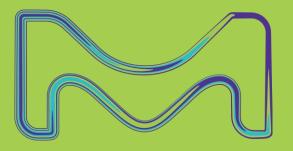
Merck KGaA, Darmstadt, Germany 21 Roadshow

Marcus Kuhnert, CFO Matthias Heinzel, CEO Life Science

August 2021



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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 as well as the impact of future regulatory or legislative actions.

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Agenda

- **Business overview**
- **O2** Transforming the company
- Healthcare Executing on the earnings phase
- Life Science Focusing on profitable growth
- **Electronics Leveraging portfolio shift**
- **Sustainability**
- **Guidance & executive summary**



Business overview

Ol

Three high-tech businesses competing in attractive markets



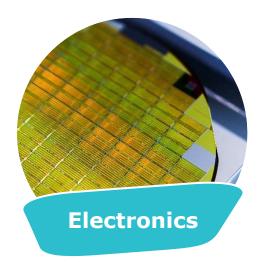
Leading in specialty pharma markets

- Biologics and small-molecule prescription medicines against cancer, multiple sclerosis, infertility
- Research focus: Oncology, Immunology & Immuno-Oncology
- Successful portfolio management: e.g. divestment of Consumer Health and Allergopharma



Leading life science company

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



Leading company in high-tech solutions

- High-tech solutions and materials for electronics
- Broad portfolio of decorative and functional solutions



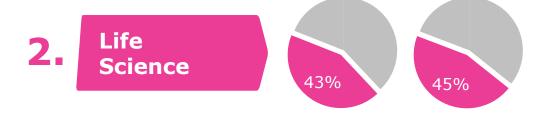
Group today – three strong pillars as basis for profitable growth





Global specialty innovator poised for above-industry growth

- Resilient base business backed by excellent life cycle management
- Strong growth from new products, late-stage pipeline assets with blockbuster potential
- Rigorous cost discipline and value-maximizing pipeline prioritization



Diversified industry leader poised for above-market growth

- Portfolio advantage and outperformance drive above-market growth
- Strengthen core: products (PS), chemistry (RS), lab water (AS)
- Establish new pillars: PS services, gene editing and novel modalities



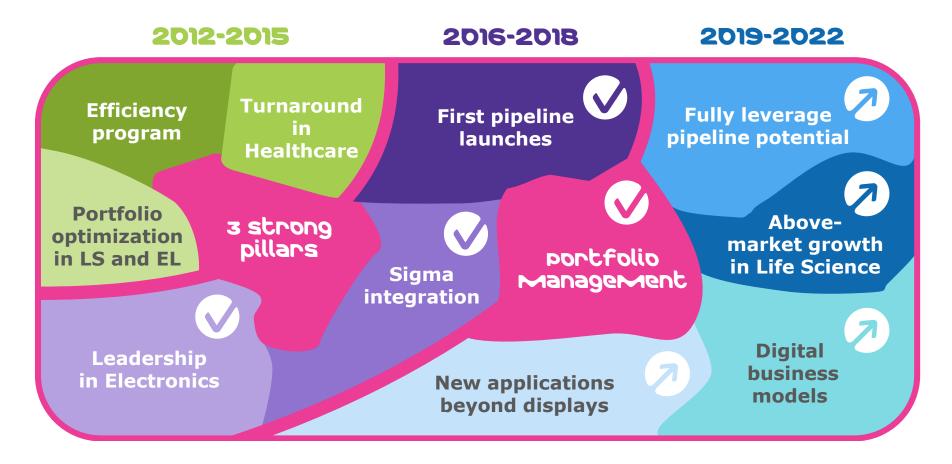
Leading electronics player poised for accelerating growth

- Growing semiconductor share as key driver for acceleration
- More resilient growth through rising diversification
- Strict cost discipline in maturing parts of the portfolio

¹EBITDA pre share excluding Corporate & Others; 2020 EBITDA pre restated for €365 m patent litigation provision release Acronyms: PS = Process Solutions, RS = Research Solutions, AS = Applied Solutions



The 2016 vision – a strategic agenda until 2022





Executing on the growth and expansion phase of the 2016-22 strategic agenda





2021 and beyond - poised for growth in a challenging environment



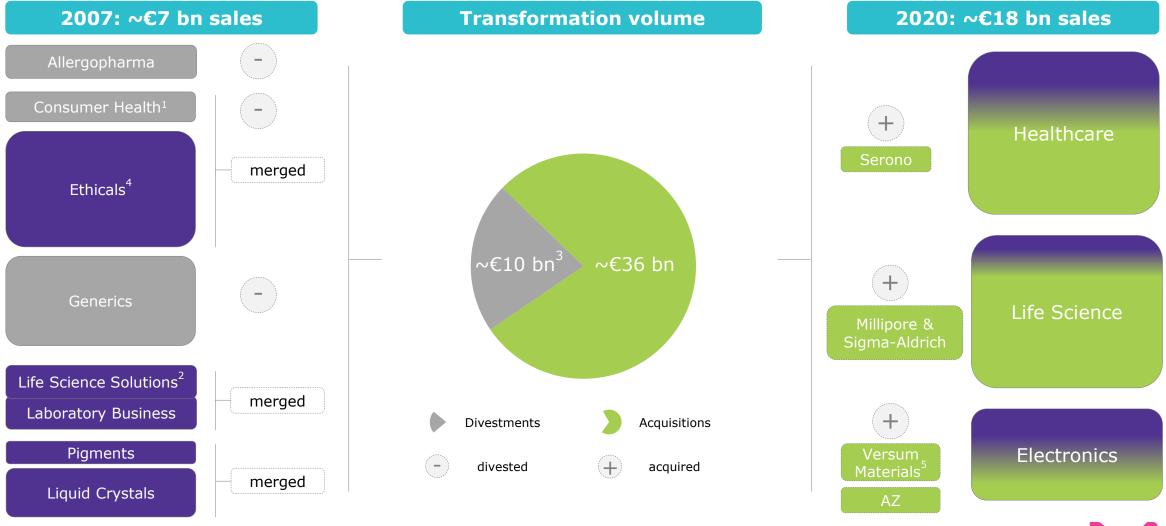
Acronym: VBP = volume based procurement



the company

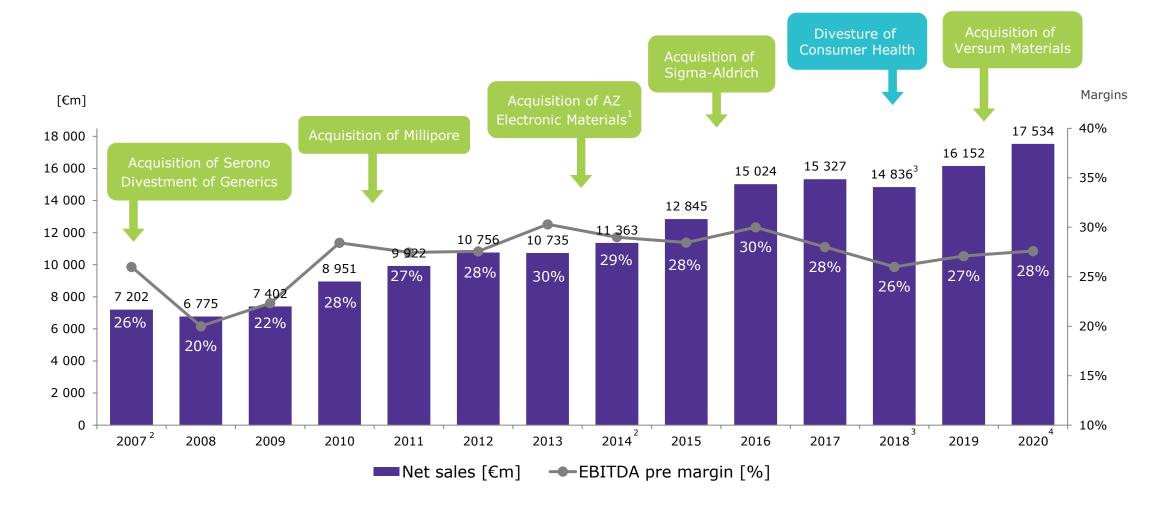
02

We have added scale and strengthened the attractiveness of our portfolio





Continue to transform to a science and technology focused company

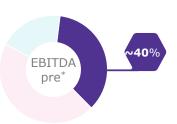




Clear set of priority goals



Healthcare



- Deliver on ambition to keep core business at least stable until 2022
- Transitioned from investment to earnings phase in 2019
- Foster successful Bavencio[®]
 and Mavenclad[®] ramp up
- Stringent pipeline execution

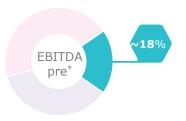


Life science



- Strengthen position as differentiated player in a highly attractive market
- Maintain consistent abovemarket growth trajectory and superior profitability





- Deliver ambition of 3-4% CAGR
- Implement 5-year transformation program & focus on continuous seamless integration of Versum
- Maintain strong cash generation and cash conversion



Three-pillar structure – positioned to win in high-growth markets

Global economy¹



Global **GDP**

~3% to 4%

End markets¹



Global pharma industry ~4% to 5%



Global life science industry



Global electronics industry ~4%



∑ ~4% to 5%

Focus market areas¹



Oncology: ~10%

Immunology: ~5% to 9%



Biologics: ~10% to 12%

Services: ~7% to 8%



Semi materials: ~4% to 6%

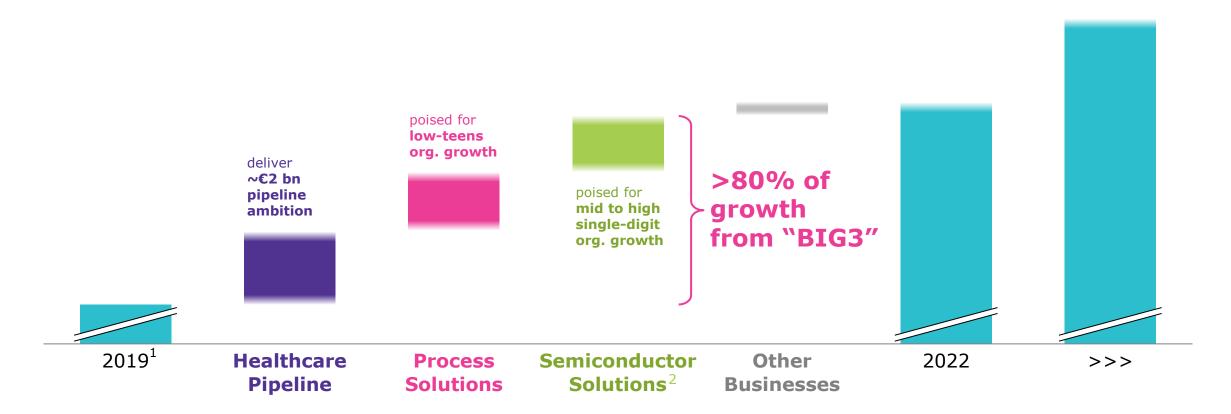




Purposefully positioned in attractive markets with secular growth above global GDP ...further focusing investments on attractive sub-segments

¹ Company estimates of mid-to-long-term growth outlook based on industry forecasts and reports from public research institutes (e.g. IMF, IQVIA, EvaluatePharma, Prismark, etc.)

Three main drivers of growth to 2022 and beyond



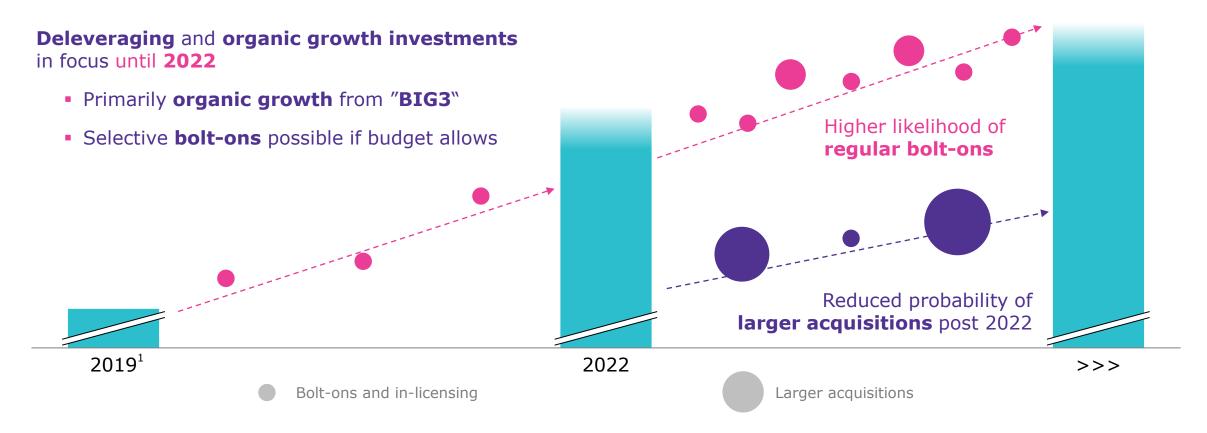


Beyond 2022: further significant growth potential from "BIG3" and increasing contributions from other businesses

¹ 2019 Group sales of €16.2 bn; ² Including Versum portfolio effect



Portfolio strategy - from transformation to evolution





Strong portfolio: significant organic growth potential to 2022 and beyond ...and higher likelihood of regular bolt-ons post 2022

¹ 2019 Group sales of €16.2 bn

Healthcare

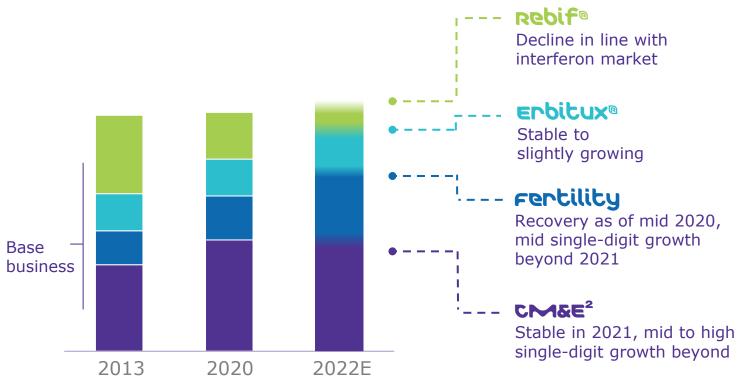
Executing on the earnings phase

03

Healthcare: Base Business

Ambition to keep base business ~stable throughout 2021 and 2022

Healthcare base business net sales until 2022



- Maintain solid track record of patient retention
- Integrate into joint franchise with Mavenclad®
- Drive EM¹ growth and mitigate competitive / price pressure in EU by clear branding
- Continued China NRDL inclusion secured through successful renegotiation in late 2020
- Drug demand driven by emerging markets growth and demographics
- Leverage competitive strengths (e.g. broad and innovative portfolio, security of supply)
- Drive recovery after COVID-19
- Increasing prevalence of diabetes and cardiovascular diseases
- Mitigate VBP pressure in China through EM growth, effective life cycle management, and portfolio expansion



¹ EM: emerging markets; ² Cardiovascular, Metabolism and Endocrinology (new Franchise name as of Q1 2021)

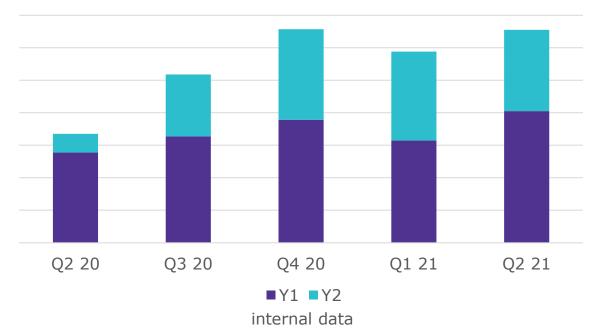
Healthcare: Mavenclad® US

Q2 Y1 patients growing QoQ with expanded prescriber base



New patient volume growing 23% QoQ, now comprising over 60% of total patient volume





- New patients growing QoQ both in terms of volume and % of total patients
- New prescribers growing QoQ by 23%
- **Total patient growth in H2 2021** to be driven by:
 - Higher volume of H2 2020 return patients
 - Continued new patient growth



Healthcare: Mavenclad®

Independent real-world data (RWD) differentiates Mavenclad®

- A high-efficacy DMT that demonstrates full antibody response to COVID-19 vaccination
- Differentiated vs. other high-efficacy therapies in light of COVID-19 vaccinations for MS patients

Patient population		Total N=125	Protective humoral immunity ^a
DMT treated patients	Mavenclad®	23	100% (p = 0.99) ^b
	Ocrelizumab	44	22.7% (<i>p</i> < 0.0001) ^b
	Fingolimod	26	3.8% (<i>p</i> < 0.0001) ^b
Untreated MS patients		32	100%
Healthy subjects		47	97.9%



In the first-ever real-world data study of its type all patients on Mavenclad® who received a mRNA COVID-19 vaccine were able to mount a full antibody response, similar to healthy subjects and untreated people with MS, irrespective of lymphocyte counts¹

DMT = disease-modifying therapy

^aProtective humoral immunity defined as a index value higher than 1.1 using EUROIMMUN semiquantitiative ELISA for IgG specific for the recombinant S1 subunit of SARS-CoV-2 spike protein bFisher's exact test to detect differences in categorical variables between DMT-treated patients with MS and untreated patients with MS

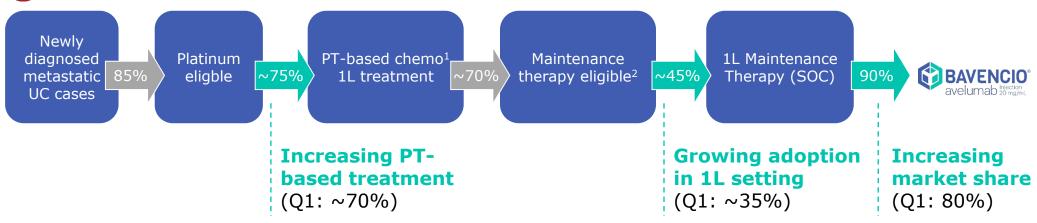


^{1.}Achiron et al. Ther Adv Neurol Disord https://doi.org/10.1177/17562864211012835

Bavencio® UC 1L launch: Increasing the adoption of 1L maintenance therapy in both U.S. and ex-U.S.



