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Agenda

- Business overview
- **O2** Healthcare Funding for success
- Life Science Focusing on profitable growth
- Performance Materials Maintaining leadership and innovation
- **Executive summary and guidance**

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

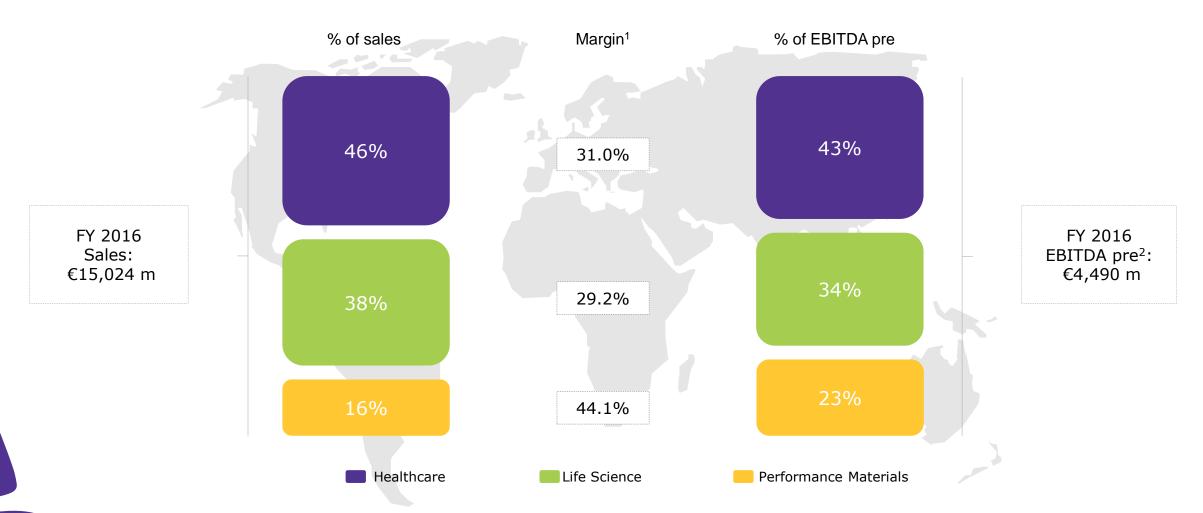
- Tools and services for biotech research & production
- Tools and laboratory supply for the academic research and industrial testing



Market leader in specialty materials

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

Strong businesses with attractive margins



We keep focus on our major priorities and goals per 2018



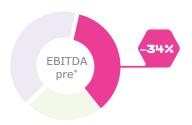
Healthcare



- Maximize growth of existing franchises
- Deliver pipeline: one product launch or indication p.a. from 2017



Life science



- 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 also beyond displays

Financial performance: solid organic growth amid pipeline investments

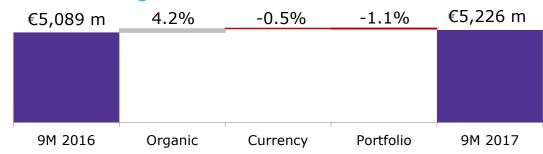
Healthcare P&L

[€m]	9M 2016	9M 2017
Net sales	5,089	5,226
Marketing and selling	-1,878	-2,033
Administration	-202	-226
Research and development	-1,078	-1,188
EBIT	1,314	1,375
EBITDA	1,947	1,847
EBITDA pre	1,631	1,566
Margin (in % of net sales)	32.0%	30.0%
EBITDA EBITDA pre	1,314 1,947 1,631	1,37 1,84 1,56

Comments

- Rebif declines due to competition in U.S. & EU
- Erbitux shows organic decline volume increase in growth markets outweighed by competition and price reductions in Europe
- General Medicine portfolio posts double-digit organic growth driven by strong performance in growth markets and repatriation in China
- M&S reflects investments for launches & Glucophage repatriation China
- R&D costs increase due to pipeline development
- Profitability benefits from royalty swap, Bavencio milestones and organic performance, but more than offset by higher R&D and M&S costs

