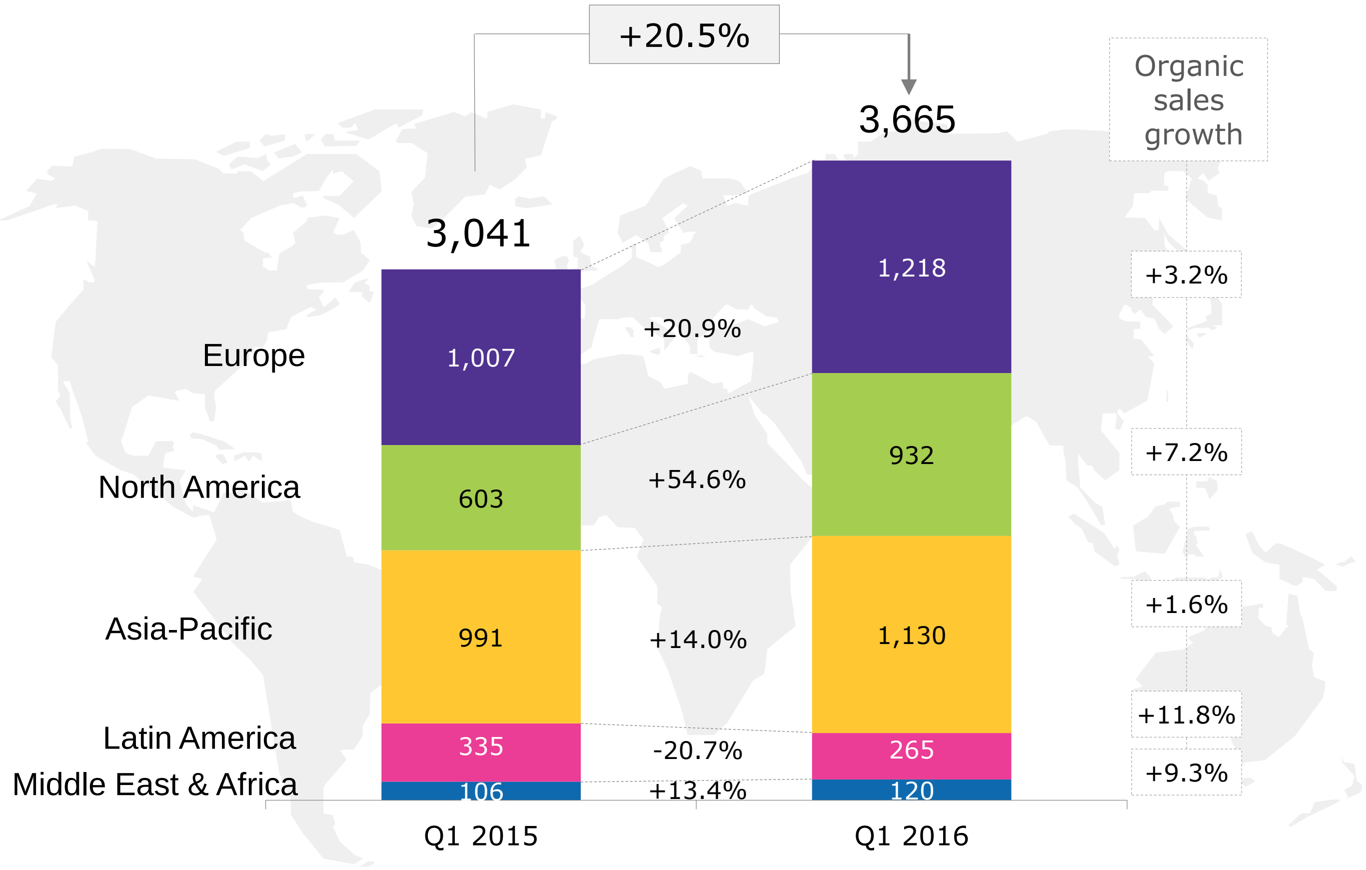
*General Medicine and CardioMetabolic Care
Totals may not add up due to rounding

Q1 2016 sales split reflects Sigma's footprint in North America



Organic growth in all regions

Regional breakdown of net sales [€ m]



Regional development

- Organic growth in Europe driven by Life Science, partially offset by Rebif
- North America benefits from strong Fertility franchise and Life Science as well as Rebif price increases
- Asia-Pacific shows slight organic growth as good demand in Fertility is offset by display supply chain destocking
- Strong organic growth in LatAm driven by Consumer Health and new reimbursement for Rebif in Brazil

Totals may not add up due to rounding

02 FINANCIAL OVERVIEW

MERCK

Q1 2016: Overview

Key figures

[€m]	Q1 2015	Q1 2016	Δ
Net sales	3,041	**3,665**	20.5%
EBITDA pre	853	**1,084**	27.0%
Margin (in % of net sales)	*28.0%*	*29.6%*	
EPS pre	1.12	**1.54**	37.5%
Operating cash flow	279	**352**	26.5%

[€m]	Dec. 31, 2015	March 31, 2016	Δ
Net financial debt	12,654	**12,072**	-4.6%
Working capital	3,448	**3,726**	8.0%
Employees	49,613	**50,262**	1.3%

Comments

- EBITDA pre & margin increase driven by Sigma, end of Rebif commission expenses and organic performance
- Strong EPS pre growth due to higher EBITDA pre; LY financial result burdened by LTIP* charges
- Operating cash flow reflects strong business performance; LY impacted by one-time tax payment
- Net financial debt reduction driven by cash-in for Kuvan & operating cash flow
- Working capital reflects increase in business activity

*Long Term Incentive Plan
Totals may not add up due to rounding

MERCK

Reported figures reflect solid business performance and Kuvan divestment

Reported results

[€m]	Q1 2015	Q1 2016	Δ
EBIT	480	**849**	76.8%
Financial result	-101	**-68**	-32.1%
Profit before tax	379	**780**	>100%
Income tax	-94	**-187**	99.2%
Tax rate (%)	*24.8%*	***24.0%***	
Net income	282	**591**	>100%
EPS (€)	0.65	**1.36**	>100%

Comments

- EBIT reflects increased EBITDA pre and Kuvan disposal gain amid integration costs and D&A from Sigma
- Financial result contains Sigma financing interest expenses; last year adversely affected by LTIP
- Tax rate within guidance range of ~23-25%

