Merck KGaA, Darmstadt, Germany

Berenberg Pennyhill Conference 2012

Eva Schäfer-Jansen

Merck KGaA, Darmstadt, Germany Investor Relations

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Merck KGaA

Darmstadt · Germany



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A healthy portfolio





Pharmaceuticals

Development, manufacturing and marketing of prescription and over-the-counter drugs



Life Science Tools

Products and solutions to support life science research, pharmaceutical development and drug production



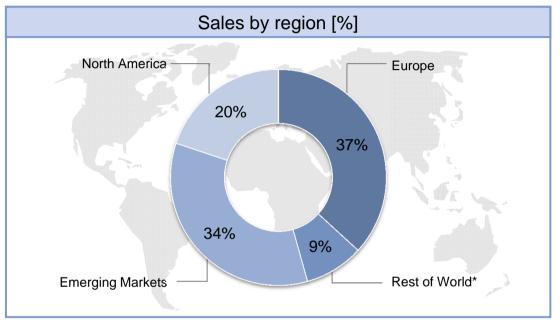


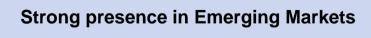
Speciality Chemicals

Number one supplier of liquid crystal mixtures used in LCD panels



Geographically diverse business





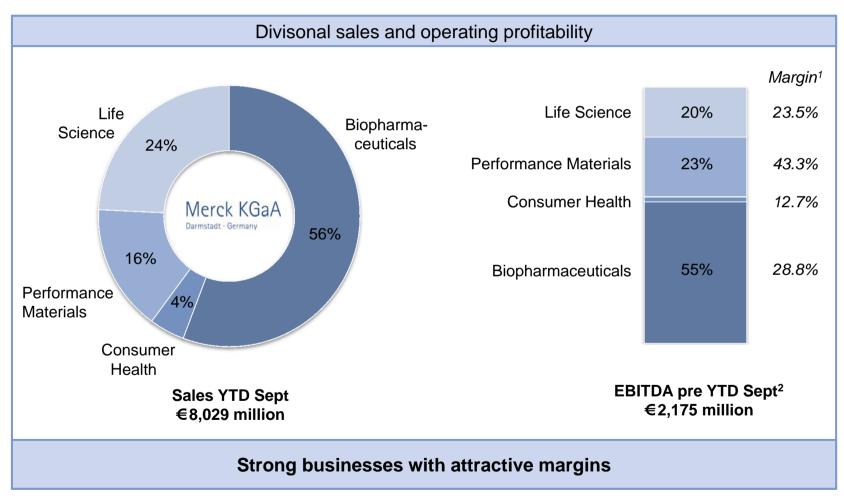


^{*} Africa, Australia, Japan





A balanced portfolio of four divisions



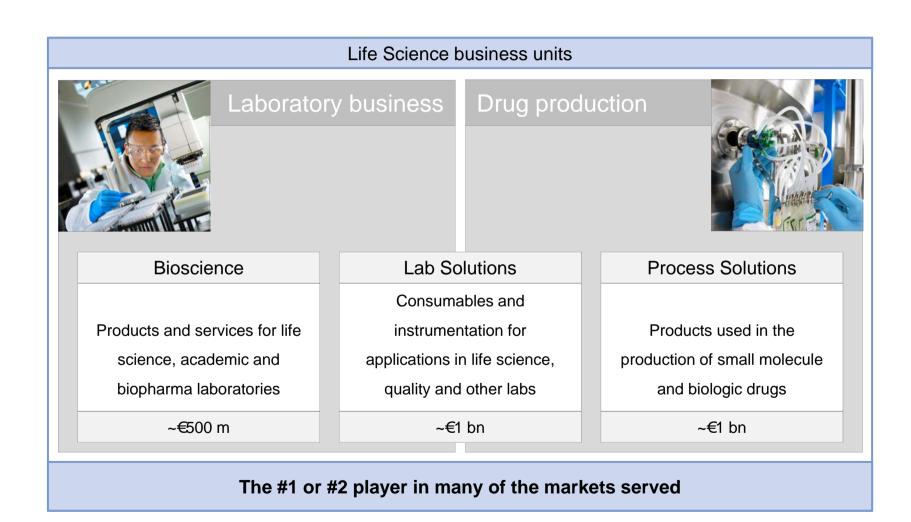
¹ EBITDA pre one-time items as % of sales