Net sales bridge

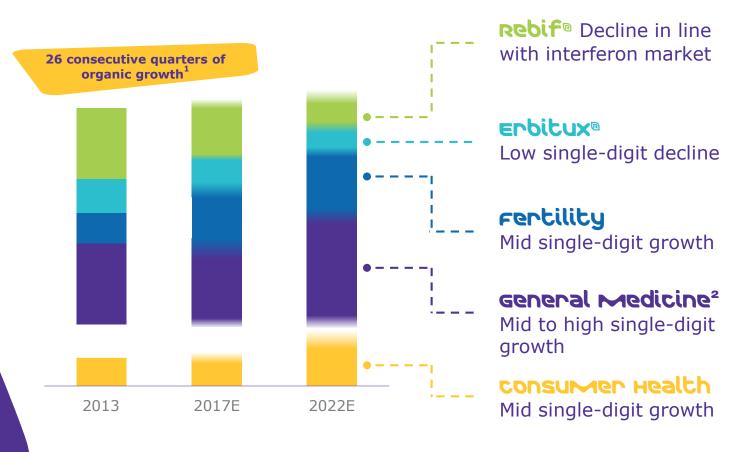


Key achievements 2017

- √ Mavenclad: approved in Europe (RMS)
- √ Bavencio: approved in two indications (MCC/UC)
- ✓ BTK inhibitor: comprehensive phase IIb program started
- ✓ Anti-PD-L1/TGF-ß trap: broad phase Ib program started
- ✓ Early stage: innovative assets in-licensed

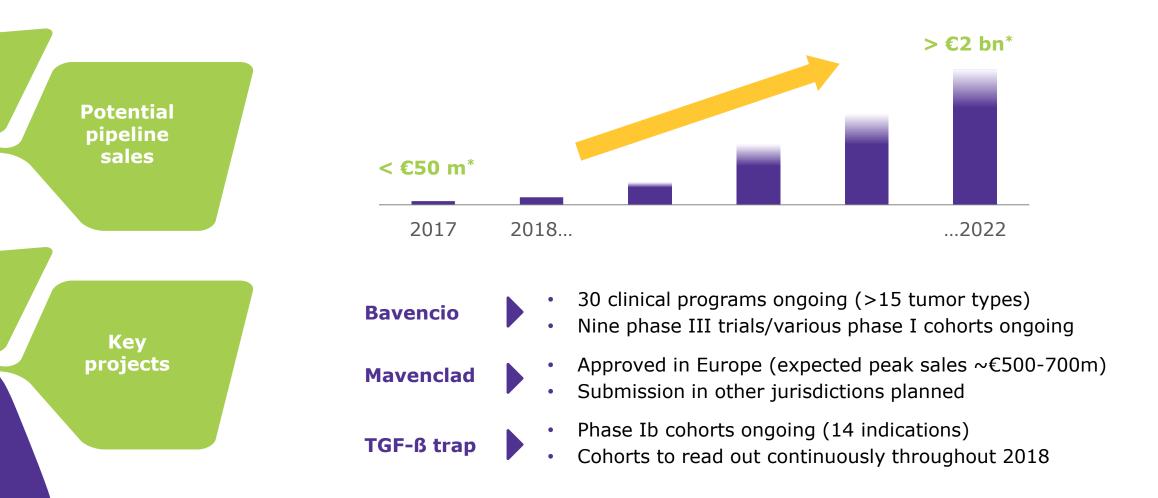
Base business: ambition to keep sales organically stable until 2022

Healthcare base business net sales until 2022



- Maintaining solid track record of patient retention
- Driving emerging markets growth
 - Drug demand driven by emerging markets growth and demographics
- Emerging markets demand growth enhanced by new launches, e.g. GlucophageXR® China
- Continuously build emerging markets presence/ invest in key strategic brands