Healthcare: Good organic sales growth amid ramp-up in R&D investments

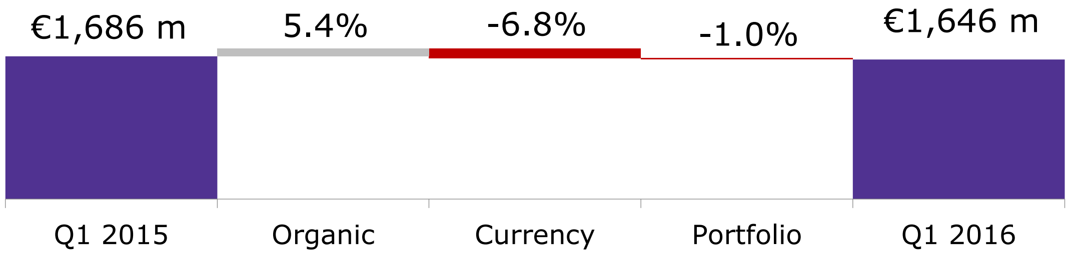
Healthcare P&L

[€m]	Q1 2015	Q1 2016
Net sales	1,686	**1,646**
Marketing and selling	-660	**-613**
Administration	-66	**-71**
Research and development	-348	**-378**
EBIT	268	**641**
EBITDA	449	**829**
EBITDA pre	461	**508**
Margin (in % of net sales)	*27.3%*	***30.9%***

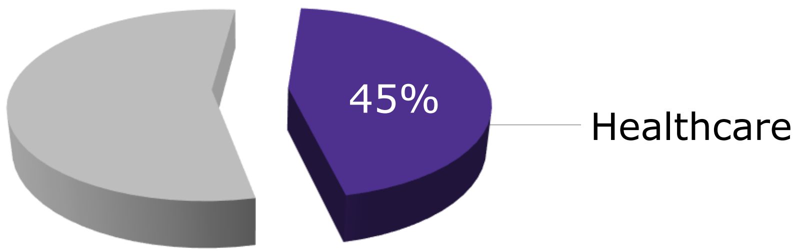
Comments

- Organic growth driven by strong Fertility, GM, CH and Xalkori
- Rebif still impacted by ramp-up of orals in Europe, while U.S. pricing and first contribution from PDP* in Brazil support performance
- Erbitux shows moderate organic growth benefiting from pick-up in Brazil, but also low comparables; EU remains competitive
- Marketing & selling reflect end of commission expenses for Rebif (U.S.) partially offset by reinvestments in sales force & launch preparations
- R&D spend starts to increase as avelumab Phase III trials progress
- EBIT reflects Kuvan disposal gain of €324 m
- Profitability improves mainly due to good organic growth, ceased Rebif commission expenses amid phased R&D cost ramp-up

Net sales bridge



Q1 2016 share of group net sales

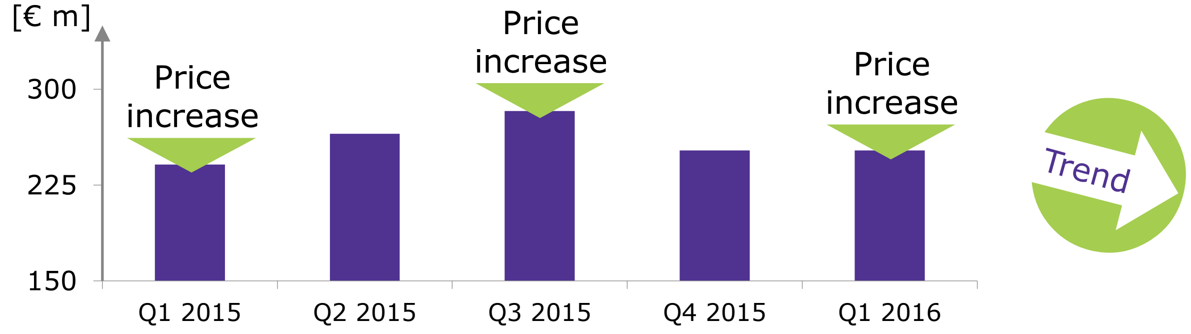


*Productive Development Partnership
Totals may not add up due to rounding

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

Rebif sales evolution

North America

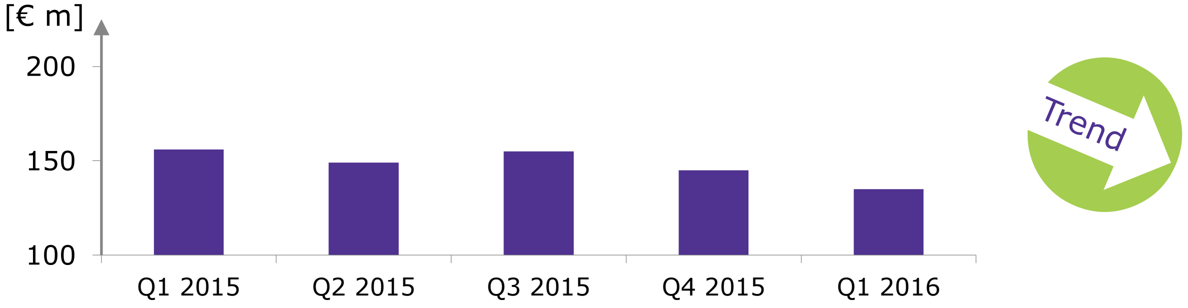


Q1 drivers

1.6% org.

- Price
- Volume
- FX

Europe



Q1 drivers

-13.1% org.

- Price
- Volume

Rebif performance

- Rebif sales of €422 m in Q1 2016 reflect organic decline of -1.5% and negative FX effects from LatAm
- Market shares within interferons stable due to high retention rates and known long-term track record
- U.S. price increases and market share stabilization outweigh decline of interferon class
- Phased market entry of orals in Europe causes ongoing volume decline
- Initiation of Productive Development Partnership (PDP) in Brazil supports Rebif growth

Avelumab: JAVELIN clinical development program initiated as of Oct 2015 and continuously expanded to further indications in 2016

Trial led by Merck KGaA Darmstadt, Germany | Trial led by Pfizer | Both

Ongoing trials

Phase I

- Solid tumors – Dose escalation and expansion phase
  - Completed efficacy N=150 / cohort: NSCLC 1L; Gastric; mBreast; NSCLC 2L
  - Signal detection N=50 / cohort: CRC; CRPC; Ovarian; Melanoma; Urothelial/ Bladder; Meso-thelioma; ACC; RCC; Hodgkins Lymphoma[1]
  - Expanded efficacy N=100-150 / cohort: Ovarian; SCCHN; Bladder; Gastric 3L; RCC 1L/2L
- Combination trials
  - avelumab+4-1BB NSCLC, SCCHN, Melanoma[1]
  - avelumab+Inlyta RCC 1L[1]
  - avelumab+Xalkori/Iorlatinib NSCLC 1L ALK+