U.S. – 1 year into launch, continued progress across the entire treatment flow:







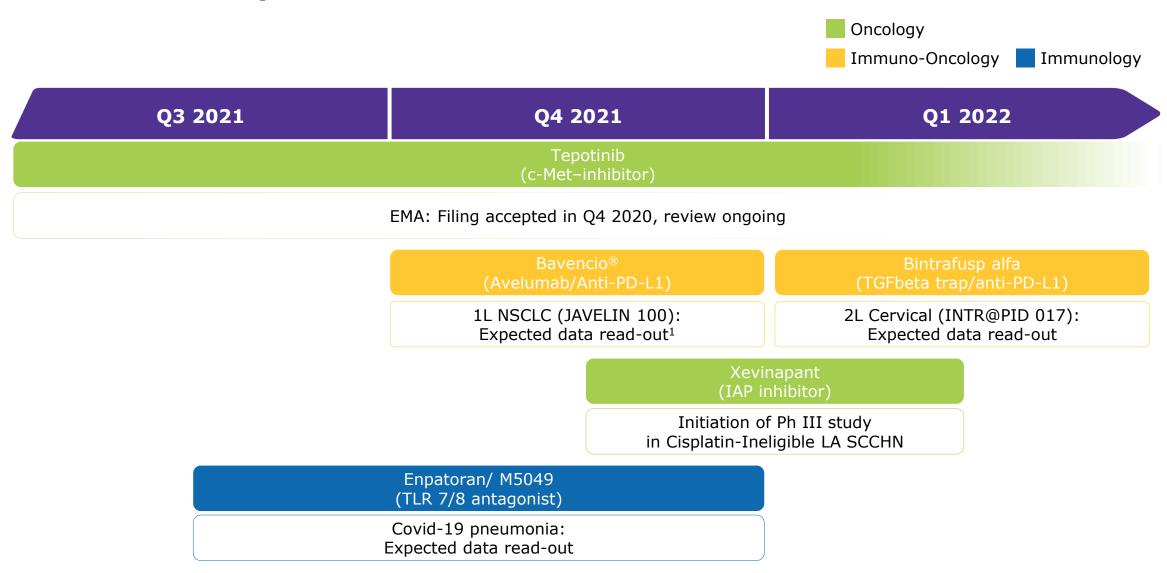
Europe & Japan – Recently approved, encouraging uptake:

- Now approved in 45 markets and reimbursed in ~1/3
- Strong initial uptake in key launch markets (e.g. Japan, France, Germany) with guideline recommendations and KOL support, on track to become SOC



^{1:} Carboplatin or Cisplatin, 2: Complete / partial response or stable disease based on clinical trial data; Acronyms: PT = Platinum, SOC = Standard of care

Healthcare catalysts



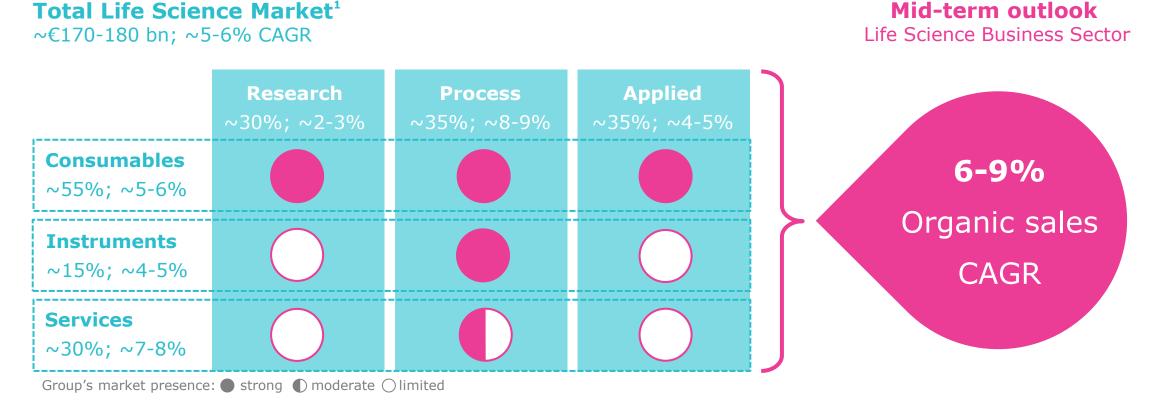
Acronyms: EMA = European Medicines Agency, LA = locally advanced, SCCHN = Squamous cell carcinoma of the head and neck, NSCLC = Non-small cell lung cancer, TLR = Toll-like receptor, 1: Clinical timelines are event-driven and may be subject to change



Focusing on profitable growth



Building growth momentum with focus on attractive market segments





Upgrading mid-term financial ambition to 6-9% organic sales CAGR

¹ Company estimate of the market segments, based on industry forecast over 5-year horizon; all growth rates in 3x3 Matrix indicate external market growth

Improved mid-term outlook driven by market and portfolio focus

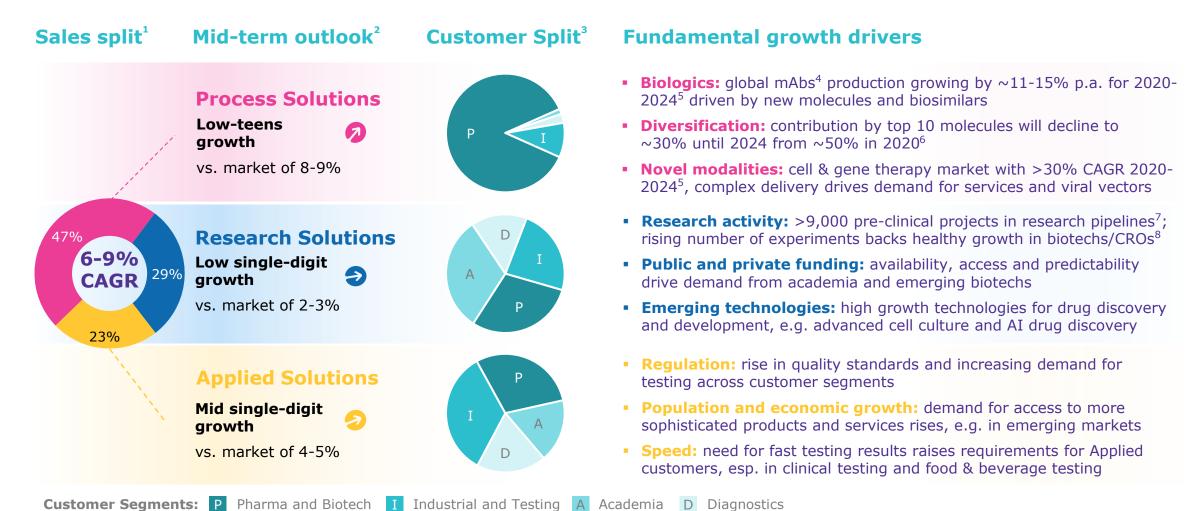






¹ Company estimate based on industry forecast over 5-year horizon; pre COVID-19 impact

All business units contributing to above-market growth



¹ Based on H1 2020, CAGR is organic mid-term ambition; ² growth rates are organic CAGRs pre COVID-19; ³ indicative only; ⁴ mAbs = monoclonal antibodies; ⁵ Source: company estimate based on industry forecasts; ⁶ Source: EvaluatePharma; ⁷ Source: statista; ⁸ CRO = Contract Research Organization

25

Critical offering in the fight against COVID-19



products feed into...

www.sigmaaldrich.com/covid-19

VIRUS **DETECTION**

- Leading critical component provider for Molecular and Serological diagnostic kits
- PCR reagents, kits and tools for all stages of assay development

VIRUS CHARACTERIZATION

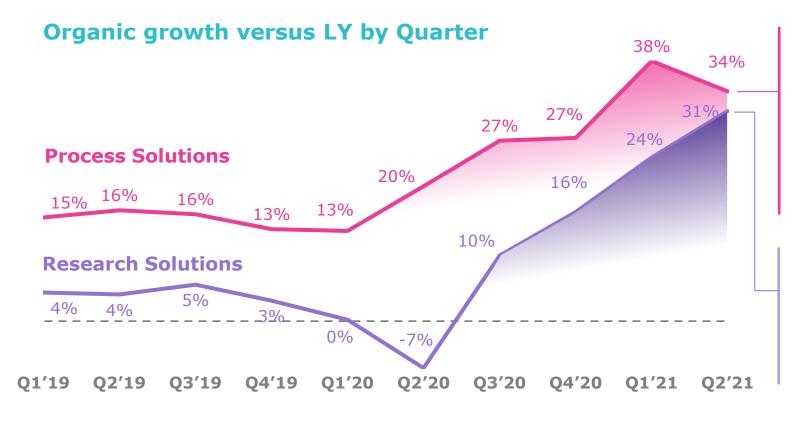
- Offering is among largest biologic reagents and hardware portfolios
- Effective vaccines and therapies start with reliable virus characterization
- Highest quality reagents needed for understanding of viral attachment, genomics, or proteomics

VACCINE & THERAPY **PRODUCTION**

- Supporting global COVID-19 vaccine and therapy response effort:
 - Upstream and downstream research and scaling
 - End-to-End solutions
 - Biosafety Testing Services



Life Science: Upside potential for Process Solutions materializing amid increasing capacity; Research Solutions gaining momentum as well



- Successful capacity ramp-up continuing
- Roughly half of growth since Q2 2020 has been COVID-19 related
- Strong underlying demand
- Order intake > +60%
- Strong core business
- Parts of growth attributable to ongoing recovery post lock-downs
- Slightly less than half of growth since Q2 2020 has been COVID-19 related



Key factors for 2021 guidance remain:

- Further progress of capacity expansions & optimizations
- Sustainable demand growth; both Covid-19 and underlying

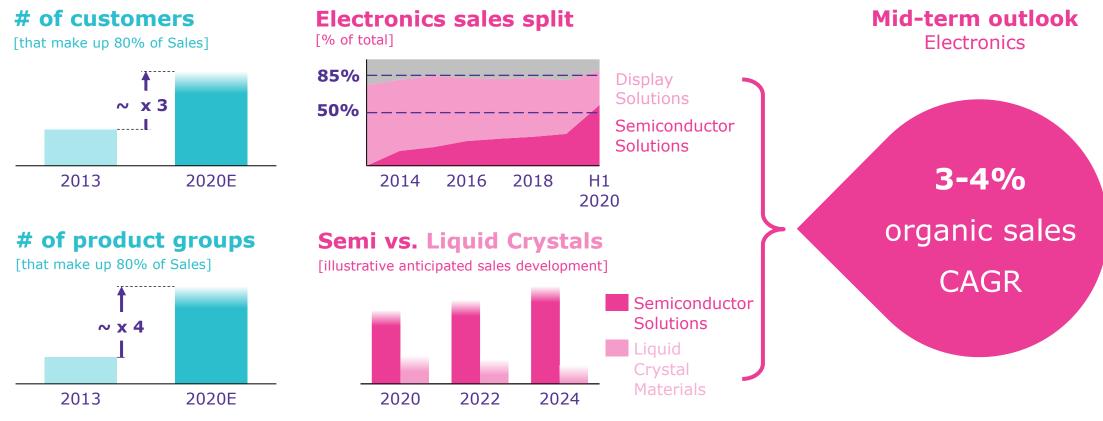




Leveraging the portfolio shift



Portfolio shift leads to greater resilience and accelerated growth





Updating mid-term financial ambition to 3-4% organic sales CAGR

Portfolio refocus drives mid-term guidance upgrade to 3 to 4% CAGR

Mid-term outlook² **Business Split**³ Sales split¹ **Semiconductor Solutions Materials** Mid- to high single-DS&S digit growth **Display Solutions** 2020 Low single-digit LC **OLED** 33% € 3.5bn decline Other, 11% **Surface Solutions** Auto

Low single-digit

growth

Fundamental growth drivers

- Continued market growth due to technological advances (Artificial intelligence, 5G, Big Data and cloud, Internet of Things) serving customers in Logic, Memory, Packaging and others
- 4 to 6% market growth⁴
- 200 to 300bps above-market growth from share gains & better portfolio (incl. 100 to 150bps additional growth from integration top-line synergies)
- Driven by trend to bigger TV size, higher resolutions, more mobile devices
- 3 to 4% growth of total LCD m² area⁵, while price pressure continues
- 18 to 22% growth of total OLED m² area⁵ with slight to moderate market share gains
- OLED material market to exceed LC material market by 2021⁶
- Well balanced exposure to automotive and cosmetics end market
- Drivers: rising living standards, higher disposable income in growing markets & higher demand for high value products at reasonable prices
- Light vehicle production and relevant cosmetics end markets returning to growth in 2021 and reaching 2019 levels by 2022 and beyond⁷

Cosmetic

Other

¹ Based on FY 2020, CAGR is organic mid-term ambition; ²growth rates are organic CAGRs; ³indicative only ⁴Source: Jan 2020 IC Insights 2018-2024 CAGR for wafer starts in million units; ⁵Source: Omdia Display Market Outlook, Q1 2020; ⁶Internal Business Intelligence; ⁷Sources: LMC Automotive Light Vehicles Forecast, Aug 2020 & Euromonitor BPC (Beauty & Personal Care) Aug 2020

5-year transformation program Bright Future is well on track



Cultural change addressed in three dedicated initiatives focused on customer centricity, market-driven innovation and corporate culture

1 New mid-term CAGR guidance starting 2020



Strategic roadmap materializing

---Measures for a bright future -----



Darmstadt

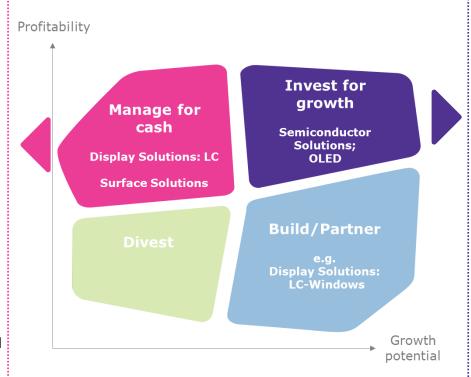
- In Darmstadt focus on R&D and production
- Immediate bottom line contribution from 2019 onwards
- Reduce the number of FTEs by ~15%
 = ~400 FTEs



 Chilworth site during September 2019 successfully closed



- Shut down of Electronics activities at Atsugi site started (to be completed during 2021)
- R&D and production activities in Atsugi transferred and consolidated in other PM locations in Asia
- Consolidation of site structure in Japan





- Leading supplier of high-purity process chemicals, gases and equipment serving semiconductor manufacturers
- Track record of accelerated growth and industry leading profitability
- Creating a leading electronic materials player with attractive long-term prospect

INTERMOLECULAR®

- Leading in advanced materials innovation
- Acquisition to strengthen semiconductor technology offering
- Application specific materials expertise with that perfectly complement Group's business and technology portfolio



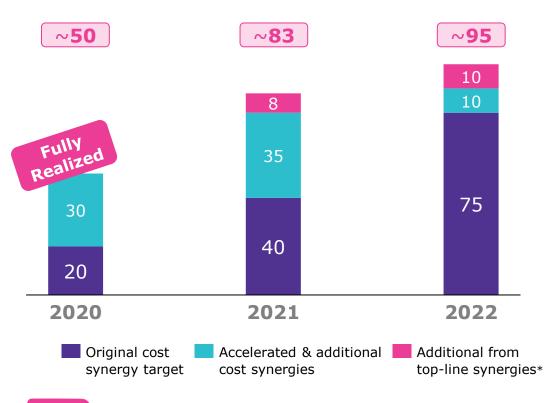
Both transactions successfully closed



Bottom-line management to support margin ambition of 30% in the long-term

Successful integration drives substantial synergy upgrade and acceleration

EBITDA pre impact of synergy ramp-up [€ m]



Sources of synergies



- Transform country setup
- Streamline duplicate structures



- Optimize production and supply chain network
- Achieve savings through joint procurement



- Integrate corporate & administrative functions
- Cost savings due to U.S. company delistings

Original target for 2022 is now being addressed for 2021

^{*}Top-line synergies from cross-selling, new products introductions and overarching initiatives

sustainability

06

Sustainability strategy enhanced, leveraging strengths with clear commitment to new targets

who we are

Targets

integration

Linked to steering and operations

Board compensation system

Currently built into part of Executive

Part of the overall strategy

Innovation Power

- We are a leading science and technology company with curious minds dedicated to human progress
- Long track-record in offering innovative products in attractive markets and serving important megatrends

Pioneering Products

- Well-equipped for developing new product classes: Portfolio of lifeimproving products in all businesses
- Enabling customers incl. scientists and developers to design next-gen products

Responsible Governance

- Resilient operations; sustainable leadership and risk-mitigation approach
- Responsibility is in our DNA: reflected by legal form, corporate governance and long history of more than 350 years

Goal #1: **Dedicated to human progress**

In 2030, we will advance human progress for more than 1 bn people through sustainable science & technology.