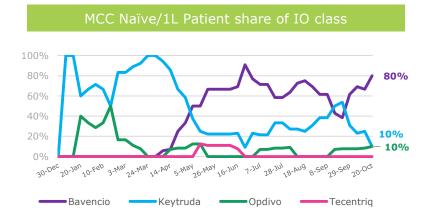
Development pipeline: on track to deliver the €2bn ambition by 2022

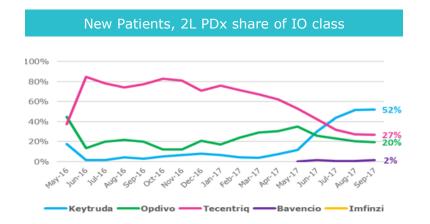


Bavencio: targeted launches picking-up continuously









on track to reach ~€20 m sales in 2017

 Successful uptake since accelerated approval³

 Targeted go-to-market strategy (conditional approval)

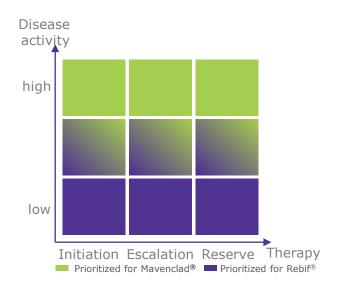
¹mMCC = metastatic Merkel cell carcinoma; ²mUC = metastatic urothelial cancer; ³Accelerated FDA approval for mMCC on March 23, 2017, and for mUC on May 9, 2017; Continued approval for these indications in the U.S. is contingent upon verification and description of clinical benefit in confirmatory trials

Data sources: IMS claims data

Mavenclad: launch ramping-up successfully

Position Complementary offering

Posology Unique value proposition





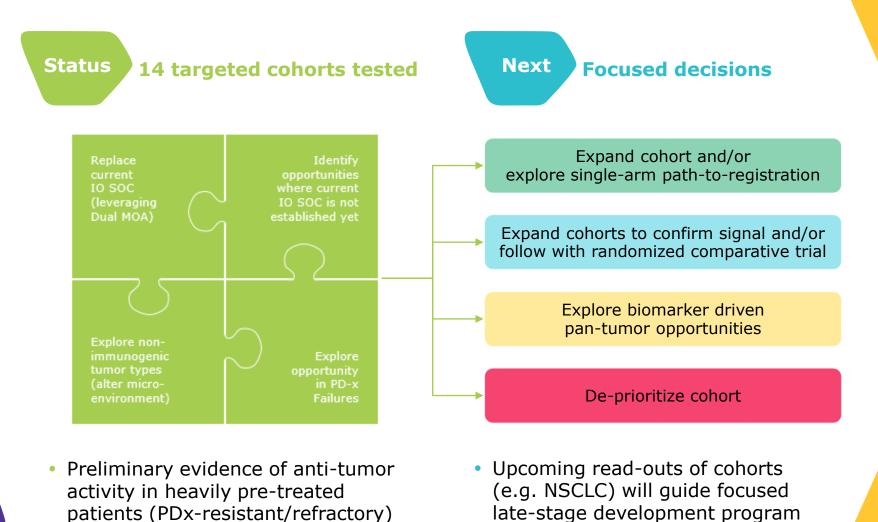
MAVENCLAD®

on track to reach high single digit €m sales in 2017

- Mavenclad label: ~60-70% of RRMS
- Rebif & Mavenclad complementary at patient and physician level

- Selective immune reconstitution.
- 2 years treatment for 4 years efficacy
- Positive NICE recommendation in UK

Anti-PD-L1/TGF-ß trap: reading-out throughout 2018



anti-pp-L1/ TGF-B trap on track to inform focused development

program for 2018

Growth funded through active portfolio management

Kuvan divestment

- Returning rights for Kuvan (rare disease) to BioMarin
- +324m€ divestment gain

Biosimilars divestment

- Selling Biosimilars activities
- +156m€ upfront, up to 500m€ milestone payments/royalties

Consumer Health

- Announced evaluation of strategic options for OTC business*
- Financials subject to deal model

rocused ræd on three specialty therapeutic areas



Nov '14 Jan '17 Jun '17 Sep '17 Sep '17

Pfizer alliance

- Co-promotion agreement to jointly develop/commercialize avelumab (immuno-oncology)
- +850m\$ upfront plus regulatory/ commercial milestone payments