² Including Corporote and Others (- € 161 million)





A leader in life science tools products





Highly specialized chemical business dominated by display materials



Source: DisplaySearch, Quarterly Worldwide Flat Panel Forecast Report Q3 2012





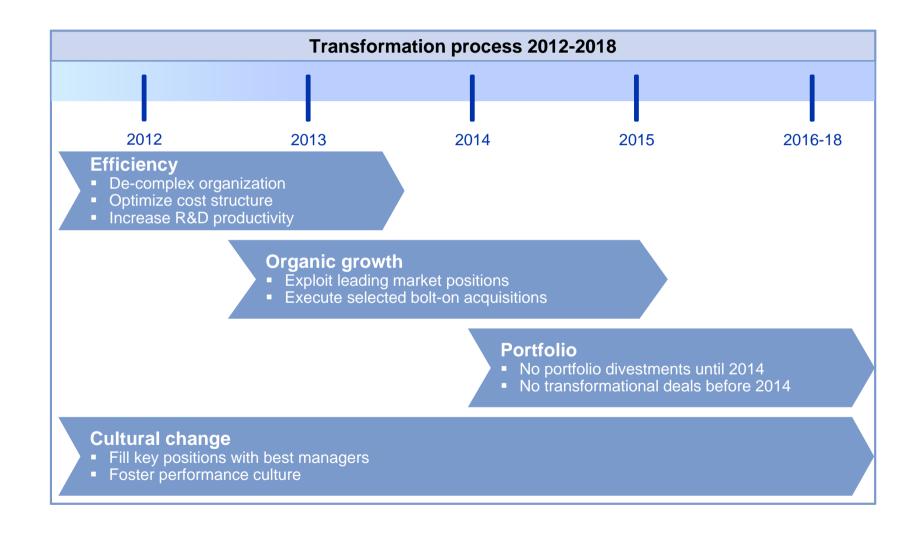
Improving its operating model and competitiveness







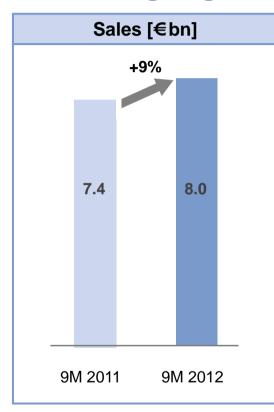
Focus on efficiency

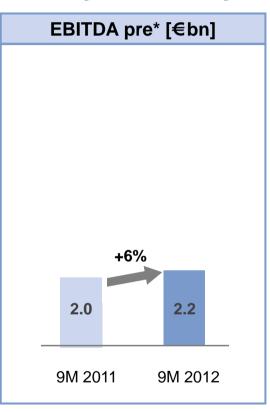


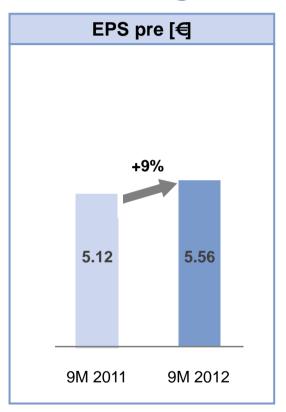


The Group

We are positioned to report solid numbers in challenging economy and a year of change