- and technology for our customers
- Sustainable innovations
 Impact of our technologies and products on health and well-being

Steered & Reviewed

Executive Board

Integrated

- Supervisory Board
- Corporate Sustainability Committee

Goal #2: Creating sustainable value chains

By 2030, we will integrate sustainability into all our value chains.

- Sustainability
 Sustainable culture & values and transparent supply chain
- Securing our social license to operate in all regions

Goal #3: Reducing our ecological footprint

By 2040, we will achieve climate neutrality and reduce our resource consumption.

- Climate change & emissions
- Water & resource intensity

Communicated

- Development and reporting of meaningful KPIs
- Annual Report, Sustainability Report
- Investor events

Potential to increase sustainable value for business and society

and benefit High-IMPact SDGs where we can contribute Goal 3 GOOD HEALTH AND WELL-BEING **Business opportunities** Good Health > We are able to contribute with dedicated Develop a new range of sustainable products & services, and products, know-how, partnerships and benefiting from our innovation power Well-being initiatives in pharma, science and technology. Open up additional customer groups and expand regional reach Decent Work ➤ Our ambition of future growth considers and Economic health and safety of employees also in the **Risk management** Growth supply chain. Reduce risks through higher awareness and longer-term view Secure supply chain resilience Industry, Our innovation power will lead to more Innovation **Partnerships** sustainable products and processes in various and industries. 2 Contribute as supplier of choice to customers' ESG strategy Infrastructure Improve ESG impact of our suppliers RESPONSIBLE CONSUMPTION Responsible > Being a responsible supplier, we will also 2 • Increase depth, meaning, and strategic focus of partnerships Consumption challenge suppliers to support in reaching and **Operations** 3 company targets. Production Increase attractiveness as employer 17 PARTNERSHIPS FOR THE GOALS Reduce costs of capital To unleash even more power, we foster **Partnerships** collaborations with capable partners to sum Benefit from grants and reliefs (politics, insurance, etc.) for the Goals 3 up know-how for more sustainable impact. Incentivize through integrated compensation schemes

Reduce our environmental footprint: Environmental targets 2020 have been achieved, new targets set

Achievements 2020

Reduce scope 1+2 emissions



2020 target¹: -20% 2020 result¹: -25%

Emissions target 2020 achieved!

25% overall reduction for Scope 1 and 2 emissions in 2020 relative to 2006 (planned: 20%)

Reduce water in stressed areas



2020 target²: result²: -27%

Water target 2020 achieved!

- Water use in stressed areas reduced by 27% in 2020 vs. 2014 (planned: 10%)
- By 2020, all production sites⁴ successfully implemented sustainable water management system

Reduce Group Waste Score



2025 target³: result³: -5% -4.6%

Waste target ongoing & on track!

Based on Group Waste Score, reduced environmental impact by 4.6% vs. 2016 (planned: 5% by 2025)

¹versus 2006 baseline, excluding Versum Materials

²versus 2014 baseline ³versus 2016 baseline

 4 Sites > 70.000 m 3 /a

New targets from 2021

- Aiming for climate neutrality (scope 1 to 3 emissions) by 2040
- Lower scope 1 and 2 GHG⁵
 emissions by 50% and to source
 80% of purchased electricity from
 renewable sources until 2030 vs.
 2020 baseline
- Absolute reduction of 1,500 kt⁶
 scope 3 CO₂ equivalents by 2030
- Enhancing water efficiency and improve the new Group water intensity score by 10% by 2025 vs. 2019 baseline
- Minimize negative environmental impacts, harmful emission residues should be lowered below a scientifically defined threshold by 2030

⁵GHG = Greenhouse Gas ⁶corresponds to ~30% of 2019 scope 3 emissions (current estimation incl. Versum Materials)



Next steps towards achieving ESG targets

AGENDA 2020-2022

Analysis of requirements: Strategy, business, regulation, stakeholders

Develop SBV tool² to measure product sustainability value

Link ESG¹ to board compensation

Build effective data platform for internal steering

Develop ESG KPIs for reporting

Further incorporate ESG in R&D, controlling, M&A and supply chain

Decide on dedicated investments and initiatives to achieve targets





Dedicated to human progress

Creating sustainable value chains

Reducing our ecological footprint

¹ESG: Environmental, Social, Governance



²Sustainable Business Value: Dive in deeper and read the research article on the <u>SBV method</u>

Executive Sumary

07

Full-year 2021 guidance

Net sales:

Organic: +12% to +14% YoY

FX: -2% to -4% YoY ~€18.8 - 19.7 bn

EBITDA pre:

Organic: +21% to +25% YoY (excl. Biogen1)

FX: -2% to -4% YoY ~€5.6 - 6.0 bn

EPS pre: ~ €7.80 - 8.50



¹Q3 20 reversal of the provisions for the patent litigation proceedings for Rebif in the amount of ~€365 m; Guidance including Biogen – organic EBITDA pre: +12% to +17%

Group

Looking beyond 2021

Life Science

Upgrading the COVID-19 PS floor for 2022 to €700 m

- Higher 2021 overall COVID 19 contribution of
 at least €1 bn
- Process Solution expecting more than €900 m COVID-19 related sales in 2021
- Current strong additional demand and forward visibility warrant upgrade

Healthcare

Updating
2022 pipeline sales to
€1.6 bn to €1.8 bn

- Mavenclad[®] peak sales of €1 to 1.4 bn unaffected; phased outward due to COVID-19
- Ramp-up dependent on continued recovery of dynamic market
- Anticipated positive impact of vaccines data

Electronics

guidance also **positive** signal for 2022 and beyond

- Continued strong semi market based on underlying accelerating demand
- Above mid-term guidance for 2021, further upgraded organic sales growth
- Positive market signals also anticipated to carry on mid-term



Executive SUMMary

steady earnings growth



Group

Successfully driving transformation into a leading science and technology company



setup

Three-pillar structure strengthened further as a resilient basis; COVID-19 crisis as another proof point



with high margins and a low risk profile

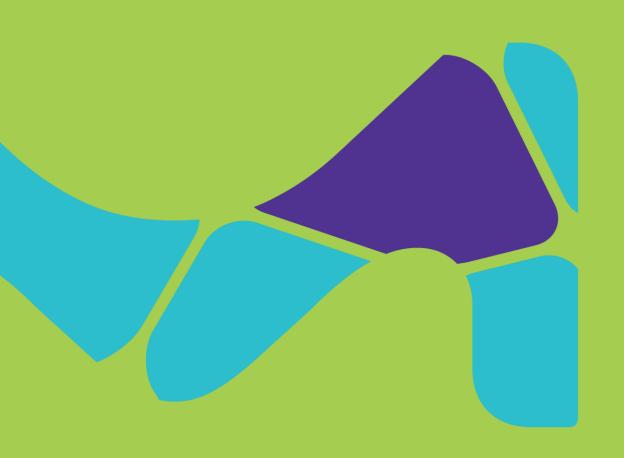
Growth Engines

Healthcare pipeline, Process
Solutions and Semiconductor
Solutions will be key drivers
of growth to 2022
and beyond



Execution

Delivery on strategic priorities ensures profitable growth; regaining financial flexibility with higher likelihood of regular bolt-ons post 2022



xibneqqa



FX sensitivity per business sector



Sales

- Global presence
- ∼35% of sales in Europe

Costs

- High cost base in EUR from main manufacturing sites
- HQ costs and large part of R&D costs in EUR





Sales

 Balanced regional sales split between EU, NA and RoW

Costs

- Extensive manufacturing and research footprint in the U.S.
- Global customer proximity requires broad-based sales force





Sales

- ~75% of sales in Asia-Pacific
- Industry is USD-driven

Costs

- Main production sites in NA, APAC and Germany, several R&D and production facilities in Asia
- Improved natural hedge post Versum acquisition



FX impact on EBITDA pre²
Low High



Acquisition history - Strengthening leadership positions

Year	Company	Volume	Sector
2021	Amptec	n/a	Life Science
2020	Resolutions Spectra Systems	n/a	Life Science
	FloDesign Sonics	n/a	Life Science
2019	Versum Materials Inc., USA	€5.8 bn	Electronics
	Intermolecular Inc., USA	€62 m	Liectionics
	Grzybowski Scientific Inventions Ltd., USA	€7 m & €1 m	Life Science
2017	Natrix Separations Inc., Canada	€12 m & €7 m	Life Science
	BioControl	€161m	Life Science
	Sigma-Aldrich	€13.1 bn	Life Science
2015	Qlight Nanotech Ltd. (Israel)	n/a	Life Science
	Ormet Circuits Inc. (USA)	n/a	Electronics
2014	AZ Electronics	€1.9 bn	Electronics
	Peer+	n/a	Liectionics
2012	Biochrom AG	n/a	Life Science
	CellASIC (USA)		
2011	Amnis (USA)	n/a	Life Science
	heipha Dr. Müller GmbH / Hycon branch of Biotest AG	n/a	
2010	Beijing Skywing Technology (China)	€14.5 m	Life Science
	Millipore Corporation (US)	€5.1 bn	
2009	Taizhu (China)	€26.3 m	Electronics
2008	Bio-Fyt Pharma (Belgium)	€30.0 m	Healtchare
	SeQuant	€6.9 m	Life Science
2006	Serono (Switzerland)	€10.3 bn	Healthcare
	Agribiotics Holdings (Canada)	€21.9 m	Electronics
2005	Survac (Denmark)	€10.8 m	Healthcare
	Avecia (UK) incl. Covion (Germany)	€49.3 m	Electronics





Divestment history - Focusing on profitable core areas

Year	Company	Volume	Sector
2020	Litec-LLL GmbH Allergopharma	€3 m n/a	Electronics Healthcare
2018	Consumer Health	€3.4 bn	Healthcare
2018	Flow Cytometry business	€62.5 m	Life Science
2017	Biosimilars	€656 m	Healthcare
2016	Pakistan Healthcare business	n/a	Healthcare
2014	Discovery and Development Solutions	€22.6 m	Life Science
2012	Mobile Energy Business (Germany)	n/a	Electronics
2010	Crop BioScience Business (USA, CA and Argentina) Théramex (Monaco and Italy)	€208.2 m €269.3 m	Healthcare
2007	Generics	€4.9 bn	Healtchare
2006	ITO/CF Business (Taiwan)	€29 m	Electronics
2005	Electronic Chemicals Business	€270 m	Electronics
2004	Lab distribution VWR (US)	\$1.68 bn	Healthcare
2004	50% holding of Biomet	\$300 m	Electronics





Successful crisis management increasingly mitigates pandemic impact



Guidance upgrade proof point of excellent crisis management and strong business performance

Group

2020 – strong resilience in times of global crisis

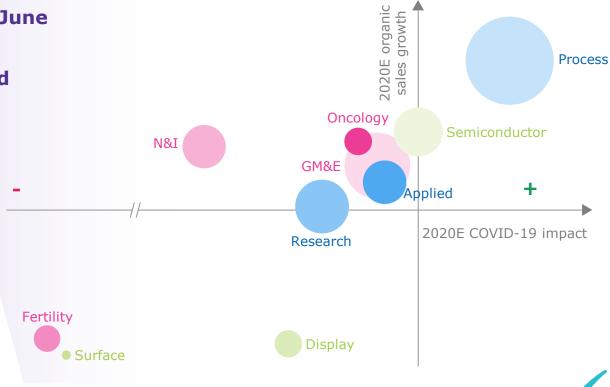
2020 guidance confirmed; recovery started in June

- Most businesses growing despite COVID-19
- Largest business growing and positively affected
- Smallest businesses with biggest impact

Delivery on priorities during crisis

- √ Health & safety of employees
- ✓ Business continuity
- ✓ Contributions to public health and society
- ✓ Sustainability aspects further enforced





CMD 2019

Group- steady earnings growth with high margins and a low risk profile



¹ Indicative only and based on guidance from August 6: slight to moderate organic sales and EBITDA pre growth, COVID-19 with up to a mid single-digit impact on sales of which 50-60% hitting EBITDA pre

2021 business sector guidance¹

Healthcare



Net sales

- Organic: +7% to +10%
- Mainly driven by Mavenclad[®], Bavencio[®] and recovery of Fertility
- Base business organically around stable

EBITDA pre

- Organic: +15% to +18% YoY (excl Biogen²)
- FX: -5% to -7% YoY
- ~€2,050 2,150 m

Net sales



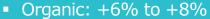
Process Solutions as main growth driver

EBITDA pre

- Organic: +30% to +34% YoY
- FX: -1% to -3% YoY
- ~€3,050 3,200 m

Electronics

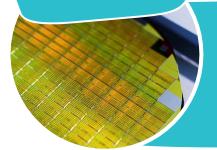
Net sales



- Strong Semiconductor Solutions contribution
- OLED with high growth

EBITDA pre

- Organic: +9% to +12% YoY
- FX: -2% to -4% YoY
- ~€1,070 1,130 m





Additional financial guidance 2021

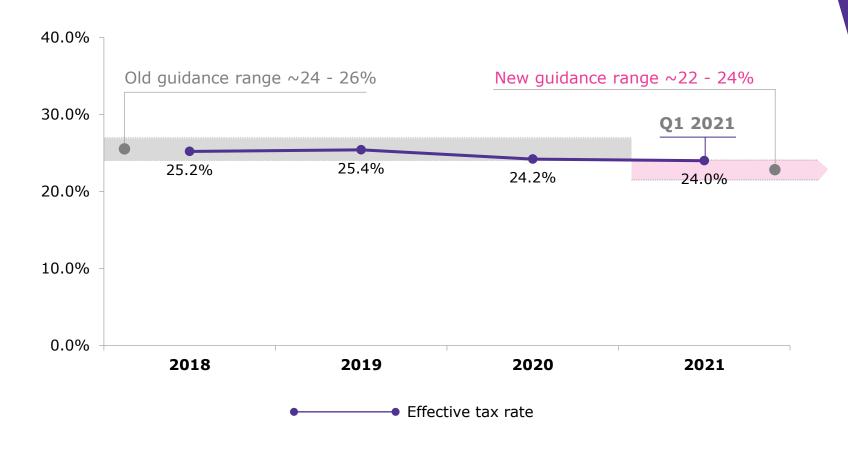
Further financial details

Corporate & Other EBITDA pre	~ €-450 to -500 m
Interest result	~ €-220 to -245 m
Effective tax rate	~22% to 24%
Capex on PPE	~€1.4 to 1.5 bn
Hedging/USD assumption	FY 2021 hedge ratio ~70% at EUR/USD ~1.17
2021 Ø EUR/USD assumption	~1.19 to 1.23



Effective tax rate guidance lowered to new range of 22% to 24%

Tax rate development 2018-2020 and from 2021 onwards



Rationale for update

Strong profit growth in Life Science results in different profit contributions worldwide, leading to a lower overall tax rate

New **resulting underlying tax** rate
used for EPS pre
calculation is now 23%



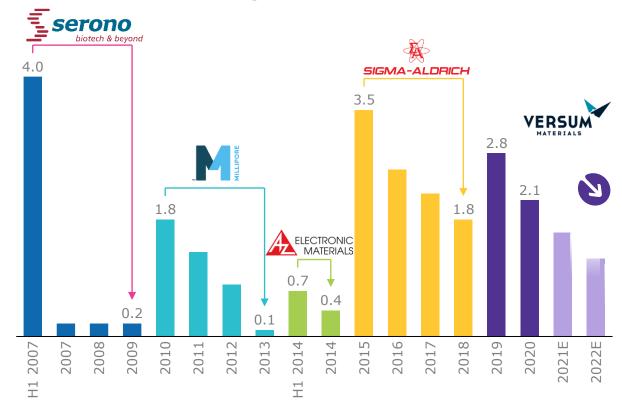
Group

Focus on organic growth and further deleveraging

Proven swift deleveraging after major acquisitions

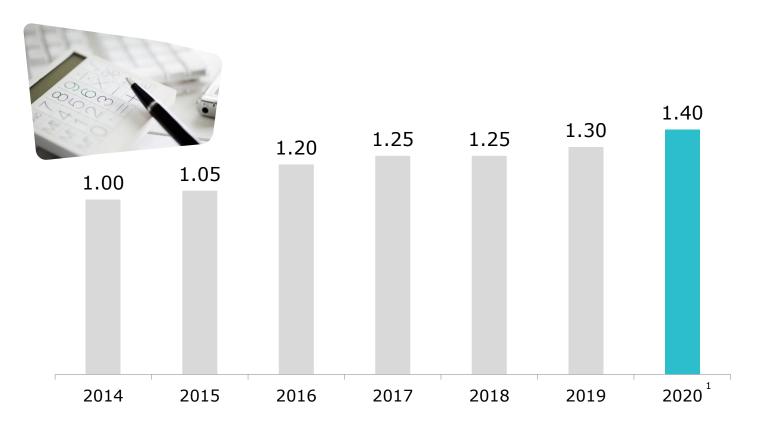
- Deleveraged to ~2x net debt/EBITDA pre already in 2020
- M&A on hold until 2022; only smaller deals to be realized if budget available
- New mid-term capex ceiling of ~€1.4 – 1.5 bn reflects increased focus on organic investment
- Dividend policy mirrors
 sustainable earnings trend

Net debt / EBITDA pre track record & outlook



Sustainable dividend growth

Dividend development 2014 - 2020



2020 dividend

- Dividend of €1.40 (+8% YoY) per share approved¹ by Annual General Meeting
- Payout ratio of 23.1% of EPS pre² in 2020; aiming for 20-25% of EPS pre
- Dividend yield³ of 1.0%

³Calculated with 2020 year-end share price of € 140.35 per share.