Vertex in-licensing

- Access to two promising clinical programs targeting DNA damage and repair
- -230m\$ upfront plus royalties on future sales

F-star collaboration

- Develop and commercialize five bi-specific IO assets (incl. anti-LAG-3/PD-L1 asset)
- Up to -115m€ upfront, plus R&D funding/milestones

Innovative deal Models to realize full potential of assets

Long-term: pipeline optionality / short-term: five priorities (H1 2018)

Phase I Phase II Phase III Registration avelumab M2698 tepotinib avelumab - anti-PD-L1 mAb cladribine tablets p70S6K & Akt inhibitor anti-PD-L1 mAb c-Met kinase inhibitor Non-small cell lung cancer 1L1 lymphocyte targeting Solid tumors Solid tumors Non-small cell lung cancer avelumab - anti-PD-L1 mAb agent M3814 avelumab Non-small cell lung cancer 2L2 Relapsing multiple sclerosis6 tepotinib DNA-PK inhibitor anti-PD-L1 mAb c-Met kinase inhibitor avelumab - anti-PD-L1 mAb Solid tumors Hematological malignancies Hepatocellular cancer Gastric cancer 1L-M^{1M} M9831 (VX-984) M9241 (NHS-IL12)4 avelumab - anti-PD-L1 mAb avelumab - anti-PD-L1 mAb DNA-PK inhibitor Cancer immunotherapy Gastric cancer 3L3 Merkel cell cancer 1L1 Solid tumors Solid tumors avelumab - anti-PD-L1 mAb M6620 (VX-970) M7824 sprifermin Ovarian cancer platinum resistant/refractory ATR inhibitor anti-PD-L1/TGFbeta trap fibroblast growth factor 18 avelumab - anti-PD-L1 mAb Solid tumors Solid tumors Osteoarthritis Ovarian cancer 1L1 M4344 (VX-803) M4112 atacicept avelumab - anti-PD-L1 mAb ATR inhibitor Cancer immunotherapy anti-Blys/anti-APRIL fusion protein Urothelial cancer 1L-M1M Solid tumors Solid tumors Systemic lupus erythematosus Oncology avelumab - anti-PD-L1 mAb M3541 atacicept Renal cell cancer 1L1 Immuno-Oncology M1095 (ALX-0761)5 ATM inhibitor anti-Blys/anti-APRIL fusion protein avelumab - anti-PD-L1 mAb anti-IL-17 A/F nanobody Solid tumors IgA nephropathy Immunology Locally advanced head and neck cancer Psoriasis M8891 abituzumab – anti-CD51 mAb Neurology M6495 MetAP2 inhibitor Systemic sclerosis with interstitial lung disease, anti-ADAMTS-5 nanobody Solid tumors General Medicine evobrutinib Osteoarthritis M7583 BTK inhibitor BTK inhibitor Rheumatoid arthritis M5717 Hematological malignancies PeEF2 inhibitor evobrutinib Pipeline as of November 1st, 2017 Malaria BTK inhibitor Pipeline products are under clinical investigation and have not been proven Systemic lupus erythematosus There is no quarantee any product will be approved in the sought-after indication. evobrutinib BTK inhibitor Multiple sclerosis

Prioritization/ Externalization

Read-out of 14 indications

Two phase III decisions

Two phase III read-outs

One potential US submission

3



5

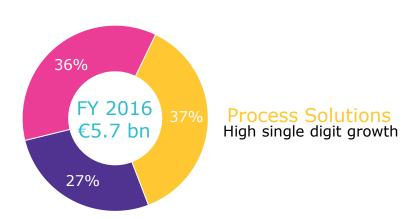
Life Science

Business on track to deliver above market organic growth (~4%)