Registrational

- avelumab MCC 2L[2]
- avelumab NSCLC 1L (PDL1+)
- avelumab NSCLC 2L
- avelumab+Chemo Gastric 1L MN
- avelumab+Chemo Gastric 3L
- avelumab+Chemo Ovarian 2L Plat Res/Ref
- avelumab+Chemo Bladder 1L MN
- Started in 2016:
  - avelumab+Inlyta RCC 1L
  - avelumab+Chemo Ovarian 1L

Planned trials in 2016

Phase I

- Additional studies in planning

Registrational

- avelumab MCC 1L
- Additional studies in planning

Acronyms: CRC = colorectal cancer; CRPC = Castrate Resistant Prostrate Cancer; ACC = Adrenocortical Cancer; SCCHN = Squamous Cell Carcinoma of the Head and Neck, RCC = Renal Cell Carcinoma; Avelumab is an investigational agent. Avelumab is the proposed non-proprietary name for the anti-PD-L1 monoclonal antibody (MSB0010718C); [1]Trials currently initiating; [2]Phase II trial

MERCK

ASCO: We look forward to presenting the results from our avelumab trials at the upcoming scientific meeting in further detail*

1
MCC (2L+)

- Largest trial of a PD(L)1 agent in Merkel cell carcinoma as of today
- Clinical activity with durable responses in high unmet-need patient population (previously treated with chemotherapy)

2
NSCLC (1L)

- Single-agent avelumab showed clinical activity
- Higher trend in PD-L1+ patients suggested
- Phase 3 trial is underway (JAVELIN LUNG 100)

3
Ovarian

- Single-agent avelumab showed clinical activity in heavily pre-treated patients
- Largest reported dataset of patients with advanced ovarian cancer treated with anti-PD-L1 as of today
- Phase 3 trials are underway (JAVELIN OVARIAN 100 and 200)

4
Gastric (1L MN & 2L)

- Treatment with avelumab as a 1st line maintenance and 2nd line treatment showed clinical activity
- Largest reported dataset of patients with advanced gastric cancer treated with anti-PD-L1 as of today
- Disease control rate was also observed in a SwM (switch-maintenance therapy) group

5
Urothelial / Bladder

- Treatment with avelumab showed clinical activity
- Phase 3 trial is underway (JAVELIN BLADDER 100)

6
Mesothelioma

- Largest reported dataset of patients with advanced unresectable tumors treated with anti-PD-(L)1 as of today
- Treatment with avelumab showed clinical activity in pre-treated patients

7
mACC

- First study to date of an anti-PD-(L)1 agent in this rare tumor type (advanced adrenocortical carcinoma)
- Treatment with avelumab showed clinical activity in pre-treated patients

All cohorts are part of the Phase I trial solid tumors: JAVELIN Solid Tumor NCT01772004
*To date; source: ASCO abstracts

MERCK

Newsflow: Upcoming pipeline catalysts

avelumab	Data readout Phase II MCC	H1 2016
CLADRIBINE	Expected EMA filing	H1 2016
sprifermin	Data readout Phase II	H2 2016
atacicept	Data readout Phase II	H2 2016
M7824 (PD-L1 – TGF-beta)	Dose escalation outcome	H2 2016
tepotinib	Data readout Phase II	H1 2017

Life Science: A strong start to 2016

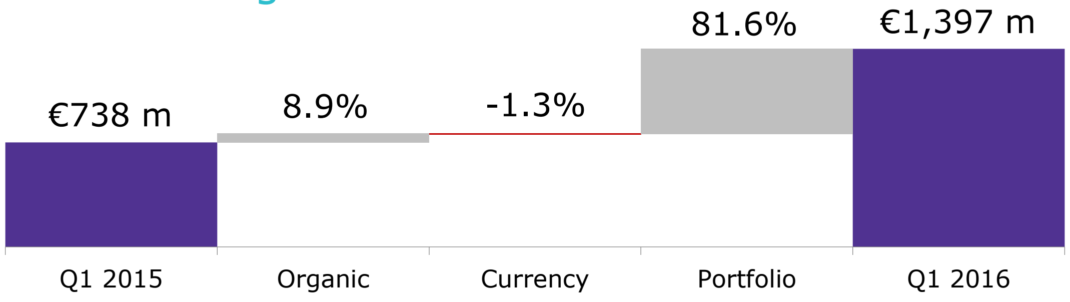
Life Science P&L

[€m]	Q1 2015	Q1 2016
Net sales	738	**1,397**
Marketing and selling	-233	**-421**
Administration	-31	**-63**
Research and development	-45	**-62**
EBIT	83	**105**
EBITDA	164	**284**
EBITDA pre	184	**393**
Margin (in % of net sales)	*25.0%*	***28.1%***

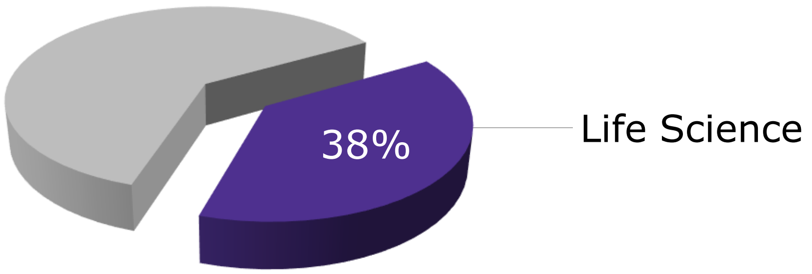
Comments

- Very strong growth of Process Solutions driven by increasing production of large molecules across global and regional accounts
- Applied Solutions shows moderate organic growth, driven by bio-monitoring products for pharma & demand for analytical testing
- Research Solutions benefits from increased research on biologics as well as solid demand for high-value chemicals
- Absolute costs higher due to Sigma, but improve in relation to sales
- Profitability reflects Sigma, favorable product mix and synergies

Net sales bridge



Q1 2016 share of group net sales



Totals may not add up due to rounding

Performance Materials: Healthy profitability amid destocking of display industry inventories

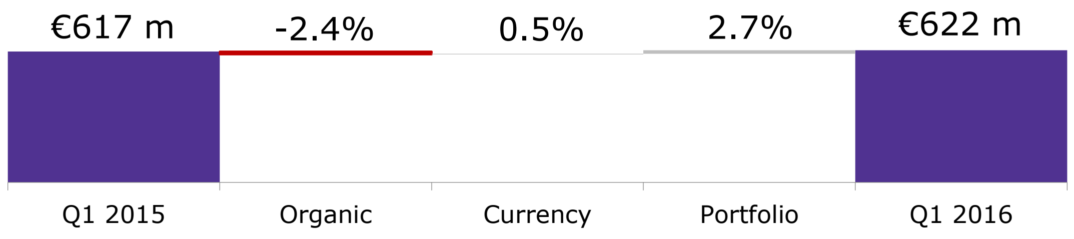
Performance Materials P&L

[€m]	Q1 2015	Q1 2016
Net sales	617	**622**
Marketing and selling	-46	**-58**
Administration	-18	**-16**
Research and development	-47	**-48**
EBIT	214	**207**
EBITDA	273	**267**
EBITDA pre	277	**273**
Margin (in % of net sales)	*44.8%*	***43.9%***