¹April 28, 2021: Pay Date

²Excluding Biogen provision release, including the provision release the ratio is 20.9%

Governance

Merck's KGaA, Darmstadt, Germany ownership structure

Shareholders of Merck KGaA, Darmstadt, Germany

- Shareholders hold ~30% of Group's total capital
- Share capital is divided into 129,242,252 bearer shares and one name share without nominal value
- Listed at German Stock Exchange's Prime Standard, member of the DAX 30
- Only bearer shares entitled to vote at Group's Annual General Meeting: one share, one vote

Supervisory Board

Its Chairman leads AGM, representing interests of Group's shareholders:

- Approving financial statements of the Group
- Working together with Executive Board, receiving reporting on progress regarding business and financial development at the Group, Containing 16 members: 8 employee representatives, 8 independent representatives (2 family representatives and 6 elected shareholder representatives)
- 2 committees: Nomination and Audit Committee

Nomination Committee
Defines criteria and makes
proposals for new candidates

Audit Committee
Control systems, external
auditing, financial statements

Finance Committee financial statements, external auditing

70%

E. Merck KG, Darmstadt, Germany: Partners' Meeting & Family Board • E. Merck KG, Darmstadt, Germany (99.9% The Family)

- E. Merck KG, Darmstadt, Germany (99.9% The Family) holds ~70% of Merck KGaA's, Darmstadt, Germany total capital
- These 70% of the total capital are not entitled to vote at Merck's KGaA, Darmstadt, Germany AGM
- A Family Board represents the entrepreneurial interests of the Family (elected by partners' meeting)

Board of Partners

Elected by Family Board, and takes over some duties of a usual Supervisory Board:

- Appointment, dismissal and supervision of Executive Board members of Merck KGaA, Darmstadt, Germany
- Approves extraordinary business transactions and annual financial statements of the Group
- Contains 5 members of the Family Board and 4 external members of the business community
- 3 committees: Finance, Personnel and R&D Committee

Personnel Committee Executive board members, contracts, compensation R&D Committee research activities of all sectors: HC, LS, EL



Executive board compensation

Pay for performance reflecting the company's long-term strategy

		Long-Term Incentive Plan					
riable	40-50%	 Reflecting the long-term strategy for Company's growth and (from 2022) sustainability ambition 4 years performance cycle: 3 years target achievement + 1 year holding period Based on virtual Group Share Units (Grant € divided through start share price, multiplied with the end share price) Financial targets: 50% Group Share Price vs. DAX + 25% EBITDA pre margin + 25% Organic sales growth From 2022 multiplied with sustainability factor (0.8-1.2) reflecting KPIs from each of the sustainability goals Corridors for each target and achieved targets published transparently ex-post in the compensation report Maximum cap: Maximum pay out 250%, maximum € cap for LTIP for each board member published Claw-back allows to retain amounts allocated from the Long-Term Incentive Plan Performance of Group share price vs. the DAX CBITDA pre margin in relation to target value Organic sales growth in 25% Organic sales growth in 25% Oseach Sustainability factor Claw-back allows to retain amounts allocated from the Long-Term Incentive Plan 					
Va		Profit Sharing					
		• Three-years average profit after tax of the E. Merck KG, Darmstadt, Germany, multiplied with individual permille rate					
	25-35%	• From 2021 reduced individual performance factor of 0.8-1.2 can increase (bonus) or decrease (malus) the amount based on a set of control incl. the 3 sustainability goals, disclosure of catalogue and reasons for if performance factor ≠1.0					
		 Individual permille rate for each board member and maximum € cap for each board member published 					
		 Staggered incentivization and minimum threshold value and maximum limit for profit after tax (0.75/2.0 bn €) 					
=	6-9%	 Mandatory personal investment in Group Shares amounting to one third of the net payment of the profit sharing (4 year holding period) Pension Entitlements Defined contribution 					
	0-3%	Additional Benefits Mainly contributions to insurance policies, personal security expenses, company car					
sic		Basic Compensation					
Bas	1F 200/	Fixed and non-performance related compensation					
_	15-20%	Paid in 12 equal monthly installments					
		 1.4 million € for the chairman / up to 1.1 million € for the members of the executive board 					
	Maximum	n total compensation: reduced to €11.5 m Chairman, €9.5 m other executive board members					

External stakeholders assess our engagement



As of 2021, we received an MSCI ESG* Rating of AAA.

*Environment, Social, Governance



2021, we received an **ESG Risk Rating** of 19.5 and **Sustainalytics: low risk** of experiencing material financial impacts from ESG factors.



Since 2008, Group is part of FTSE4Good Index, measuring the performance of companies with strong ESG practices (top 15).



In 2020, Group has once more achieved **prime status** by **ISS Oekom**.



In 2019, the Group share was again included in STOXX Global ESG Leaders Index.

a sustainability index based on key environmental, social and governance criteria.



We have been **reconfirmed** as a constituent of the **Ethibel Sustainability Index** (ESI) **Excellence Europe** since May 2020, based on VigeoEiris.



Group for the second time received platinum status in 2021, among the **top 1% of companies**. **EcoVadis** annually examines ~75,000 suppliers from 160 countries.



In the 2021 Access to

Medicine Index Group
ranked eighth place. We
were recognized for our
performance in R&D, where
we ranked fifth.



CDP Climate: In 2020, we scored **"B"** (2019: C). **CDP Water**: In 2020, we received a **"B"** (2019: B).



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Group

Regular portfolio review remains key to success

strong track record

- Acquisitions and divestments are part of Company's history
- Licensing and partnerships remain on our agenda
- All prior transactions earned their cost of capital



pefining portfolio guard rails

- Three strong pillars with no business marginalized
- Leading market position in attractive markets
- Focus on innovation and sustainability through science and technology



clear financial M&A criteria

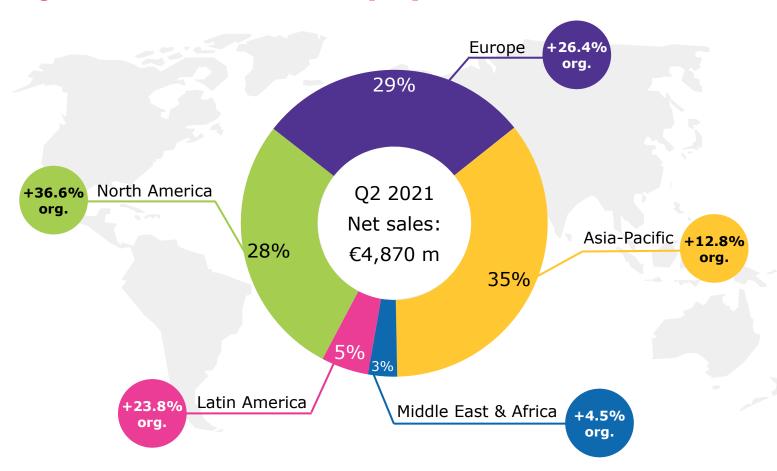
- Supporting profitable growth strategy
- IRR > WACC
- EPS pre accretive
- Maintain investment grade rating



- Current set-up is strong and organic investment opportunities are attractive
- Expect to regain financial flexibility by 2022 to pursue external growth opportunities
- Targeted and more regular bolt-on approach more likely than large transformative deals

Life Science demand and Healthcare recovery drive particularly strong growth in North America and Europe

Regional breakdown of net sales [€m]



Regional organic development

- APAC: Strong growth across all sectors particularly in Process Solutions,
 Fertility and Semiconductor Solutions
- Europe: Strong demand in Process Solutions and strong recovery in Fertility against heavily impacted Q2 2020 drive 26% growth
- North America: Growth across all sectors, particularly strong Life Science, Fertility & Oncology (supported by Eli Lilly supply agreement)
- LATAM growth driven foremost by Fertility and CM&E
- Fertility drives growth in ME&A

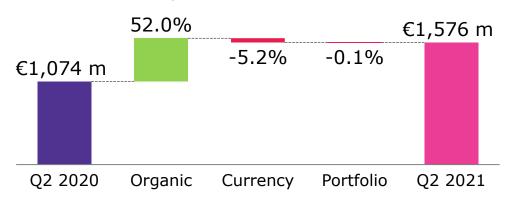


Strong Life Science demand and robust Healthcare recovery drive 23% sales and 52% EBITDA pre organic growth

Q2 YoY Net Sales				
	Organic	Currency	Portfolio	Total
Healthcare	23.6%	-4.3%	0.0%	19.2%
Life Science	28.2%	-5.0%	0.0%	23.2%
Electronics	10.3%	-5.0%	0.0%	5.4%
Group	23.0%	-4.8%	0.0%	18.2%

- Strong recovery in Fertility well above pre COVID-19 levels, organic Mavenclad[®] growth of 102% and Oncology organic growth of +49% drive +24% growth in Healthcare overall
- Record 28% organic growth in Life Science; driven by all businesses with Process Solutions up +34%; Research Solutions +31% and Applied Solutions +13% against soft comps from lockdown
- Electronics growing 10% organically, driven by strong performance in Semiconductor Solutions (+12% org.) and strong recovery of Surface Solutions while Display Solutions declines slightly

Q2 YoY EBITDA pre



- Organic EBITDA pre increases by more than 50% and more than twice as fast as sales
- Strong uptake in Life Science and Healthcare gross profit paired with continued cost discipline in all sectors vs. soft Q2 2020
- FX burden of -5% across various currencies with largest negative impact from USD and JPY; partly mitigated by hedging

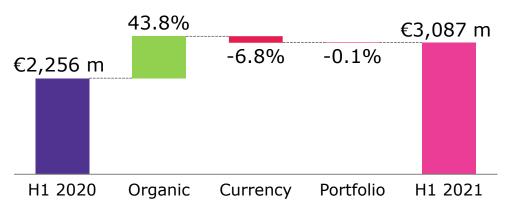


Strong performance across all sectors drives 17% organic net sales growth and 44% organic EBITDA pre growth against COVID-19-impacted H1 2020

H1 YoY Net Sales				
	Organic	Currency	Portfolio	Total
Healthcare	12.9%	-5.2%	-0.6%	7.1%
Life Science	27.5%	-5.6%	0.0%	21.8%
Electronics	5.0%	-4.7%	0.0%	0.3%
Group	17.4%	-5.3%	-0.2%	11.9%

- Healthcare: +13% org. growth vs. COVID-19-impacted H1 2020, driven by strong recovery in Fertility, growth in Mavenclad[®] & Bavencio[®]; supported by Erbitux[®] Eli Lilly supply agreement
- Life Science: Up +28% as strong base business across all BUs is boosted by additional COVID-19 demand in Process and Research Solutions against lockdown-related soft comps
- Electronics: Grows +5% (above mid-term guidance) as strong performance in Semiconductor Solutions and recovery of Surface Solutions overcompensate stabilizing Display decline





- EBITDA pre grows faster than sales organically, largely driven by strong operating leverage in Life Science and Healthcare vs. a weaker H1 2020
- FX headwinds primarily from USD, JPY & BRL amount to burden of -5% on net sales and -7% on EBITDA pre



Credit details

Maturity profile as of June 30, 2021

Coupon [€m / USDm] 3.250% 0.125% 2.950% 1.375% 1 600 3.375% 1.625% 2.875% 1 000 0.875% 0.500% 0.005% 0.375% 1.000 1.000 800 750 750 600 500 600 550 2021 2022 2023 2024 2025 2026 2027 2028 2029 2030 2031 ■ EUR bonds ■ USD bonds ■ Hybrids (first call dates)

Credit rating information

	LT Rating	Since	Outlook	ST Rating
Moody's	Baa1	12.12.14	Stable	P-2
S&P Global	А	29.05.13	Stable	A-1
SCOPE	A-	19.10.16	Stable	S-1



Q2 2021: Overview

Key figures

[€m]	Q2 2020	Q2 2021	Δ
Net sales	4,119	4,870	18.2%
EBITDA pre	1,074	1,576	46.7%
Margin (in % of net sales)	26.1%	32.4%	6.3pp
EPS pre	1.30	2.24	72.3%
Operating cash flow	502	888	76.9%
[€m]	Dec. 31, 2020	June 30, 2021	Δ
Net financial debt	-10,758	-10,141	-5.7%
Working capital	3,938	4,222	7.2%
Employees	58,096	58,382	0.5%

- EBITDA pre & margin increase, driven by operating leverage in Life Science and Healthcare vs. LY COVID-19 burden
- EPS pre increase driven by EBIT pre, better financial result and lower tax rate vs. particularly soft Q2 2020
- Operating cash flow up 77% driven by higher profit after tax in all three sectors
- Ongoing reduction of net financial debt



H1 2021: Overview

Key figures

H1 2020	H1 2021	Δ
8,489	9,501	11.9%
2,256	3,087	36.9%
26.57%	32.49%	5.919%
2.80	4.42	57.9%
1,019	2,104	106.6%
Dec. 31, 2020	June 30, 2021	Δ
-10,758	-10,141	-5.7%
3,938	4,222	7.2%
58,096	58,382	0.5%
	8,489 2,256 26.57% 2.80 1,019 Dec. 31, 2020 -10,758 3,938	8,489 9,501 2,256 3,087 26.57% 32.49% 2.80 4.42 1,019 2,104 Dec. 31, 2020 June 30, 2021 -10,758 -10,141 3,938 4,222

- Strong performance across all sectors drives +12% growth despite -6% FX
- EBITDA pre & margin increase, driven by operating leverage in Life Science and Healthcare vs. pandemic-impacted LY; further supported by Erbitux[®] Eli Lilly supply agreement (+€49 m net sales)
- EPS pre above last year driven by strong operating performance, supported by better financial result & lower tax rate
- Operating cash flow more than doubles as strong EBITDA pre growth supported by favorable net working capital



Q2 2021: Reported figures

Reported results

[€m]	Q2 2020	Q2 2021	Δ
EBIT	491	1,049	113.6%
Financial result	-102	-95	-7.3%
Profit before tax	389	955	145.4%
Income tax	-100	-208	107.9%
Effective tax rate	25.7%	21.8%	-3.9pp
Net income	290	745	157.1%
EPS (€)	0.67	1.71	155.2%

- EBIT more than doubles, driven by strong performance across all sectors vs. Q2 2020 COVID-19 burden
- Improved financial result largely driven by lower interest expense from deleveraging
- Effective tax rate benefitting from boosted results in Life Science (better country mix)
- Strong EBIT growth, improved financial result and lower tax rate drive higher net income & EPS



H1 2021: Reported figures

Reported results

[€m]	H1 2020	H1 2021	Δ
EBIT	1,207	2,092	73.3%
Financial result	-201	-154	-23.3%
Profit before tax	1,006	1,939	92.6%
Income tax	-259	-444	71.2%
Effective tax rate (%)	25.8%	22.9%	-2.9pp
Net income	746	1,492	100.0%
EPS (€)	1.72	3.43	99.4%

- EBIT increase driven by strong growth and operating leverage in all business sectors, particularly Life Science
- Improved financial result largely driven by lower interest expenses in line with deleveraging
- Effective tax rate in the middle of updated guidance range
- Doubled net income and EPS reflect positive development of EBIT, financial result and tax rate



Cash flow statement

Q2 2021 – Cash flow statement

[€m]	Q2 2020	Q2 2021	Δ
Profit after tax	289	747	458
D&A	559	419	-140
Changes in provisions	-54	88	142
Changes in other assets/liabilities	-166	-217	-50
Other operating activities	-13	19	32
Changes in working capital	-112	-168	-56
Operating cash flow	502	888	386
Investing cash flow	-216	-241	-25
thereof Capex on PPE	-194	-256	-62
Financing cash flow	-302	-1,059	-757

Cash flow drivers

- Operating cash flow up €386 m driven primarily by higher profit after tax in all three sectors, particularly Life Science
- Delta in D&A driven by COVID-19 related impairments in Q2 2020
- Provisions up, driven by litigation accruals, pension fluctuations, and LTIP*
- Higher outflow from working capital but growing considerably slower than sales
- Higher investments, particularly CAPEX on PPE in line with ongoing capacity expansion
- Financing cash flow explained by net repayment of bonds, bank liabilities and commercial papers

^{*} Long Term Incentive Plan

Totals may not add up due to rounding

Cash flow statement

H1 2021 – cash flow statement

[€m]	H1 2020	H1 2021	Δ
Profit after tax	747	1,495	748
D&A	991	843	-148
Changes in provisions	-38	55	93
Changes in other assets/liabilities	-189	-56	133
Other operating activities	-24	25	48
Changes in working capital	-468	-256	212
Operating cash flow	1,019	2,104	1,086
Investing cash flow	-504	-587	-83
thereof Capex on PPE	-532	-564	-33
Financing cash flow	239	-1,054	-1,293

Cash flow drivers

- Operating cashflow more than doubles as strong profit after tax is further boosted by favorable working capital
- Lower depreciation & amortization vs.
 H1 2020 which was elevated by COVID-19-related impairments in EL
- Changes in provisions elevated by mid double-digit €m litigation accrual
- Changes in other assets and liabilities largely explained by Q1 tax positions
- Favorable lower increase in working capital driven by higher inventories in 2020 to secure supply amid COVID-19
- Financing cash flow explained by net repayment of bonds, bank liabilities and commercial papers