Solid top-line and profitability improvement in first year of transformation

^{*} EBITDA pre one-time items

Biopharmaceuticals

Berenberg Pennyhill Conference 2012

Annalisa Jenkins

Global Head of Drug Development & Medical

London, December 7, 2012





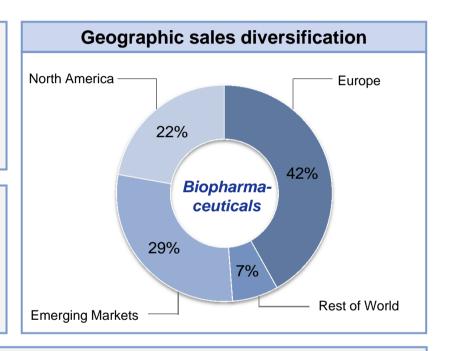
Our business today



Lack of patent cliff provides resilient revenue stream



Portfolio of strong brands continue to generate growth

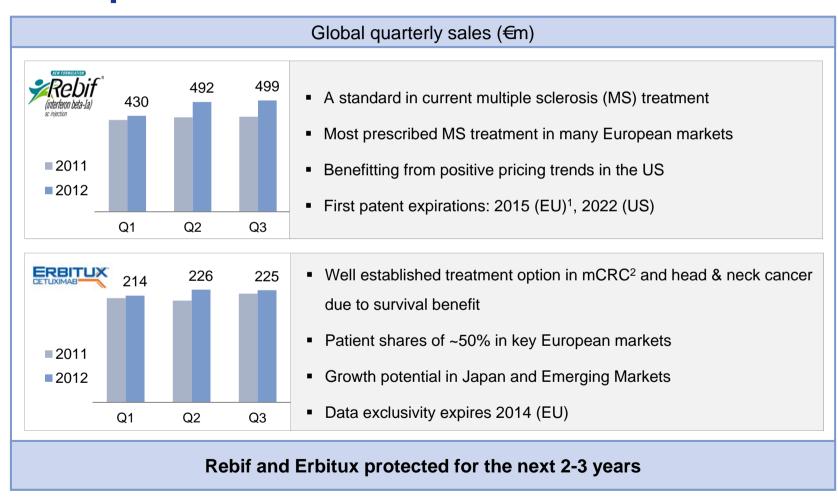




Rebuilding the pipeline to provide for future growth



Key products Rebif and Erbitux continue to be important assets



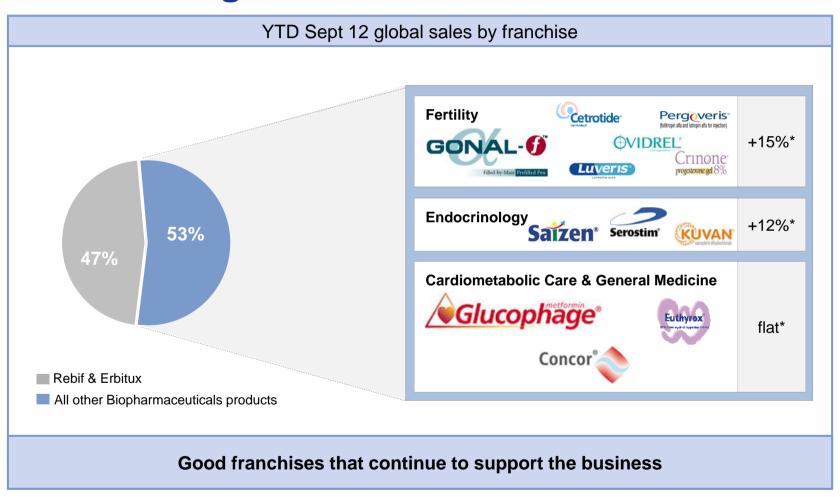
¹ First formulation, next generation (HSA-free) formulation is marketed in Europe

² KRAS wild-type, ~60% of the total patient population



Biopharmaceuticals

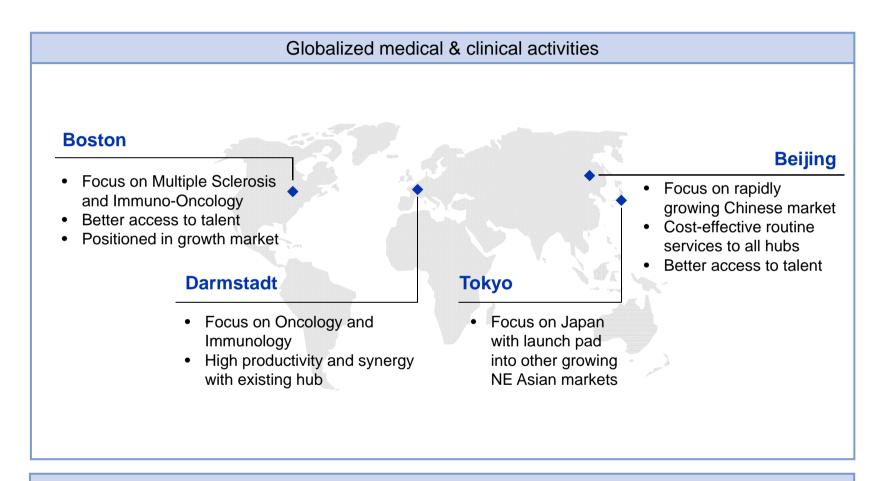
Despite patent expiry many of our products continue to grow