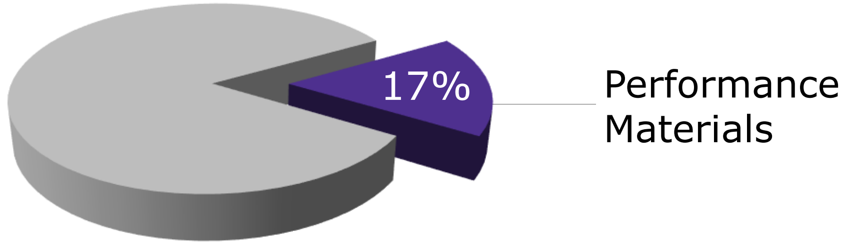
Comments

- Organic growth of OLED, ICM, and Pigments mitigates LC softness
- Liquid Crystals impacted by indicated inventory correction in supply chain
- OLED continues to grow on industry capacity expansion & investments
- Integrated Circuit Materials (ICM) show solid growth mainly driven by dielectric materials for chip production
- Pigments & Functionals post moderate growth esp. due to cosmetic API*
- Marketing & selling reflect contribution from Sigma's SAFC Hitech and sales force IT initiative
- Healthy profitability reflects leading market position with highly differentiated products, despite destocking and negative mix effects

Net sales bridge

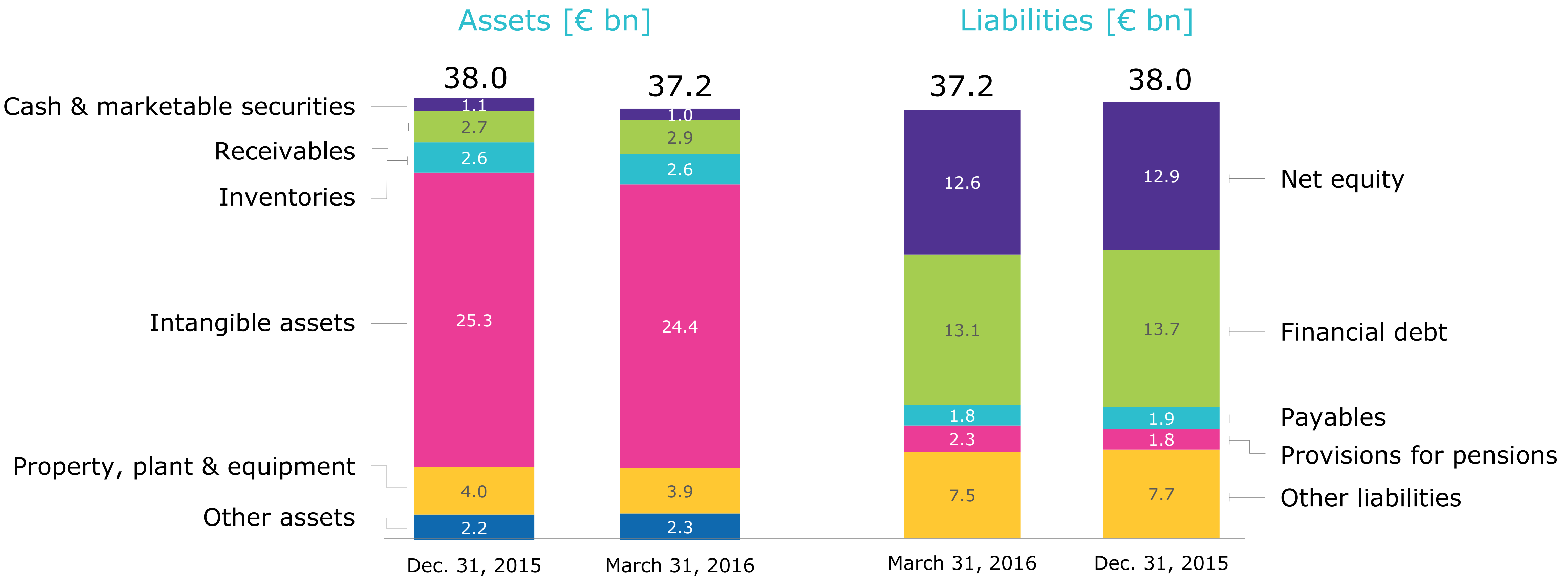


Q1 2016 share of group net sales



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

Balance sheet – deleveraging initiated after Sigma acquisition



- Total assets decline by 2.2%, equity ratio remains at 33.8%
- Reduction of intangible assets reflects D&A (€0.3 bn) and FX (€0.6 bn)
- FX development accounts for ~€0.6 bn total equity decrease
- Net financial debt reduced by €0.6 bn
- Decline of interest rates increases pension provisions

Totals may not add up due to rounding

Healthy operating cash flow enables first deleveraging measures

Q1 2016 – cash flow statement

[€m]	Q1 2015	Q1 2016	Δ
Profit after tax	285	**593**	308
D&A	325	**433**	108
Changes in provisions	90	**21**	-69
Changes in other assets/liabilities	-231	**-34**	197
Other operating activities	-20	**-394**	-374
Changes in working capital	-172	**-266**	-94
Operating cash flow	279	**352**	73
Investing cash flow	392	**284**	-108
thereof Capex on PPE	-75	**-160**	-85
Financing cash flow	2,288	**-572**	-2,860

Cash flow drivers

- Profit after tax includes gain from Kuvan divestment, which is neutralized in other operating activities
- D&A increases due to Sigma
- Tax payment (Pfizer upfront) burdened changes in other assets/liabilities LY
- Changes in working capital reflect lower payables (Pfizer-Rebif) and higher R&D receivables from Pfizer (avelumab)
- Investing cash flow contains increased Capex and Kuvan divestment; LY with sale of financial assets
- Financing cash flow reflects first re-payments of Sigma-related bank loans; LY contains USD bond issuance

Totals may not add up due to rounding

03 GUIDANCE

Full-year 2016 guidance

Merck guidance for 2016, including Sigma-Aldrich

Net sales: ~ €14.8 – 15.0 bn

EBITDA pre: ~ €4,100 – 4,300 m

EPS pre: ~ €5.65 – 6.00

2016 business sector guidance

Net sales

- Slight organic growth
- Organic Rebif decline
- Other franchises growing

EBITDA pre

~ €1,800 – 1,900m

Net sales

- Mid single-digit organic growth
- Main driver Process Solutions
- High double-digit contribution from Sigma