Adjustments in Q2 2021

Adjustments in EBIT

[€m]	Q2 20	020	Q2 20	021
	Adjustments	thereof D&A	Adjustments	thereof D&A
Healthcare	15	0	11	3
Life Science	-15	0	-6	0
Electronics	131	112	10	5
Corporate & Other	8	0	97	0
Total	138	112	112	8



Adjustments in H1 2021

Adjustments in EBIT

[€m]	H1 20	020	H1 20	021
	Adjustments	thereof D&A	Adjustments	thereof D&A
Healthcare	-12	2	21	3
Life Science	-4	0	8	0
Electronics	165	112	27	7
Corporate & Other	25	0	103	1
Total	174	114	159	11





Financial Calendar

Date	Event
August 5, 2021	Q2 2021 Earnings release
September 9, 2021	Virtual Capital Markets Day
November 11, 2021	Q3 2021 Earnings release
April 22, 2022	Annual General Meeting
August 4, 2022	Q2 2022 Earnings release



нealthcare

China's VBP: Round 4 implementation near completion, confidence in approx. stable base business through 2021 and 2022 sustained



Round 1	Round 2		Round 3		Round 4	
2019			2020		ATE 2021	
March: First round initiated, no Group products impacted	December: Round 2 announced (incl. bisoprolol)	April: Round 2 winners granted exclusive access to 60% of total hospital market ¹ , non- winners (incl. Concor®) retain access to remaining 40% of hospital market at a gradient price cut ² + can freely compete in non-hospital/ retail market	July: Round 3 announced (incl. Metformin IR and XR) ¹	August: Bidding manufacturers), (• Access retained market (~30% of the hospita) • Continued grown	Jan: Round 4 announced (incl. calcium dobesilate³ and canagliflozin³), bidding completed on Feb 3, Invokana® & Doxium® not among winners, completed, 8 winners (all Chinese Glucophage® not among winners ed to non-hospital/retail of total market), and ~20% I market (at gradient price cut) with of Chinese metformin p.a. over the next 5 years	



China Glucophage sales represent only ~8% of the total base business (2020 net sales)
Sustained confidence in approx. stable base business (org.) through 2021 and 2022

^{1:} hospital market for bisoprolol and metformin makes up ~70% of total market, this includes urban hospitals, rural hospitals, and community health centers; 2: Concor® price cut in the high single digit %; 3: alliance products; Acronyms: VBP = Volume-Based Procurement



Phase I

M1231 Bispecific MUC1xEGFR ADC

Solid tumors

M1774 ATR inhibitor Solid tumors¹

M4076 ATM inhibitor Solid tumors

peposertib DNA-PK inhibitor Solid tumors² bintrafusp alfa TGFbeta trap/anti-PD-L1 Cervical cancer 1L

M6223 anti-TIGIT mAb Solid tumors³

enpatoran (M5049) TLR7/8 antagonist Systemic lupus erythematosus / Cutaneous lupus erythematosus

M5717 PeEF2 inhibitor Malaria

Phase II

berzosertib ATR inhibitor

Small-Cell Lung Cancer⁴

tepotinib
MET kinase inhibitor
Metastatic Colorectal Cancer

RAS/BRAF wt, MET amplified⁵
tepotinib

MET kinase inhibitor Non-small cell lung cancer, *EGFR* mutant, *MET* amplified⁶

bintrafusp alfa TGFbeta trap/anti-PD-L1 Non-small cell lung cancer 1L/2L

bintrafusp alfa TGFbeta trap/anti-PD-L1 Locally advanced non-small cell lung cancer bintrafusp alfa
TGFbeta trap/anti-PD-L1

Biliary tract cancer 1L

bintrafusp alfa TGFbeta trap/anti-PD-L1 Cervical cancer 2L

bintrafusp alfa TGFbeta trap/anti-PD-L1

Triple negative breast cancer (HMGA2 positive)

enpatoran (M5049) TLR7/8 antagonist COVID-19 pneumonia

- Oncology
- Immuno-Oncology
- Immunology
- Neurology
- Global Health

Phase III

xevinapant IAP inhibitor

Locally advanced squamous cell carcinoma of the head and neck^{7,8}

avelumab anti-PD-L1 mAb Non-small cell lung cancer 1L

evobrutinib BTK inhibitor Relapsing multiple sclerosis

Registration

tepotinib MET kinase inhibitorNon-small cell lung cancer, *MET*ex14 skipping⁹

ADC: Antibody Drug Conjugate; 1L: first-line treatment; 2L: second-line treatment

¹ Study as monotherapy and in combination with niraparib. ² Study in combination with avelumab. ³ Includes study in combination with bintrafusp alfa. ⁴ Includes studies (phase I/II) in collaboration with/ sponsored by external partners, e.g. US National Cancer Institute (NCI). ⁵ In combination with cetuximab. ⁶ In combination with osimertinib. ⊓ In unresected LA SCCHN patients eligible for cisplatin. ७ On March 01, 2021, Merck KGaA, Darmstadt, Germany announced a worldwide in-licensing agreement with Debiopharm, Switzerland, for the development and commercialization of xevinapant (Debio 1143). ⁰ As announced on November 26, 2020, the European Medicines Agency (EMA) has validated for review the application for tepotinib for the treatment of adult patients with advanced non-small cell lung cancer.

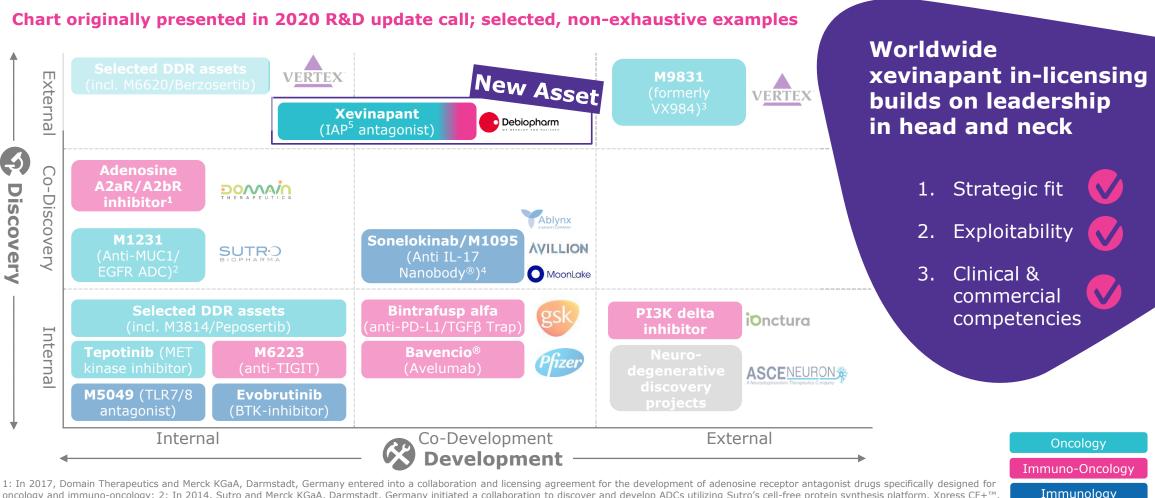
Additional information: Several combination studies (phase II) of avelumab with talazoparib, axitinib, ALK inhibitors or chemotherapy ongoing under sponsorship of Pfizer.

Unless noted otherwise, clinical programs conducted in collaboration with external partners are not shown unless Merck KGaA, Darmstadt, Germany has co-ownership of data. In such case the indication is shown in *Italics*. Pipeline products are under clinical investigation and have not been proven to be safe and effective. There is no quarantee any product will be approved in the sought-after indication.



Xevinapant

Potential to become standard of care in core area for the Group



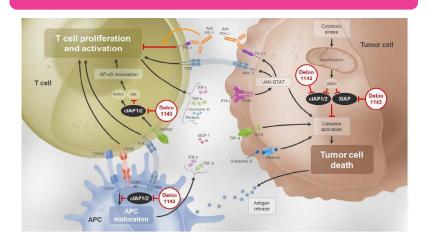
1: In 2017, Domain Therapeutics and Merck KGaA, Darmstadt, Germany entered into a collaboration and licensing agreement for the development of adenosine receptor antagonist drugs specifically designed for oncology and immuno-oncology; 2: In 2014, Sutro and Merck KGaA, Darmstadt, Germany initiated a collaboration to discover and develop ADCs utilizing Sutro's cell-free protein synthesis platform, Xpress CF+TM. Merck KGaA, Darmstadt, Germany is responsible for drug product, clinical development and commercialization of any resulting products; 3: In 2019, an exclusive license was granted to Vertex for the use of M9831 in gene-editing applications; 4: Avillion conducted Ph II of M1095 in Psoriasis, Group decided to out license sonelokinab to a new partner to initiate Phase III development in 2021 5: Inhibitor of Apoptosis Proteins



Xevinapant (Debio 1143)

Potentially first in class oral IAP antagonist with FDA BTD

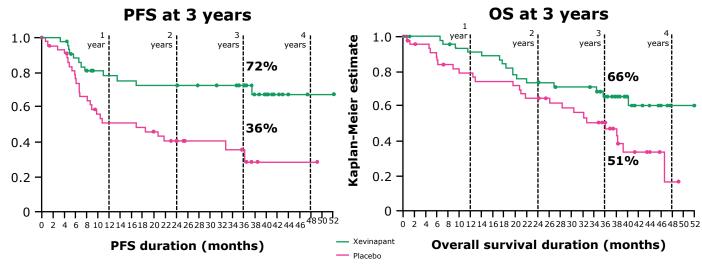
Mode of Action¹



- Oral Inhibitor of Apoptosis Proteins (IAP) antagonist: chemo-/radio-sensitizer & enhancer of anti-tumor immunity
- IAP antagonists tackling two cancer hallmarks:
 - Enhancing anti-tumor immunity
 - Lowering threshold for tumor cell death

Compelling Phase 2 data² published in *The Lancet Oncology*, and presented at ESMO 2020

- Improvement in OS statistically significant and clinically meaningful: HR 0.49 (0.26-0.92); p=0.0261
- Clinically compelling PFS improvement: HR 0.34 (0.17-0.68); p=0.0023
- Predictable and manageable safety profile without substantial additional toxicity to standard CRT



Acronyms: BTD = Breakthrough Therapy Designation; IAP = Inhibitor of Apoptosis Proteins; 1: Debiopharm; 2: ESMO 2020 - Late Breaking Abstract 39 - 3-years follow-up of double-blind randomized phase II comparing concurrent high-dose cisplatin chemo-radiation plus xevinapant or placebo in high-risk patients with locally advanced squamous cell carcinoma of the head and neck



Xevinapant

Total deal-volume of up to ~ €900 m and industry-typical sales royalties

Payment type	Amount (in €)	Accounting treatment ²
Upfront payment	~ €190 m Largest part to be capitalized as an intangible asset	
Approval milestones	Up to ~ €380 m ¹	To be paid and capitalized as an intangible asset upon approval and to be amortized once asset is ready for use
Commercial milestones	Up to ~ €330 m	To be paid and capitalized as an intangible asset, based on sales thresholds and to be amortized over remaining useful life
Sales	n/a	Merck KGaA, Darmstadt, Germany to recognize sales globally (incl. US)
R&D Costs	n/a	For ongoing TrilynX study Cash view: 50/50 cost sharing P&L view: fully shown in Company P&L 2nd study for cisplatin-ineligible patients: Group incurs 100% of cost
Royalties	n/a	Group to pay industry-typical sales royalty to Debiopharm

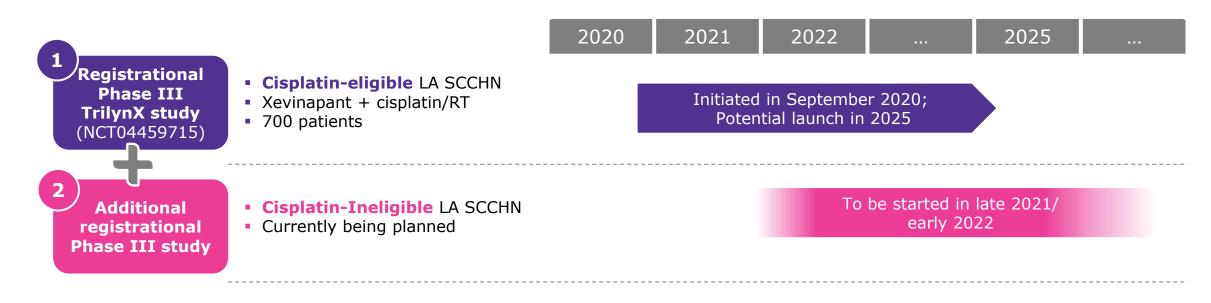
¹ thereof up to ~€300 m for focus H&N indications)



² final accounting treatment is still subject to alignment with auditors

Xevinapant (Potentially first in class oral IAP antagonist)

Two Phase III studies are designed to target the majority of unresectable LA SCCHN patients receiving systemic therapy + RT





Blockbuster potential provided success of both studies



20,000+ unresectable LA SCCHN patients in US and EU-5 each



Tepotinib (MET kinase inhibitor)

First-in-class launch in MET Exon14 sets foundation for EGFRm/METamp opportunity and exploration in other tumor types





Lay the Foundation in NSCLC - MET Exon14

(VISION study, NCT02864992 3-5% of total NSCLC population)

- Highly competitive data set presented at ASCO 2020 and published in New England Journal of Medicine in May 2020 (99 patients with a follow up ≥9 mths)
- First-in-class launch¹ in Japan² in March 2020, Sakigake designation³ granted in 2018
- Approval by US FDA¹ received on February 3, 2021



Tap into a growing opportunity in NSCLC - EGFRmut/ METamp

(INSIGHT 2 study, NCT03940703 2-5% of total NSCLC population)

- Increased EGFRm detection with testing and treatment moving into earlier lines of therapy (ADAURA trial demonstrates a 79% reduction in the risk of death with Osimertinib in the adjuvant setting (ASCO 2020), suggesting an even greater uptake of Osimertinib)
- **METamp as the primary driver of resistance** Some publications suggest that METamp resistance post-Osimertinib could be ~25%⁴



Exprore EGFR
resistance in crc –
Tepotinib + Erbitux® combo
(NCT04515394)

Opportunity for Tepotinib to address an unmet need in metastatic colorectal cancer (mCRC) together with Erbitux® ERBITUX

1: approved for both treatment naïve and previously treated METex14 positive NSCLC patients; 2: second largest Oncology market globally; 3: SAKIGAKE designation promotes research and development in Japan, aiming at early practical application for innovative pharmaceutical products; 4: Piotrowska et al., "Landscape of Acquired Resistance to Osimertinib in EGFR -Mutant NSCLC and Clinical Validation of Combined EGFR and RET Inhibition with Osimertinib and BLU-667 for Acquired RET Fusion", AACR Cancer Discovery 2018; Acronyms: CRC = Colorectal cancer; EGFR = Epidermal Growth Factor Receptor; NSCLC = Non-small cell lung cancer



Tepotinib (MET kinase inhibitor)

Tapping into the rapidly evolving EGFRmut/METamp market – Encouraging INSIGHT 1 data





INSIGHT 2 – Tepotinib + Osimertinib in Osimertinib Relapsed METamp NSCLC

- Study design recently amended to reflect evolved and future standard of care:
 - Target population Inclusion criteria adjusted to focus solely on 1L Osimertinib failures
 - Testing Streamline patient enrollment based on current gold standard method (TBx FISH)
 - Increasing METamp prevalence Some publications suggest that METamp resistance post-Osimertinib could be ~25%¹
- Estimated primary completion date: November 2022

Tepotinib + Erbitux® (Cetuximab)
Addressing a significant medical need in 2L

metastatic colorectal cancer (mCRC)

- Opportunity for Tepotinib to address an unmet need in CRC together with Erbitux®
- Estimated primary completion date: March 2023

A solid foundation - Encouraging INSIGHT 1 data (18-months follow-up presented at WCLC 2019)²

Endpoint	Tepotinib + gefitinib	Chemotherapy
Primary - PFS (HR 0.13 [90% CI 0.04, 0.43])	16.6 m	4.2 m
Secondary - ORR (OR 2.67 [90% CI 0.37, 19.56])	66.7%	42.9%
Secondary - OS (HR 0.09 [CI 0.01, 0.54])	37.3 m	13.1 m