^{*} Organic sales growth YTD Sept 12



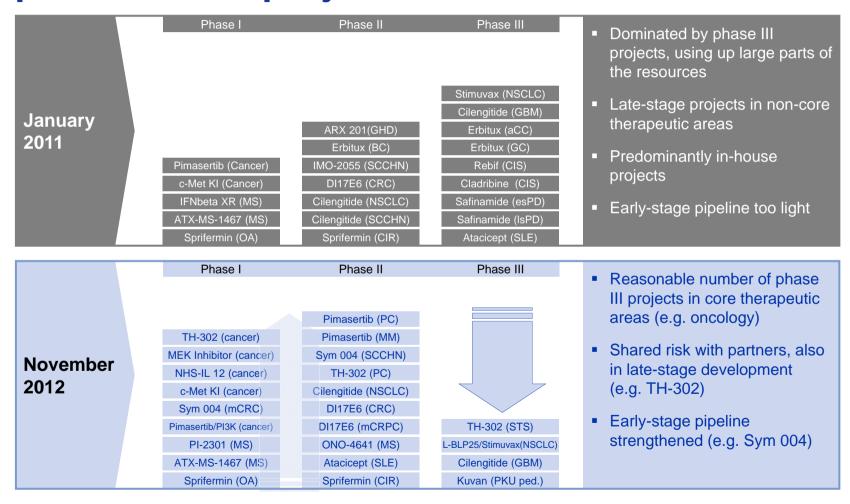
Global R&D Centers



Future growth opportunities in Western and Emerging Markets

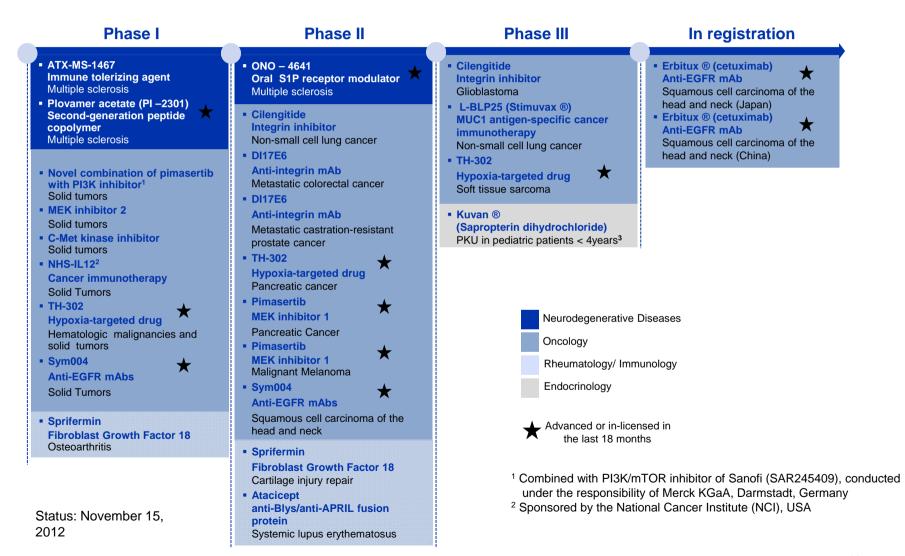


Rebuilding our pipeline with an emphasis on phase I and II projects





A pipeline with a strong focus on oncology





Examples of access to external capabilities

Research



- Platform to discover a novel class of antibody derived therapeutic proteins
- Collaborations in the therapeutic areas of immunology, oncology and osteoarthritis
- Currently 2 projects in pre-clinical testing