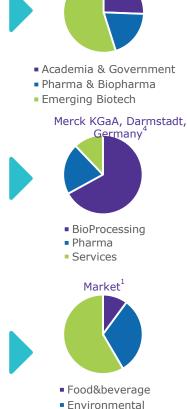
Market¹

Life Science

Research Solutions Low single digit growth







Long-term growth drivers

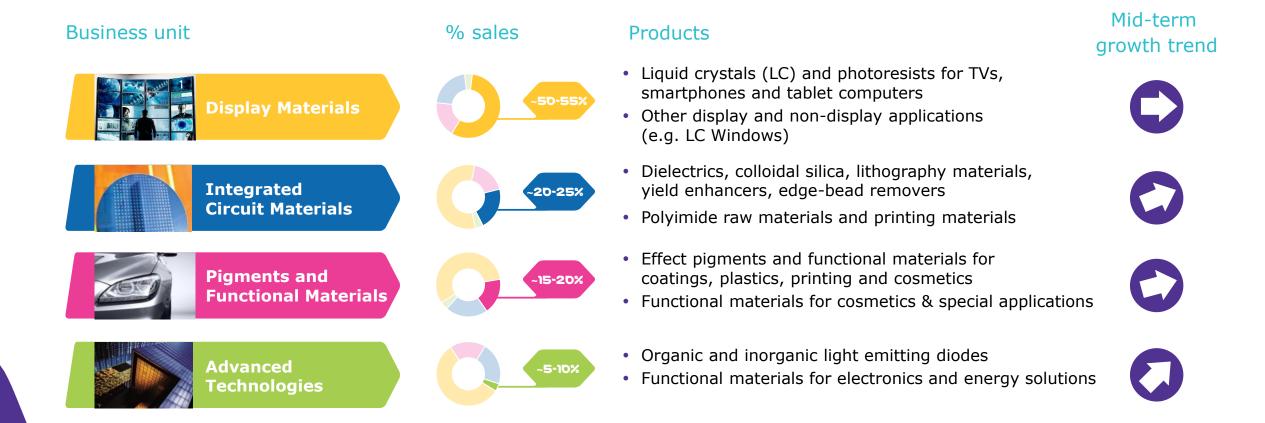
- Research activity: >3,000 projects in research pipelines², rising number of experiments and newly emerging therapies/technologies backs healthy growth in biotech and CROs³
- Public and private funding: availability, access and predictability drive demand from academia and emerging biotech customers
- Regulation: rising requirements foster long-term customer partnerships
- Biologics: mAbs production⁵ growing by ~12% p.a. for 2016-2021 driven by new molecules and biosimilars
- Diversification: contribution by top 10 molecules will decline to ~30% until 2021 from 80% today⁶
- Noval modalities: innovation in complex-to-deliver therapies, e.g. gene and cell therapy, will drive demand for single-use, end-to-end and new technology solutions
- **Regulation**: testing volumes overall are rising globally rise in quality standards and increased demand for testing across customer segments
- Population and economic growth: demand for access to more sophisticated products and services rises, e.g. in emerging markets
- Speed: need for fast testing results raises requirements for Applied customers, esp. in clinical testing and food & beverage testing

Diagnostics

¹Source: Merck KGaA, Darmstadt, Germany Factbook; ²Source: PhRMA; ³CRO = commercial research organisation; ⁴Indicative only; ⁵mAbs = monoclonal antibodies; ⁶Source: EvaluatePharma April 2017

Performance Materials

The four pillars are set for future profitable growth





Well-founded medium-term low single-digit growth profile

We are well on track to deliver on our promises



Group

Net debt reduced by >€2 bn¹ Strict financial discipline supports rating



Healthcare

Base business growing Launches and pipeline delivery materializing



Life Science

Sigma-Aldrich synergies raised and well on track Organic growth above market



Performance Materials

Market challenges well managed New technologies in test phase

Important milestones reached to deliver on our promises







Full-year 2017 guidance

Net sales: ~ €15.3 - 15.7 bn

EBITDA pre: ~ €4,400 - 4,600 m

EPS pre: ~ €6.15 – 6.50