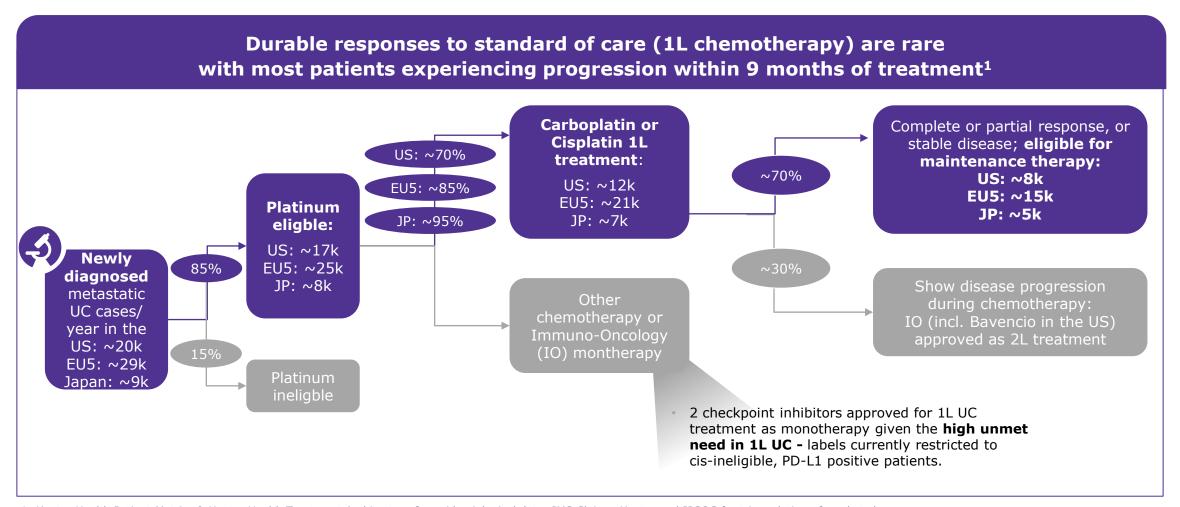
Safety: generally well-tolerated, most adverse events mild to moderate



1: Piotrowska et al., "Landscape of Acquired Resistance to Osimertinib in EGFR -Mutant NSCLC and Clinical Validation of Combined EGFR and RET Inhibition with Osimertinib and BLU-667 for Acquired RET Fusion", AACR Cancer Discovery 2018; 2: Wu et al., "Long term outcomes to tepotinib plus gefitinib in patients with EGFR mutant NSCLC and MET dysregulation: 18 month follow up", presented at WCLC 2019; Acronyms: FISH = Fluorescence in situ hybridization; TBx = Tissue Biopsy

Bavencio® (Avelumab) - Urothelial Carcinoma (UC 1L)

UC 1L maintenance treatment achieving transformative OS benefit (31% reduction in risk of death, 7 months increase in median overall survival)





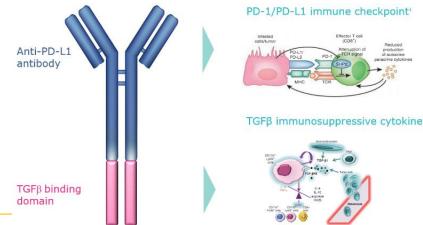


Bintrafusp alfa¹

An innovative first-in-class bifunctional fusion protein discovered in-house leading the TGF- β immuno-oncology field



- Innovative first-in-class bifunctional fusion protein designed to simultaneously target two immune suppressive pathways (blocking PD-L1 and reducing TGF-β signaling)
- Demonstrated superior anti-tumor activity in pre-clinical study compared to anti-PD-L1 alone, and anti-PD-L1 and TGF-β given in combination as separate agents
- Great excitement in IO community about M7824 uniquely addressing TGF-ß biology widely accepted as key resistance factor for anti-PDx therapies





- Tested in **14 Phase Ib expansion cohorts** across >700 patients in more than 10 tumor types
- Shown clinical anti-tumor activity across multiple hard-to-treat cancers including advanced NSCLC, biliary tract cancer, HPV-associated cancers, and gastric cancer
- PhII study M7824 monotherapy versus pembrolizumab 1L, advanced NSCLC high PD-L1-tumor expressers started in October 2018



 Multiple high priority immuno-oncology clinical development studies ongoing or expected to commence shortly, including studies in non-small cell lung and biliary tract cancers with registrational intent and most recently advanced, unresectable cervical cancer

¹proposed International Nonproprietary Name (INN)

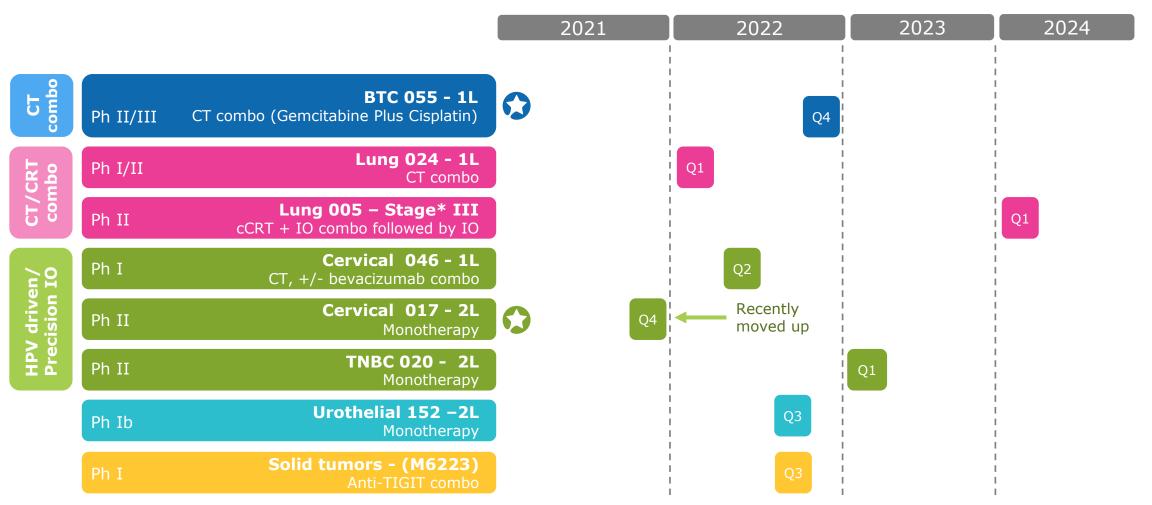




Bintrafusp alfa

INTR@PID Program: Upcoming Readouts





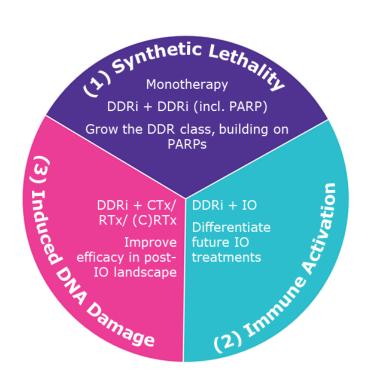
Acronyms: BTC = Biliary Tract Cancer; CT = Chemotherapy; EMT = Epithelial-mesenchymal transition; HPV = Human papillomavirus; NSCLC = Non-small Cell Lung Cancer; RT = Radiation therapy; TNBC = Triple-Negative Breast Cancer; * unresectable; All clinical timelines are event-driven and may be subject to change

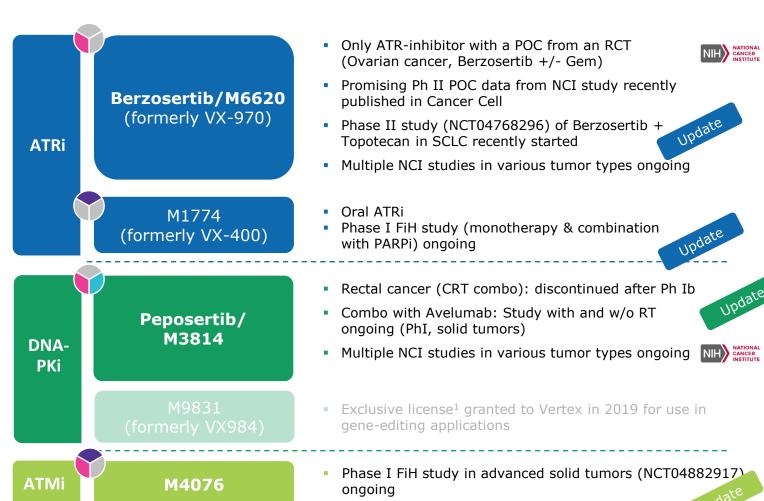


DNA Damage Response (DDR)

Leading DDR portfolio with a broad clinical program







1: incl. upfront payment + milestone/royalties on future sales; Acronyms: ATMi = Ataxia telangiectasia-mutated; ATRi = Ataxia telangiectasia and Rad3-related inhibitors; CRT = Chemoradiotherapy; DDR = DNA Damage Response; DNA-PKi = DNA-dependent Protein Kinase Inhibitor; FiH = First in Human; PARP = poly(ADP-ribose) polymerase inhibitor; POC = Proof of concept; RCT = Randomized Controlled Trial: RT = Radiation Therapy

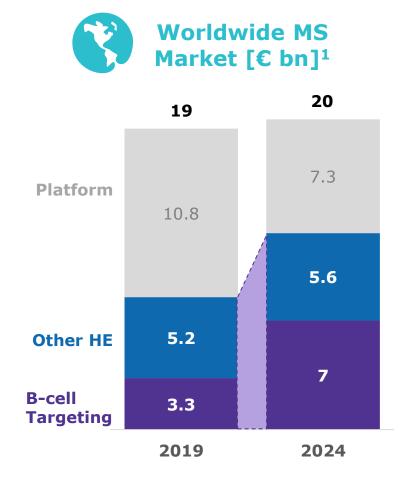


We pioneered BTKi development for MS with Evobrutinib

Potential to have 3 complementary MS branded products by 2025

Unmet need in Multiple Sclerosis (MS) – Need for new mechanisms to control disease

- ~50% of patients with Relapsing MS (RMS) continue to have ongoing disease activity over 2 years even when treated with the most effective agents
- No therapy with impact on progression mediated by CNS myeloid cells
- Systemic side effects of therapies limit patient acceptance and compliance
- All approved higher efficacy therapies associated with elevated risk of infection



B-Cell Targeting +
High-Efficacy (HE) Orals
represent >60% of MS sales

BTKi is a novel class of non-

depleting therapies selectively targeting both B-cells and innate immune cells including disease progression-relevant microglia

Merck KGaA, Darmstadt, Germany was the first to conduct a full

Phase II dose-ranging study in MS with Evobrutinib, a highly selective covalent BTKi²

Merck KGaA, Darmstadt,
Germany is a growing MS player
and could have 3 complementary
branded products by 2025 –
Mavenclad®, Rebif®, Evobrutinib

Platform agents – interferons, copaxone, DMFs and Teriflunomide; Other HE (high-efficacy) - cladribine, S1Ps, alemtuzumab; B-cell Targeting – ocrelizumab, ofatumumab, ublituximab. Includes branded products, generics and biosimilars; 1: Merck KGaA, Darmstadt, Germany internal estimates; 2: Montalban et al. NEJM 2019; 380:2406-2417; Acronyms: BTKi = Bruton's tyrosine kinase inhibitor

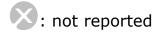
Evobrutinib stands out amongst BTK inhibitors under development

Uniquely positioned both in terms of clinical evidence and mode of action

		Fenebrutinib##	Tolebrutinib**	Evobrutinib
Ø	Long-term* efficacy on relapses			(1)
lence	Long-term* safety	×	×	(1)
l Eviden	Convenience (oral)	⊘ BID	₹ QD	♥ BID
Clinical	Exposure in CSF		(2, ##) in HV	(3) in MS
	Biomarker of inflammation and progression in MS patients (sNfL)			(3)
ical	BTK occupancy in the CNS	×	(4)	(5)
Preclinical data	Efficacy in progressive EAE model and reduction of leptomeningeal inflammation#			(6-8)



Phase III studies: Recruitment on track → Target data in-house in Q4 2023 and potential filing shortly after

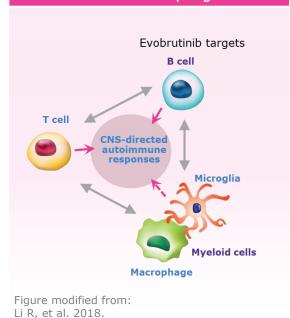




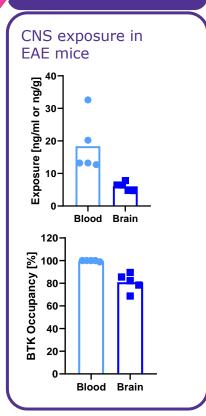
Evobrutinib targets inflammation and progression

Dual mechanism of action offers an innovative oral approach to MS therapy

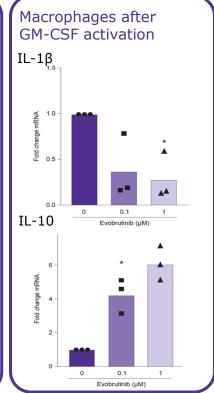
MOA on B-cells and macrophage/microglia cells is an innovative oral approach to MS therapy with potential to impact compartmentalized inflammation and progression



Evobrutinib is a CNS penetrant BTKi²

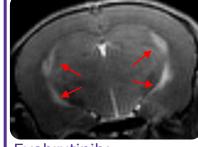


Evobrutinib shifts macrophage phenotypes from M1 pro- to M2 anti-inflammatory ³

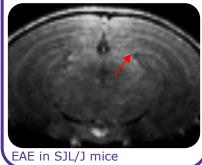


Evobrutinib reduces CNS B cells and leptomeningeal inflammation⁴





Evobrutinib:



- Important role of macrophage/ microglia cells in addition to B and T cells in both periphery and CNS¹
- Evobrutinib reaches brain to potentially impact compartmentalized inflammation²
- Effects may be mediated through effects on CNSresident and CNSmigrating cells^{3,4}



^{1.} Li et al. Nat Immunol 2018; 2. Adapted from Boschert U et al. ECTRIMS-ACTRIMS 2017; 3. Alankus YB et al. ECTRIMS 2018; 4. Sol Kim ECTRIMS 2020

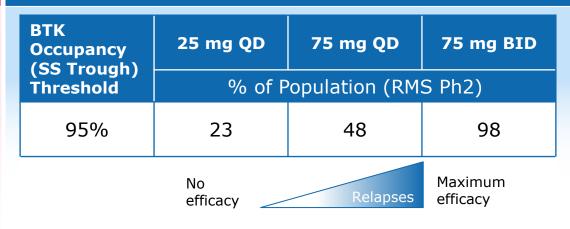
Evobrutinib is optimally dosed to offer best-in-class BTK inhibition Optimized dose selection & targeted covalent binding results in sustained BTK inhibition that is necessary for robust efficacy

Sustained BTK inhibition in CNS-resident and CNS-migrating cells resulting from covalent binding MOA and BID dosing can be critical to achieve best-in-disease efficacy¹

Targeted covalent binding leads to highly specific continuous target engagement²⁻⁶

Targeted covalent binding

BID dosing enables critical >95% BTK inhibition throughout the day in the majority of patients⁷



	, ,	20.000.7.0,	action in CNS
втк	Evobrutinib		
Covalent binding			
	Tolebrutinib ple other		
Reversible binding			
ВТК	Fenebrutinib		

Efficacy at 48 weeks ⁸					
75 mg QD Evobrutinib Fasted: ARR = 0.25	75 mg BID Evobrutinib Fasted: ARR = 0.11				

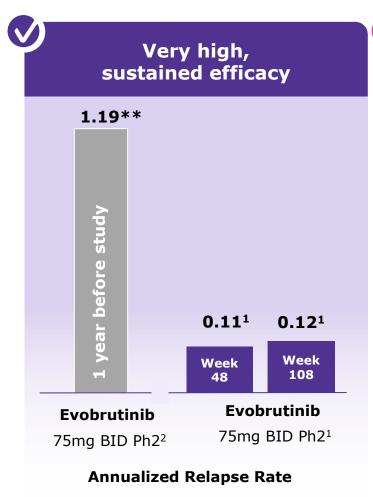


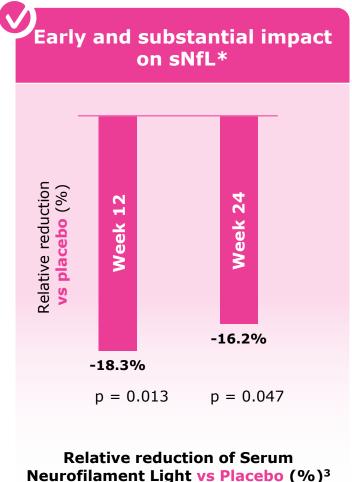
Selectivity* Sustained



Evobrutinib holds unmatched Long-Term Data among BTKi class in MS

Best-in-disease efficacy & favorable safety over 2 years in largest Phase II study in MS





- Evobrutinib is the only BTKi to have demonstrated very high, sustained efficacy and favorable safety in the largest Phase II study in MS (n=267), with an ARR of confirmed relapses of 0.12 up to 108 weeks¹
- Evobrutinib impacts sNfL levels, a biomarker of neuronal damage, reflecting disease activity and drug response in patients with MS³, starting at 12 weeks and maintained through 24 weeks²
- Evobrutinib is highly selective resulting in targeted kinase inhibition, and its safety data in >1200 patients over 2+ years supports the potential for an optimal longterm safety profile²

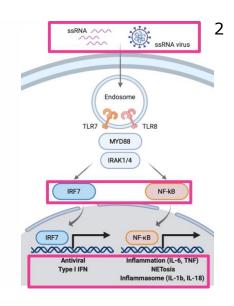


M5049 (TLR7/8 antagonist)

TLR7/8 are drivers of SLE pathology and possibly of COVID-19

Mechanism of Action¹

- M5049 (discovered in-house) is a potentially first-in class small molecule that blocks activation of Toll-like receptors TLR7 and TLR8, two innate immune sensors that detect single-stranded (ss) RNA from viruses such as SARS-COV-2, the virus responsible for COVID-19, and inflammatory self-RNAs in the context of autoimmunity
- Activation of TLR7/8 leads to immune cell activation and inflammation, which when not properly controlled can cause severe immunopathology