EBITDA pre

~ €1,620 – 1,670m

Net sales

- Organically about stable
- Including moderate destocking in liquid crystals
- Growing demand in all businesses

EBITDA pre

~ €1,100 – 1,150m

APPENDIX

Additional financial guidance 2016

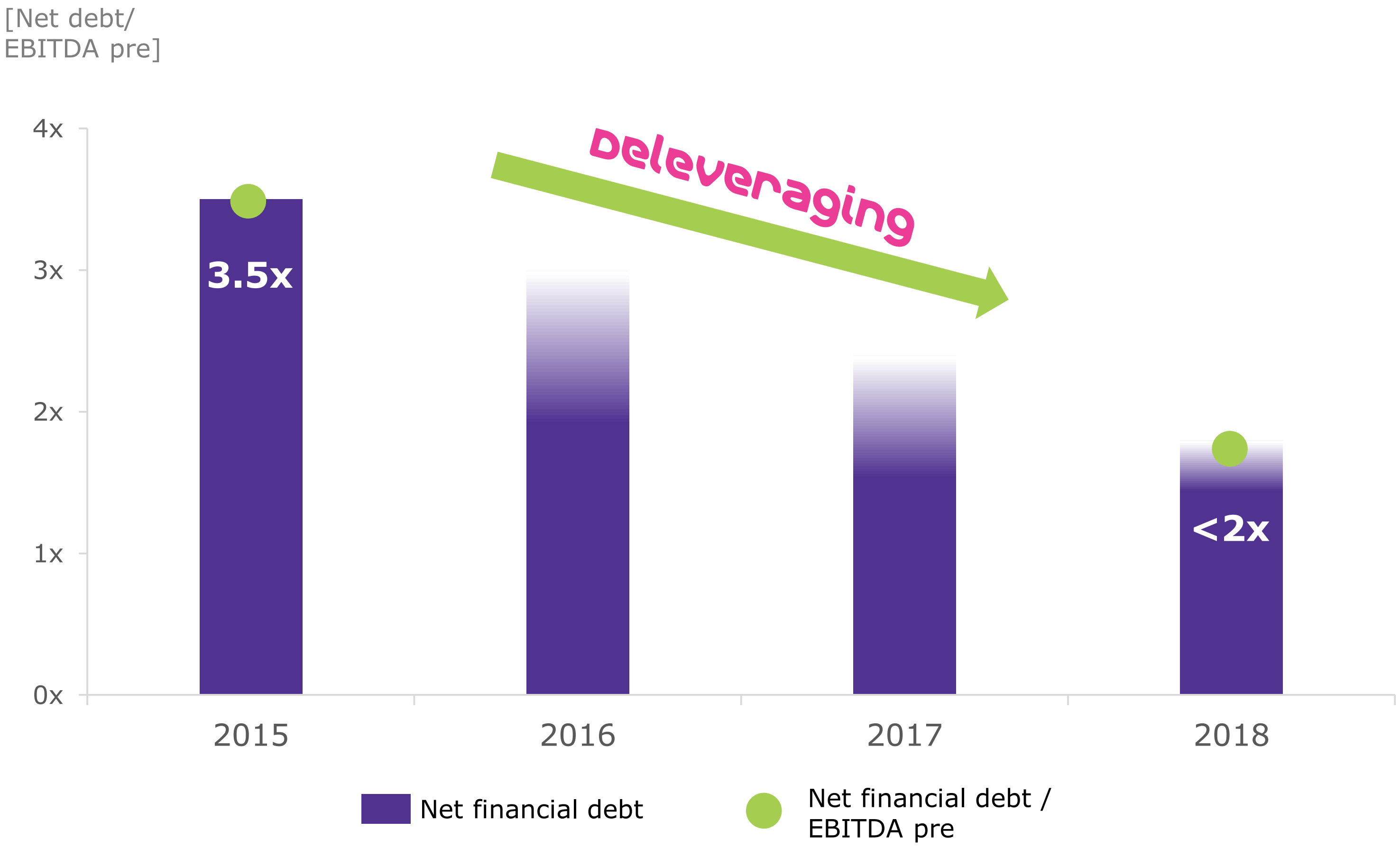
Further financial details

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 & 2017 hedge rate ~40-45% at EUR/USD ~1.10 to 1.15**
2016 Ø EUR/USD assumption	**~1.07 – 1.12**

MERCK

Strong focus on cash generation to ensure swift deleveraging

Net debt and leverage development

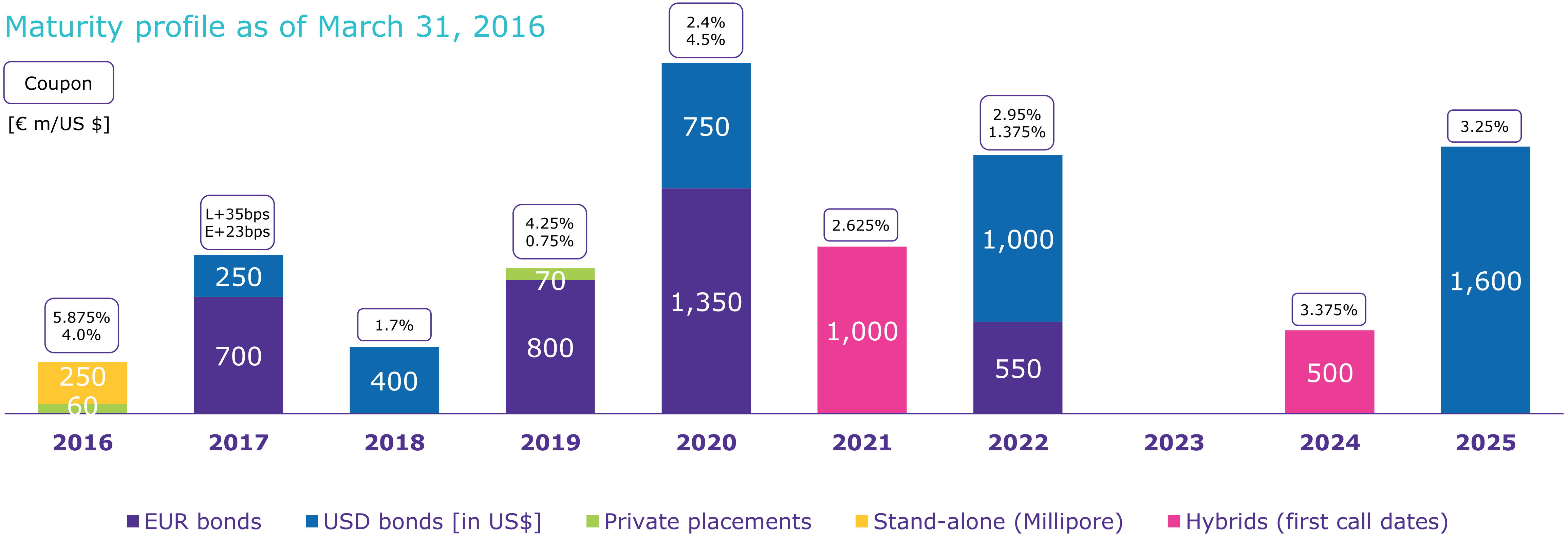


Focus on deleveraging

- Commitment to swift deleveraging to ensure a strong investment grade rating and financial flexibility
- Strong cash flow will be used to drive down leverage to expected <2x net debt/EBITDA pre in 2018
- Larger acquisitions (>€500 m) ruled out for the next two years
- Interest result in 2016: ~-€270 – -300 m