Development



- In-licensing of SYM 004, a mixture of two EGFRantibodies
- Currently in Phase I/II
- Potential to build on existing Erbitux franchise

Complementing our in-house capabilities in protein and antibody engineering

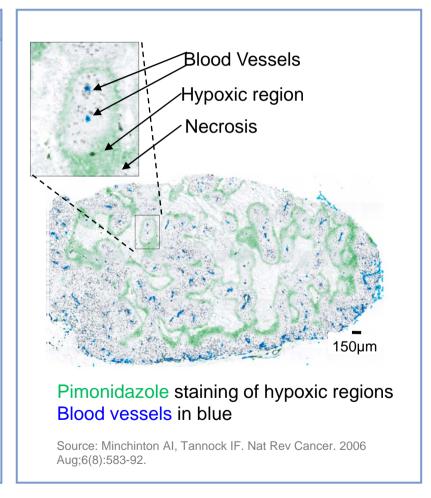
Biopharmaceuticals



TH-302: Hypoxia is a feature of tumour microenvironment

Scientific background

- Hypoxia is a feature of all solid tumours and some hematologic malignancies
- Greater tumour hypoxia is associated with poor prognosis¹
 - promotes mutagenesis, invasiveness, metastasis and drug resistance
- Tumour hypoxia underlies (chemotherapy, radiotherapy) treatment failure and relapse
- TH-302 cytotoxicity is activated by hypoxia
 - targets the regions where chemotherapy and radiotherapy fail



¹ Reviewed in Vaupel, P. & Mayer, A. Cancer Metastases Rev 26 225, 2007

Biopharmaceuticals



TH-302: pivotal trials in soft tissue sarcoma and pancreatic cancer

Mode of Action:

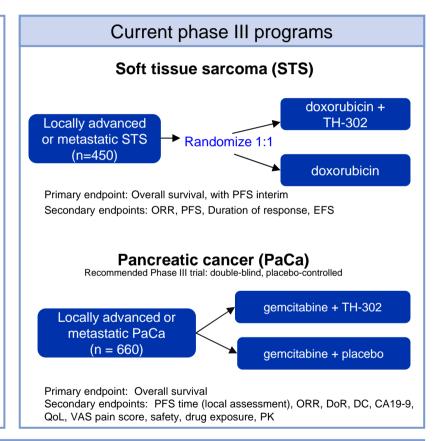
- DNA cross-linking effect that is targeted to areas of hypoxia
- TH-302 is the prodrug of an alkylating mustard that is activated in areas of hypoxia:

Soft tissue sarcoma

- Orphan drug designation granted by FDA
- A Special Protocol Assessment exists with FDA

Pancreatic cancer

 A Special Protocol Assessment recently agreed to by FDA

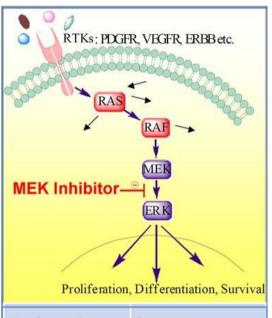


Phase I/II trials currently underway in other solid tumors as well as hematological malignancies

Special protocol assessment: written agreement with the US Food and Drug Agency that the design and planned analysis of a study can adequately address objectives in support of a regulatory submission. However, FDA's determinations for marketing application approval are made after a complete review of a marketing application and are based on the entire data in the application



Pimasertib: MAPK signaling is dysregulated in many human solid and hematological tumors



| Key Competitors | Company |
|----------------------------|-----------------------------|
| Trametinib (GSK1120212) | GSK |
| MEK162 (ARRY 162) | Novartis Array Biopharma |
| BAY86-9766 (RDEA119) | Bayer Ardea Biosciences |
| Selumetinib (AZD6244) | AstraZeneca |