Results from Phase I study in healthy volunteers (NCT03676322)¹

- Well-tolerated over the dosing interval, no significant or dose-limiting adverse event
- Pharmacokinetic parameters linear and dose-proportional from 1 to 200 mg
- Exposure-dependent inhibition of ex vivo-stimulated IL-6 secretion observed, with maximum inhibition achieved at 200 mg



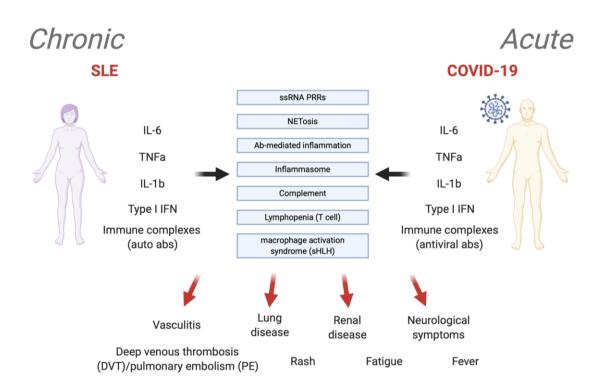
Preliminary Phase I data warrant further investigation as a potential treatment for autoimmune diseases including SLE

1: Port et al., A PHASE I, FIRST-IN-HUMAN STUDY TO ASSESS THE SAFETY, PHARMACOKINETICS AND PHARMACODYNAMICS OF SINGLE AND MULTIPLE ASCENDING DOSES OF M5049, A DUAL ANTAGONIST OF TLR7/8, IN HEALTHY SUBJECTS, Lupus Science & Medicine 2020;7(Suppl 1):A1-A131, conference cancelled due to COVID-19; 2 Adapted from ImmunoHorizons July 1, 2018 Dowling, D; Acronyms: SLE = Systemic lupus erythematosus: TLR = Toll-like receptors

M5049 (TLR7/8 antagonist)

Similarities between SLE and COVID-19

Similarities between SLE and COVID-19¹



Phase II study started in July 2020

Rational:

- Investigate if M5049 intervention at critical point in course of COVID-19 disease may prevent or ameliorate hyperinflammatory response in patients with COVID-19 pneumonia and prevent progression to 'cytokine storm'
- Successful intervention with investigational drug may reduce life-threatening complications of COVID-19, including severe respiratory symptoms often necessitating further interventions such as mechanical ventilation

Design:

- Phase II randomized, controlled clinical study
- Commenced in July 2020

Results:



Dependent on recruitment and COVID-19 infection rates

Data read-out expected in Q3/Q4 2021

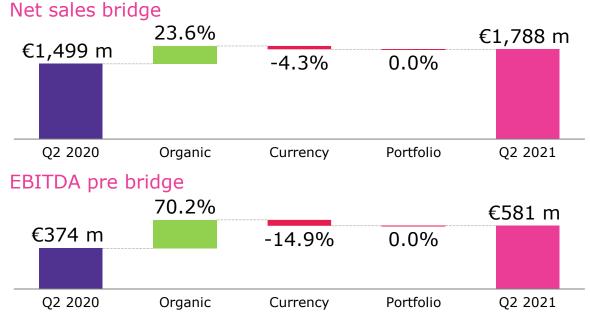


^{1:} Illustration created in-house; Acronyms: SLE = Systemic lupus erythematosus

Healthcare Q2: Strong Fertility recovery & Bavencio[®] performance; Mavenclad[®] returns to sequential growth as dynamic market picks up

Healthcare P&L

[€m]	IFRS		Pre	
	Q2 2020	Q2 2021	Q2 2020	Q2 2021
Net sales	1,499	1,788	1,499	1,788
M&S*	-409	-391	-401	-389
Admin	-81	-78	-79	-76
R&D	-366	-415	-366	-414
EBIT	269	501	284	512
EBITDA	359	572	-	-
EBITDA pre	374	581	374	581
(in % of net sales)	24.9%	32.5%	24.9%	32.5%



Comments

- Mavenclad® grows +102 % organically to €157 m, amid first signs of recovery of dynamic market and low comps; Rebif® declines -9%
- Oncology up +49%; Bavencio[®] grows +206% fueled by UC 1L launch in key markets; Erbitux[®] up +36% supported by Eli Lilly supply agreement
- Base business up +15%, driven primarily by strong Fertility recovery (+88% org.); CM&E +1% org. Glucophage VBP¹ impact compensated for by Endocrinology
- Lower absolute M&S vs. Q2 2020 with higher level of face-to-face activities amid progressing adaptation to pandemic situation
- Higher absolute R&D driven largely by ramp up of Evobrutinib and Xevinapant development activities; lower R&D as % of sales
- EBITDA pre and margin increasing with operating leverage, further supported by temporary Eli Lilly supply agreement in the U.S.



¹ Volume Based Procurement

^{*} Marketing and selling expenses

Totals may not add up due to rounding

Life science

Life Science

Capitalizing on three key life science trends







Single Use / End to End

Opened Wuxi site in 2018, and expanded Danvers facility

Viral Vectors

Expanded Carlsbad viral vector manufacturing site in 2016; further doubling of capacity planned for 2021

Antibody Drug Conjugates (ADC)

Launched ADC Express[™] for the rapid production of ADCs #1 eCommerce site in Life Science¹

- > 90% of Millipore products on eCommerce platform
- x2 net sales growth of eCommerce vs.

Manufacturing/Distribution
Nantong, Wuxi Single use

Commercial expansion
Tier 2 cities

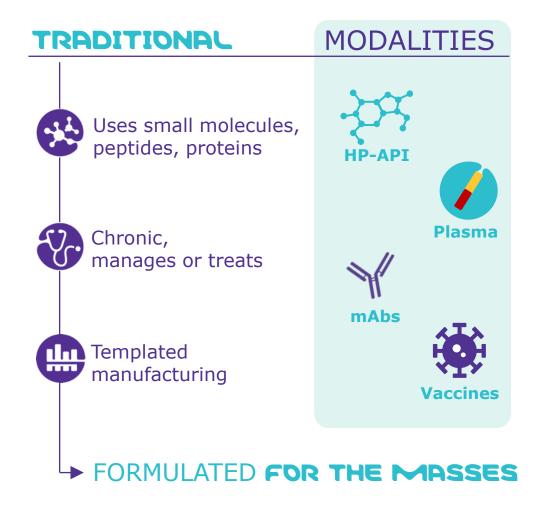
eCommerce partnership

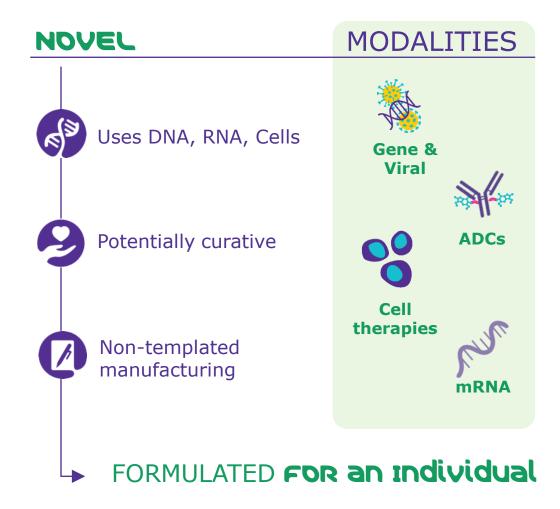




Process Solutions: Therapies are evolving from treatments to cures

Advancing traditional is critical as novel modalities develop



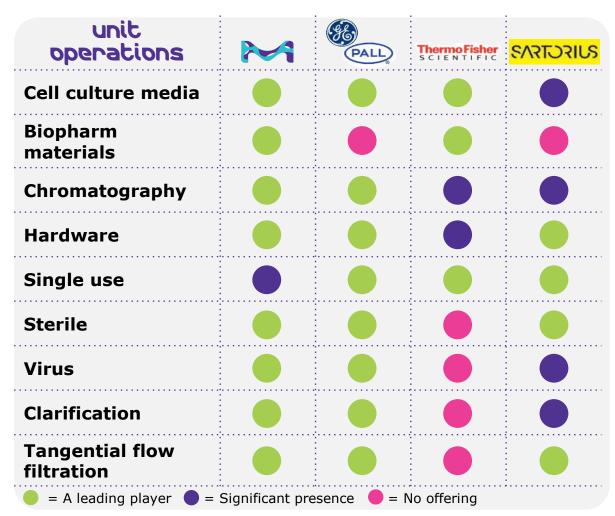


Acronyms: HP-API = highly potent active pharmaceutical ingredient; mAbs = monoclonal antibodies; DNA = deoxyribonucleic acid; (m)RNA = (messenger) ribonucleic acid; ADC = antibody drug conjugate



Process Solutions

COVID demands align with our strengths but increase supply chain pressure



COVID-19 Outlook

Type

Implications



65 programs

Bind and block virus from entering cells

- Universal templates
- A leading position for 8 out of 9 unit ops



Vaccine

199 programs
Protective **immune response**

- Multiple templates
- Leveraging Single Use



Nucleic Acid

43 programs
Leveraging human factory

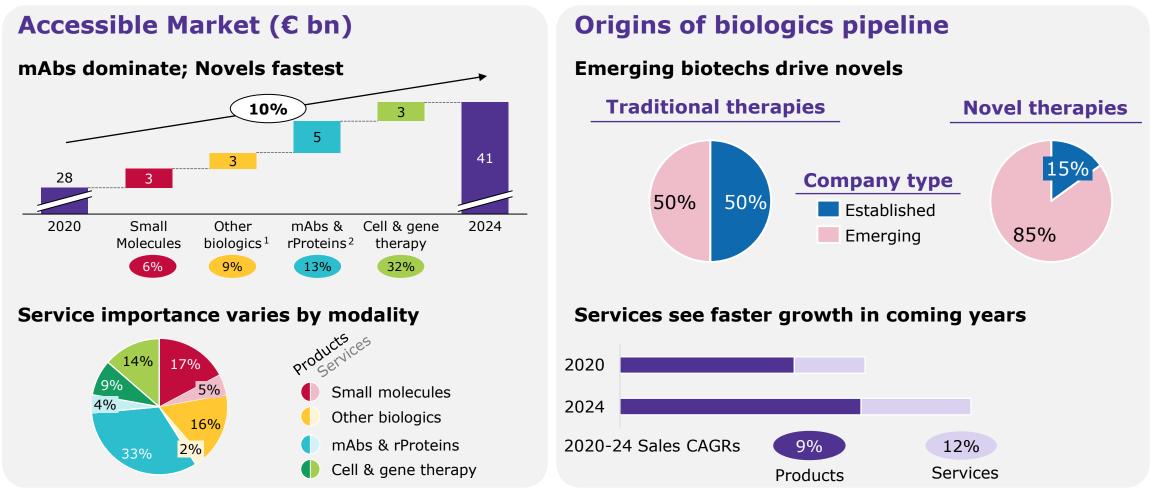
- Emerging manufacturing processes
- Lipids are critical

Sources: press releases, company reports, and internal assessments



Process Solutions

Opportunities in services to accelerate double-digit growth



Sources: Evaluate Pharma, internal market models, CSR sales data; ¹ Other biologics include plasma, vaccines, insulin, microbial and non-mAb biosimilars; ² mAbs include ADCs here; Additional acronym: rProteins = recombinant proteins



Process Solutions: Strategic direction

Innovate and invest today to continue above market growth in the future



Modality





ALL





NOVEL



Revenue driver



Near term



Mid term



Long term



Offering



Products



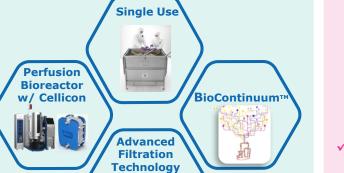
Services



Products



Portfolio





- Contract development & manufacturing organization
- ✓ Contract testing organization
- ✓ Product characterization



- Templated viral manufacturing consumables and reagents
- ✓ Autologous manufacturing system



Next-generation bioprocessing on the cards



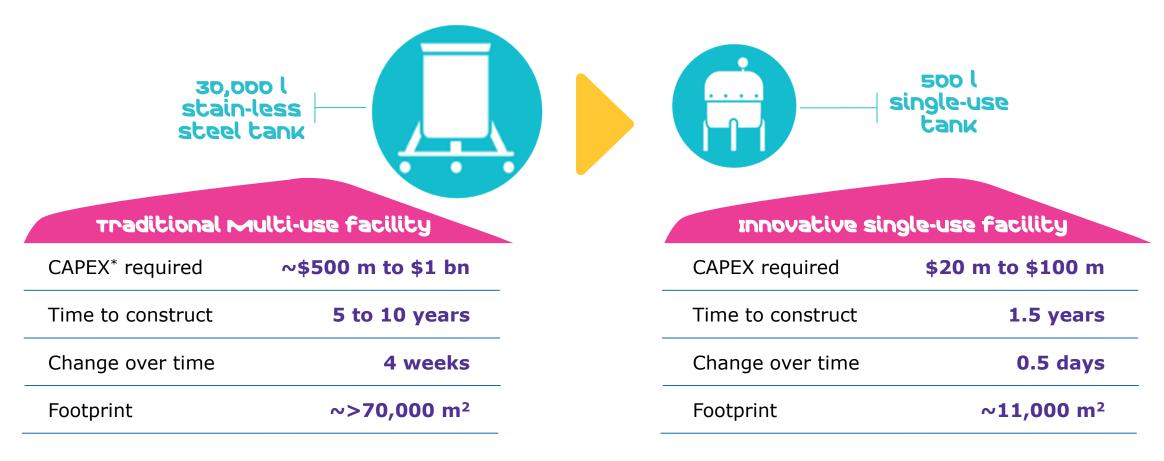


Continuous bioprocessing will ...

- be an evolution in mAb bioprocessing
- take time to establish
- leverage the present
- lead to hybrid solutions

Process Solutions

Our single-use technologies drive flexibility in modern bioprocessing





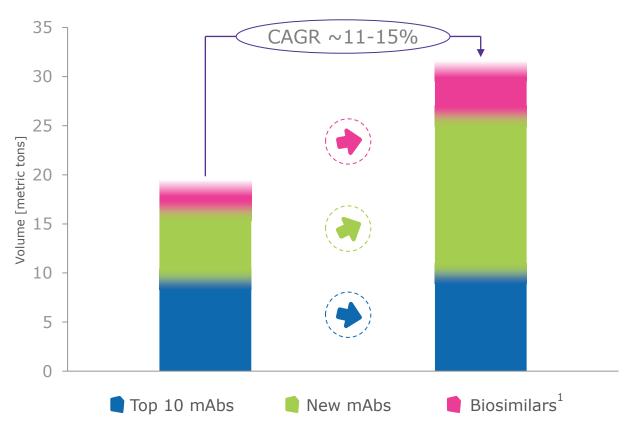
Strong demand for single-use technologies and Process Solutions' broad offering was and will remain a key source of growth for Life Science



Process Solutions

mAbs market democratization will drive diversification, change & variability

mAb volume projections 2020 to 2024



¹Biosimilars scaling factor = 2.8 based off internal estimates and McKinsey analysis; Source: company estimate based on industry forecasts, EvaluatePharma; mAbs = Monoclonal antibodies

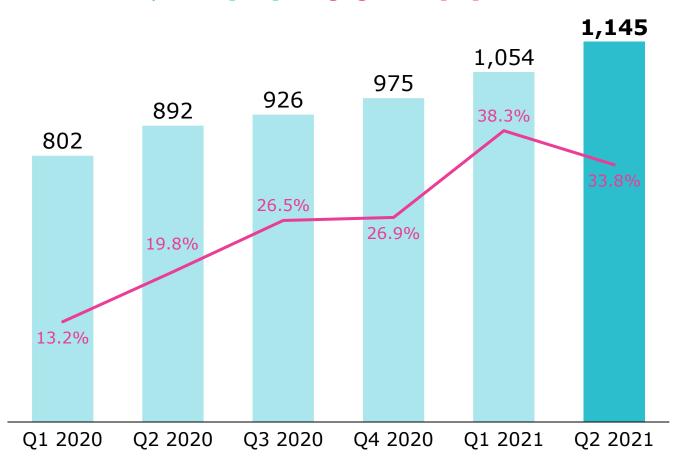
market development

- Overall mAbs market will grow ~11-15% CAGR
- Top 10 originator mAbs represent
 50% of market volume today and will decline to ~30% in 2024
- Biosimilars will gain share



BIG 3 - Process Solutions: Continued strong double-digit growth, moderating as expected against rising comps

Sales development [€m] - org. growth [%]

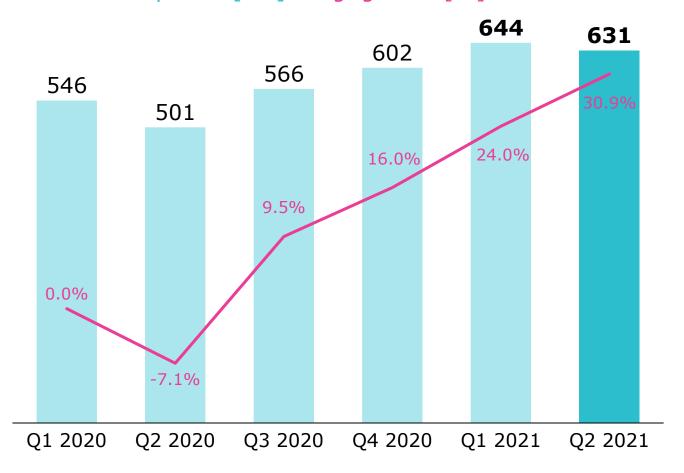


- Continued double-digit growth in the core business paired with rising COVID-19 contributions (mainly vaccine related)
- BioP as main growth driver, formulation growing fastest, services also strong
- Growth starting to moderate as expected amid rising comps (Q2 2020 with initial COVID-19 contributions)
- Sequentially higher sales as output increases on the back of ongoing productivity gains and successful capacity ramp-up
- Order intake growth remains above 60%, reflecting strong demand