*Purchase price allocation

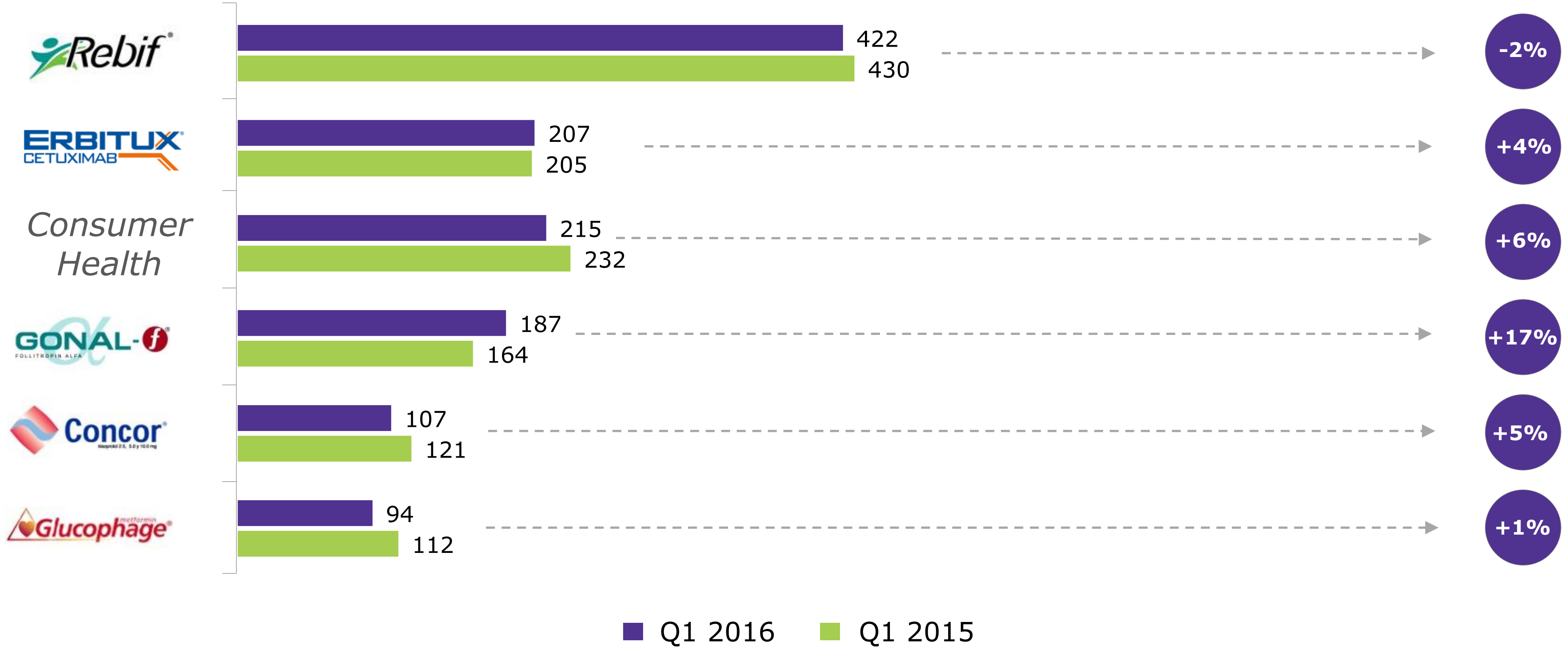
Well-balanced maturity profile reflects capital markets transactions related to Sigma-Aldrich



Financing structure enables flexible and swift deleveraging

Healthcare organic growth by franchise/product

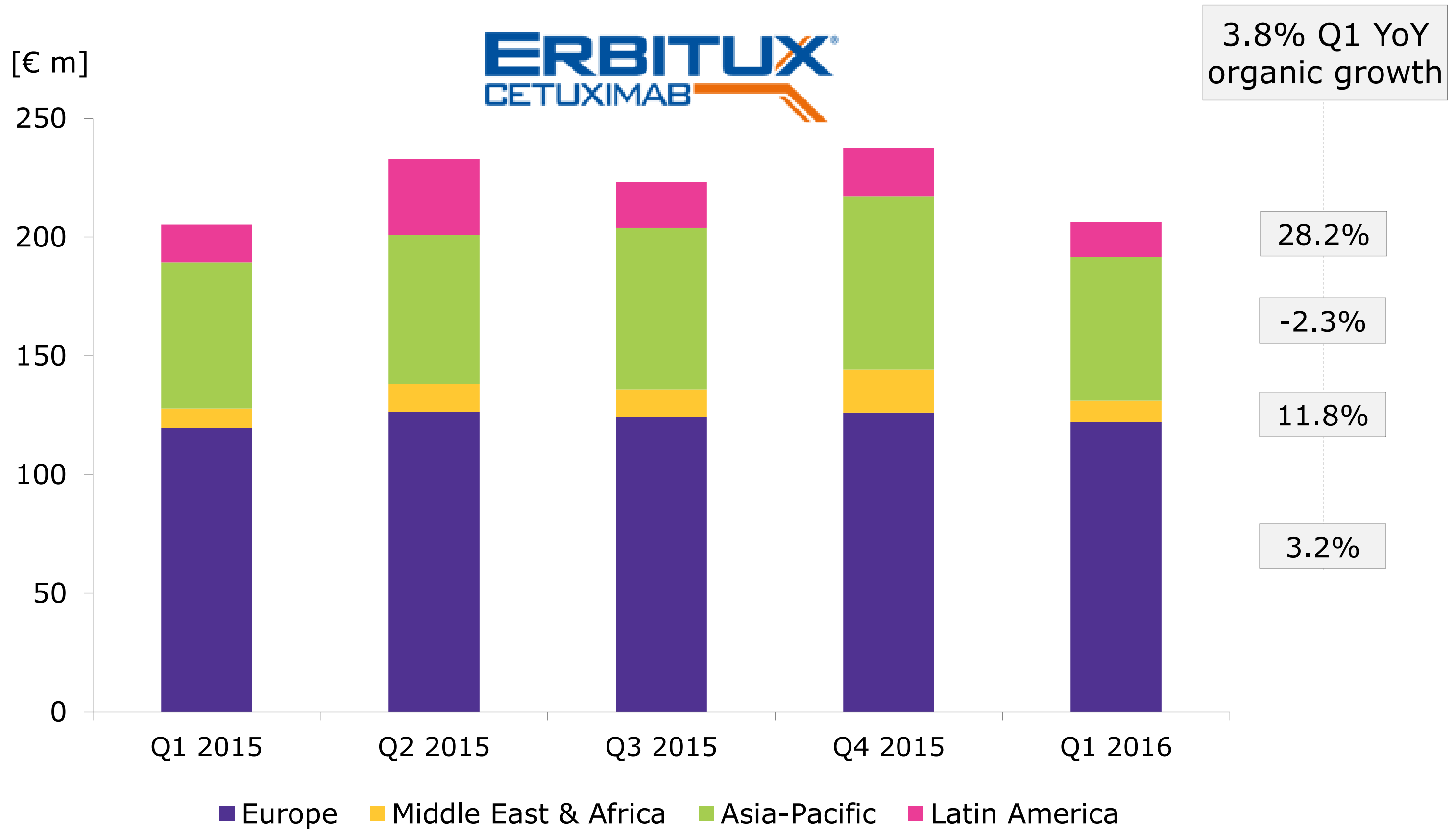
Q1 2016 organic sales growth [%] by key franchise/products [€ m]