Scientific background

- Mitogen-activated protein kinase (MAPK) cascades are key signaling pathways involved in the regulation of cell proliferation, survival and differentiation
- The crucial role of the RAF-MEK-Erk MAPK cascade in tumorigenesis has been clearly established
- Generally, MAPK pathway activation occurs in tumors by receptor tyrosine kinase (RTK) stimulation or by activating mutations in regulators of the pathway
- Evidence of anti-tumor activity following MEK inhibition has been demonstrated particularly in tumors where constitutive activation of the MAPK pathway is significantly present

Competitive field with Pimasertib being one of the most advanced compounds





Pimasertib: selective and potent MEK inhibitor

Development strategy

Mode of Action

 Highly selective, ATP non-competitive, oral, small molecule inhibitor of MEK 1 and 2

Combinations

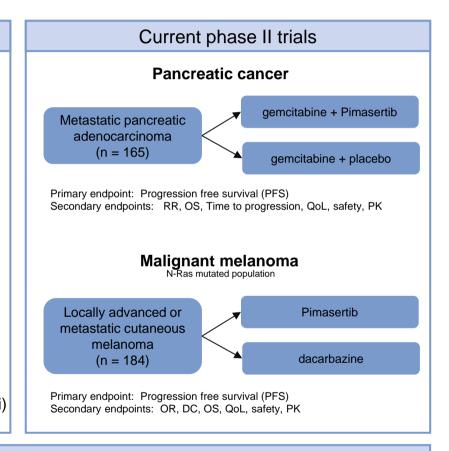
 As a preferred MEKI partner with current and future standards of care in selected indications

Monotherapy

In selected indications

Novel-novel combinations

- To provide additional benefit to patients
- To expand the combinability profile of pimasertib
- To leverage also 3rd party collaborations (e.g. Sanofi)



Employing a biomarker-driven development





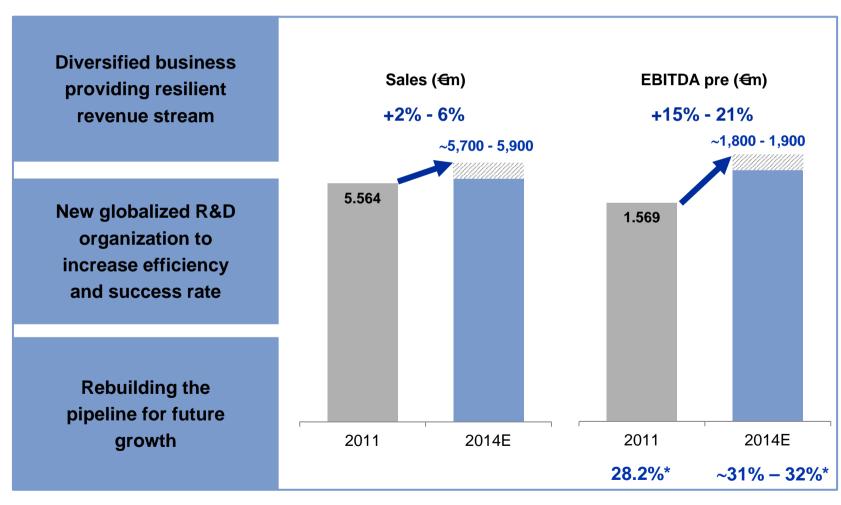
Upcoming pipeline newsflow

| Project | Indication | Current phase | Timing | Event |
|-----------------------|------------------------------|-----------------|---------|------------------------------------|
| Erbitux | SCCHN | In registration | | Approval in Japan |
| L-BLP25 (Stimuvax) | Stage III NSCLC | Phase III | Q1 2013 | Head-line result (START) |
| Cilengitide | Glioblastoma | Phase III | H1 2013 | Head-line result (CENTRIC) |
| TH-302 | Soft tissue sarcoma | Phase III | H1 2013 | Futility analysis |
| ONO-4641 | Multiple sclerosis | Phase II | 2013 | Phase III 'go/no go' decision |
| Atacicept | Systemic lupus erythematosus | Phase II | H1 2013 | Publication of results (APRIL SLE) |





Conclusion and mid-term targets



^{* %} of sales, assumes no major new product launches until end 2014

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