Totals may not add up due to rounding

Erbitux: A challenging market environment

Erbitux sales by region



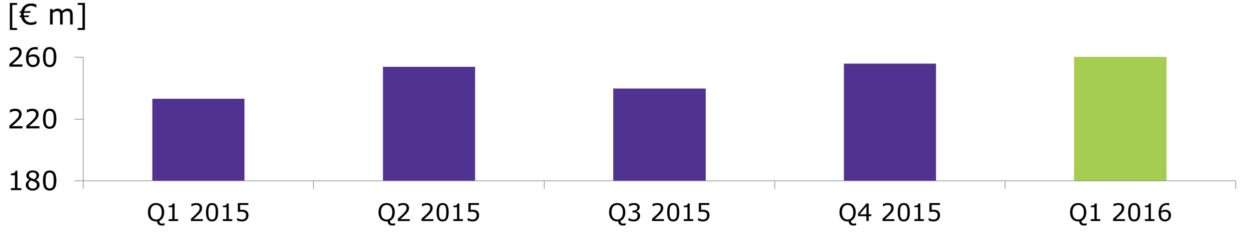
Erbitux performance

- Sales increase to €207 m due to solid volume development, but low base last year
- Europe impacted by mandatory price cuts & increasing competition offset by uptake in Russia
- Latin America shows strong growth esp. in Brazil after weak 2015
- APAC mainly reflects reduction of patient population due to label change in Japan (April 2015)

Strong organic growth in Fertility, General Medicine and Endocrinology

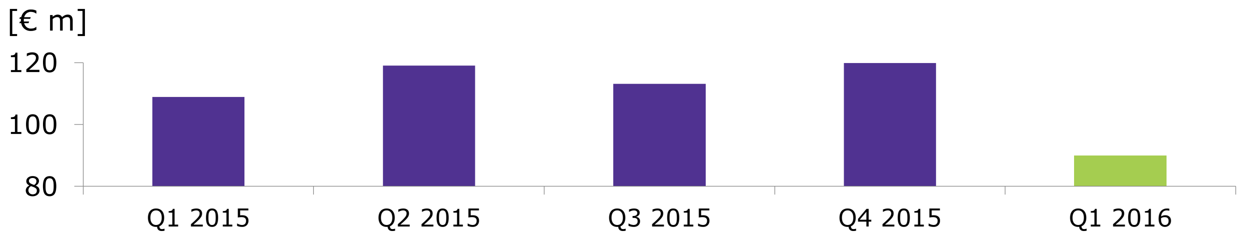
Sales evolution

Fertility



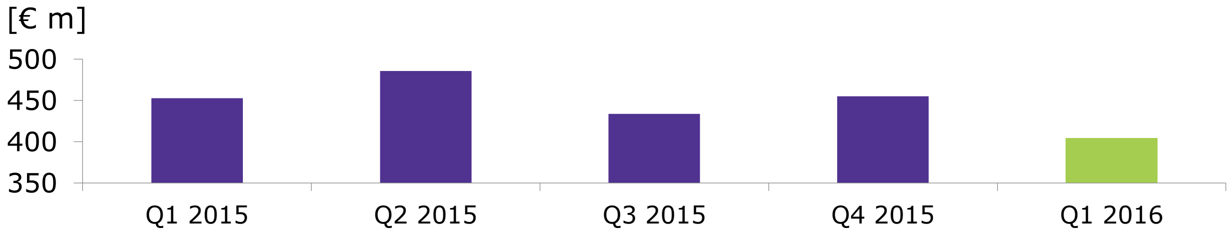
Organic

Endocrinology



Organic

General Medicine*



Organic

Q1 drivers

- Record quarter for Fertility (17% org.) driven by favorable competitive situation in U.S. and strong 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 driven by ongoing demand from China
- Glucophage benefits from successful repatriation in Russia; very strong comparables last year

*includes "CardioMetabolic Care & General Medicine and Others

Merck 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[1]
Anti-PD-L1 mAb
Solid tumors

M9241 (NHS-IL12)[2]
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[1]
Anti-PD-L1 mAb
Merkel cell carcinoma

Sprifermin
Fibroblast growth factor 18
Osteoarthritis

Atacicept
Anti-Blys/anti-APRIL fusion protein
Systemic lupus erythematosus

Phase III

Avelumab[1] – Anti-PD-L1 mAb
Non-small cell lung cancer 1L[3]

Avelumab[1] – Anti-PD-L1 mAb
Non-small cell lung cancer 2L[4]

Avelumab[1] – Anti-PD-L1 mAb
Gastric cancer 1L[3]

Avelumab[1] – Anti-PD-L1 mAb
Gastric cancer 3L[5]

Avelumab[1] – Anti-PD-L1 mAb
Bladder cancer 1L[3]

Avelumab[1] – Anti-PD-L1 mAb
Ovarian cancer platinum resistant/refractory

Avelumab[1] - Anti-PD-L1 mAb
Renal cell carcinoma 1L[3]

MSB11022
Proposed biosimilar of Adalimumab
Chronic plaque psoriasis

Registration

Cladribine Tablets[6] – Lymphocyte targeting agent
Relapsing-remitting multiple sclerosis

- Neurodegenerative Diseases
- Oncology
- Immunology
- Immuno-Oncology
- Biosimilars

Pipeline as of May 19, 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.

[1]Avelumab is the proposed International Non-proprietary Name (INN) for the anti-PD-L1 monoclonal antibody (previously known as MSB 0010718C);
[2]Sponsored by the National Cancer Institute (USA); [3] 1st line treatment; [4] 2nd line treatment; [5] 3rd line treatment
[6]As announced on September 11, 2015 Merck KGaA Darmstadt, Germany is preparing a regulatory submission to the European Medicines Agency

Exceptionals in Q1 2016

Exceptionals in EBIT

[€m]	Q1 2015		Q1 2016	
	Exceptionals	thereof D&A	Exceptionals	thereof D&A
Healthcare	12	0	-321	0
Life Science	20	0	109	0
Performance Materials	4	0	6	0
Corporate & Other	12	0	7	0
Total	48	0	-198	0

Totals may not add up due to rounding

Life Science: New reporting structure reflects customer-centric approach

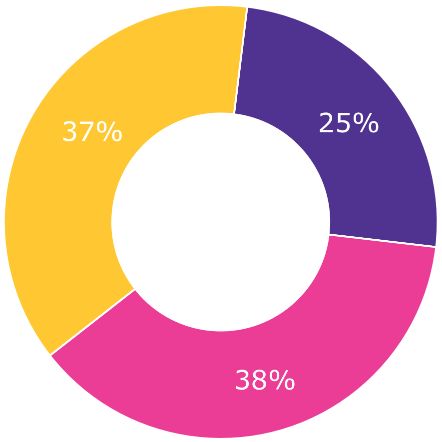
2015 product-centric structure

- Process Solutions
- Lab Solutions
- Bioscience

New 2016 customer-centric structure

- Process Solutions
- Applied Solutions
- Research Solutions

New sales breakdown as of Q1 2016

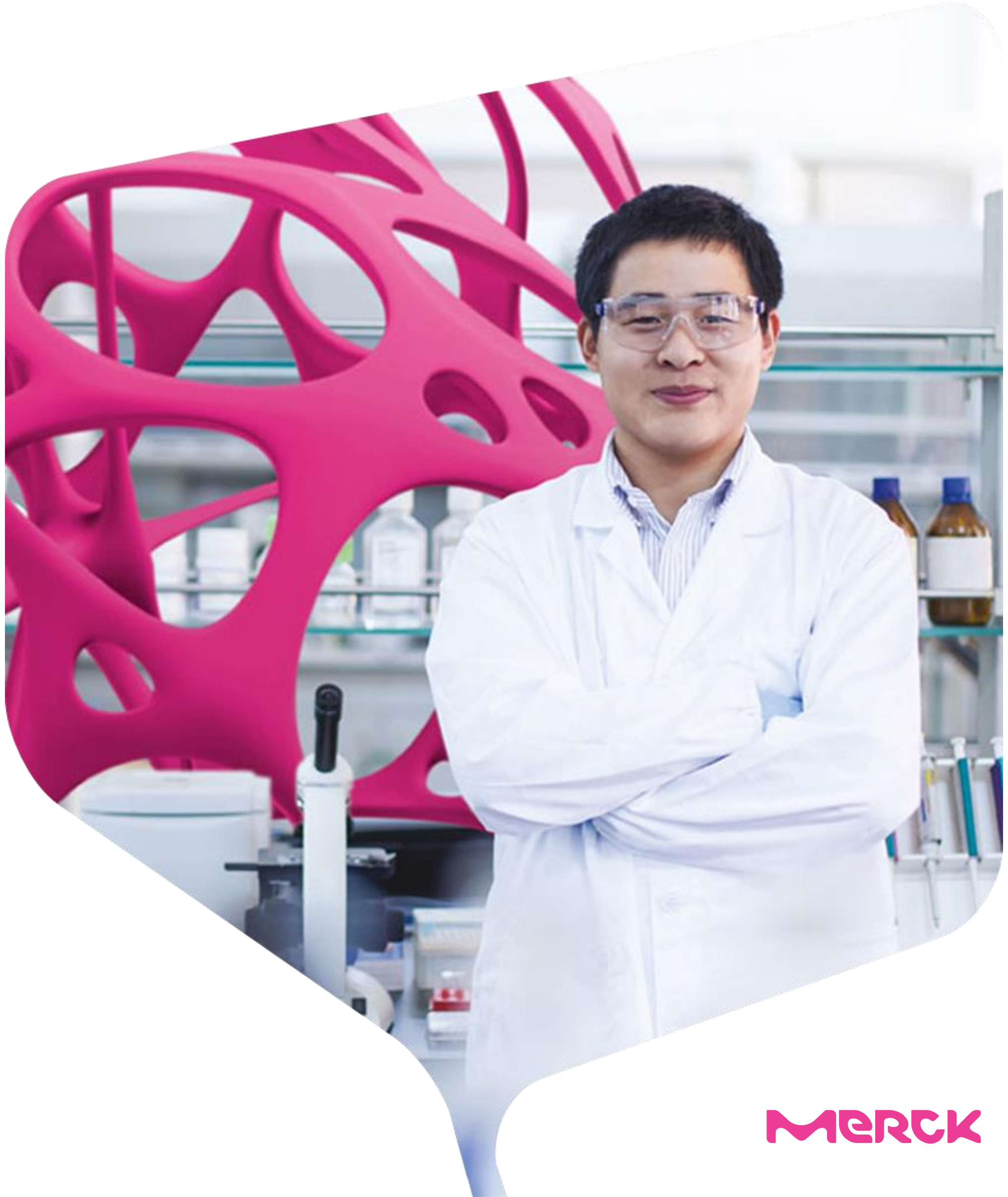


Customer-centric business approach:

- Enhanced emphasis on customers' unique needs
- Customized solutions for the specific needs of each segment from the start of product development to its completion

Financial calendar

Date	Event
June 20, 2016	R&D Update Call
August 4, 2016	Q2 2016 Earnings release
November 15, 2016	Q3 2016 Earnings release



CONSTANTIN FEST

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

SVENJA BUNDSCHUH

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

ALESSANDRA HEINZ

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

ANNETT WEBER

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

JULIA SCHWIENTEK

Institutional Investors / Analysts
+49 6151 72-7434
julia.schwientek@merckgroup.com

EVA STERZEL

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

OLLIVER LETTAU

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

EMAIL: investor.relations@merckgroup.com
WEB: www.investors.merck.de
FAX: +49 6151 72-913321