Research Solutions: Record organic growth amid ongoing business recovery and soft comps

Sales development [€m] - org. growth [%]

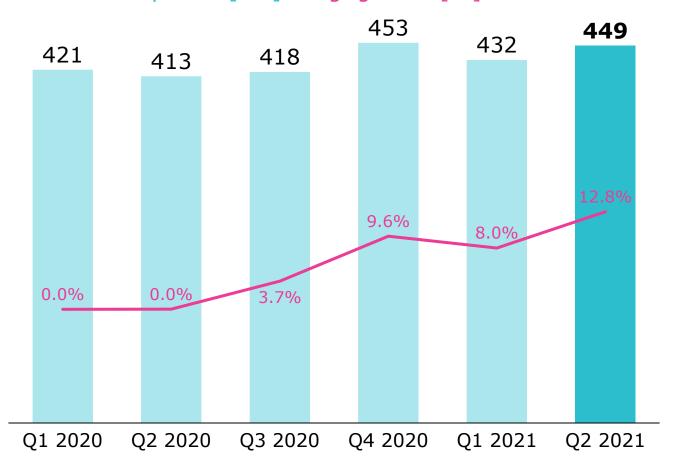


- Strong double-digit growth, accelerating further on soft comps (Q2-20 with biggest impact from lockdowns)
- Ongoing core business recovery and catchup amid rising lab activity
- Diagnostics related COVID-19 tailwinds continue albeit slowing as expected
- North America as fastest growing region, followed by Europe and APAC (given difference in comps / phasing of lockdowns last year)
- Strong rebound in academia and healthy growth in pharma



Applied Solutions: Growth accelerating as recovery is gaining traction

Sales development [€m] - org. growth [%]



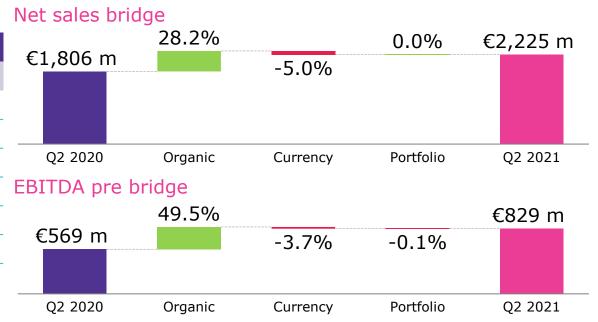
- Growth accelerating to double-digits as recovery continues amid still easy comps (H1-20 with flat growth due to lockdowns)
- Core business as main driver with broadbased performance across business lines
- COVID-19-related sales are negligible
- North America as fastest growing region, followed by Europe and APAC (given difference in comps / phasing of lockdowns last year)
- Strong rebound in Academia and ongoing recovery in Industrial



Life Science Q2: Strong core business and COVID-19 demand fuel record growth, mainly driven by Process and Research Solutions

Life Science P&L

[€m]	IFRS		Pre	
	Q2 2020	Q2 2021	Q2 2020	Q2 2021
Net sales	1,806	2,225	1,806	2,225
M&S [*]	-488	-505	-488	-505
Admin	-100	-92	-88	-84
R&D	-75	-87	-75	-87
EBIT	386	644	370	638
EBITDA	584	835	-	-
EBITDA pre	569	829	569	829
(in % of net sales)	31.5%	37.3%	31.5%	37.3%



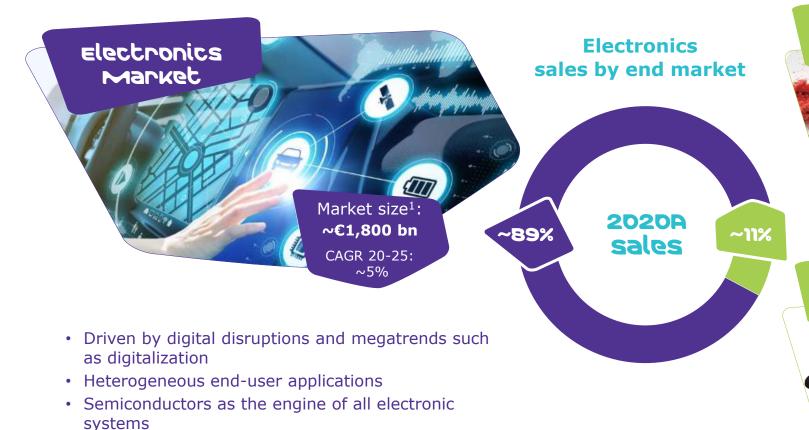
Comments

- Process Solutions: grows +34% organically, supported primarily by bioprocessing demand for COVID-19 projects; comparable base now starts including COVID-19 business
- Research Solutions: grows +31% organically against Q2 2020 COVID-19 dip, driven by recovery in base business and COVID-19 opportunities, mainly in diagnostics and pharma
- Applied Solutions: grows (+13% org.) against softest quarter of 2020

- Declining M&S in % of sales from 27% to 23% due to strong top line leverage, slightly higher in absolute terms
- Higher R&D in absolute terms with continued focused investments in high growth & emerging segments
- Business performance, operational leverage & favorable mix continue to drive strong EBITDA pre and margin expansion



Targets attractive markets – especially in the electronics space



Automotive Market

- Driven by world GDP growth
- Increasing demand in emerging markets

Market size: ~€2,000 bn

cosmetics market

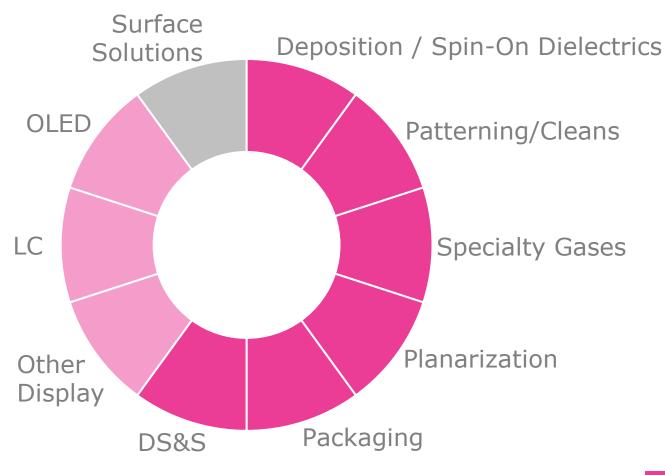
- Driven by world GDP growth
- Rising living standards and higher disposable income

Market size: ~€400 bn



¹Prismark 2021

Expected mid-term portfolio split



Mid-term the Electronics portfolio will consist of ~10 equally sized businesses

>60% of them serving chip makers

Semiconductor Solutions

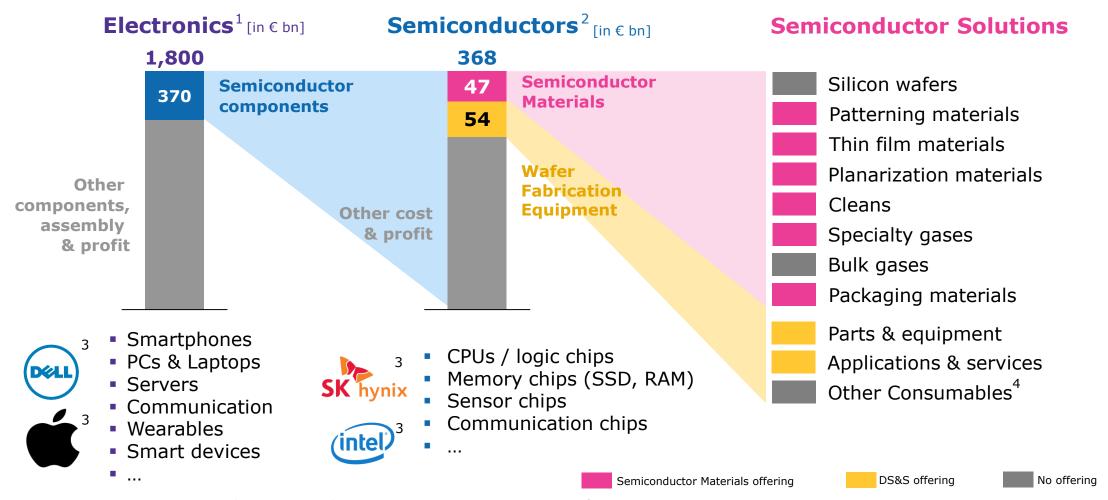


Display Solutions



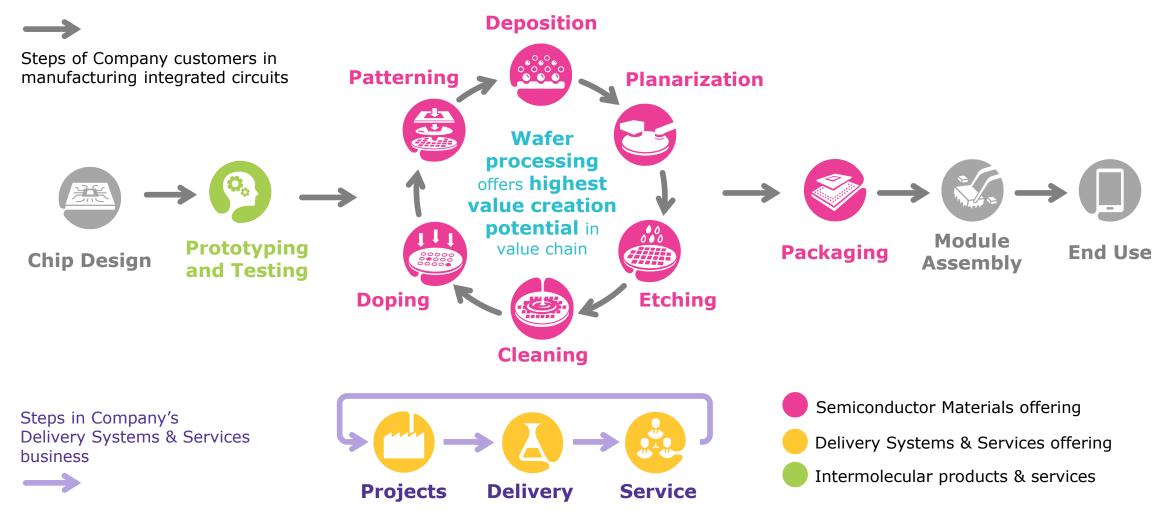
INDICATIVE Chart

Semiconductor Solutions - integrated materials player, well positioned to serve the need of customers in semiconductor fabrication



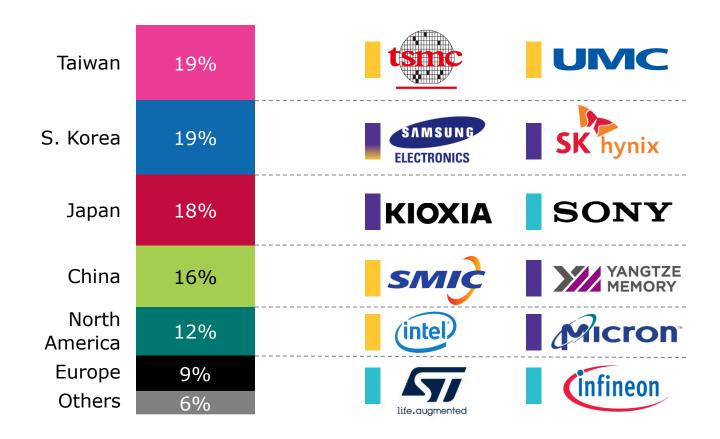
llustrative Industry P&Ls based on Sources: ¹Prismark 2021, ²Prismark 2021 & WSTS/SIA & SEMI Q1 2020; ³Representative player in the industry, non-exclusive list, not based on any underlying criteria; ⁴e.g. Filters, Pads, etc.; CPU = Central Processing Unit; RAM = Random Access Memory; SSD = Solid State Disk; CMOS = Complementary metal-oxide semiconductor

Unique comprehensive products and services portfolio offers end-to-end solutions, well-placed in high growth segments



Beyond a comprehensive portfolio Semiconductor Solutions also has an industry spanning customer base, supplying various end markets

2019 wafer capacity by region Selected customers per region 2



Semiconductor Solutions has

customers

supplying all top 10 chip makers and virtually all of the top 100³



Focus on logic chips

Focus on memory chips

Focus on other chips

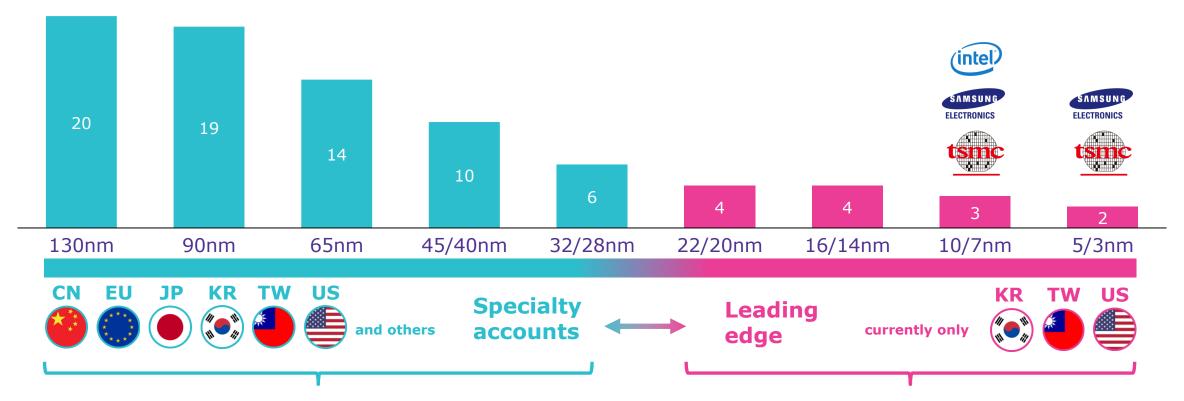
¹SEMI World Fab Forecast Q3 2020 - Dec 2019 capacity,

²Representative, non-exclusive list, not based on any underlying criteria

³Based on H1 2020 Sales

Only 3 companies are currently running volume production ≤10nm These companies have the largest market shares across all nodes

Number of companies currently running volume production per logic chip node¹



Primary focus on operation efficiency

Primary focus on innovation efficiency

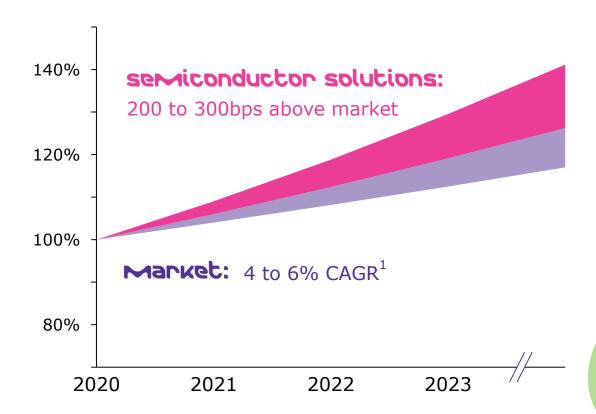
¹Source: Wikichip.org and own data; volume production as of Sep 2020; countries are listed in alphabetical order

Semiconductor Materials

Set to outgrow highly attractive semiconductor materials market

Semiconductor Solutions sales guidance vs. market

[Indexed 2020 = 100%]



¹Source: Jan 2020 IC Insights 2018-2024 CAGR for wafer starts in million units

Market

- Technological trends inevitably drive exponential data growth
- More data requires more chips and higher complexity of chips
- Rising materials value added per wafer

semiconductor solutions

- Comprehensive offering focusing on attractive materials categories
- Integration topline synergies
- Critical mass and deep customer centricity
- Better customer understanding: know-how exchange and collaboration across DS&S and Semiconductor Materials
- Cutting-edge innovation and R&D capabilities



Technology trends inevitably drive exponential data growth... ...more chips needed to generate, transfer, process & store data

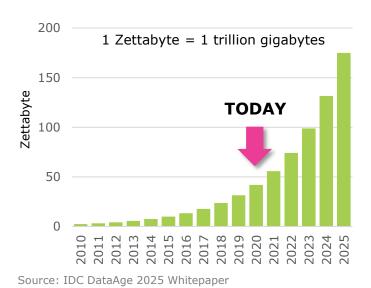
Data created worldwide

is growing +30% annually

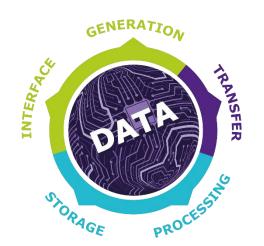
All segments of data application are affected by global data growth

Technology trends strongly impact relevance of data application segments

Size of global data sphere









Technology market growth - examples

5G Technology¹

>122% CAGR

Artificial Intelligence²

>33% CAGR

IoT Sensors³

>24% CAGR

Data Center Services⁴

>13% CAGR

Autonomous Driving⁵

>18% CAGR

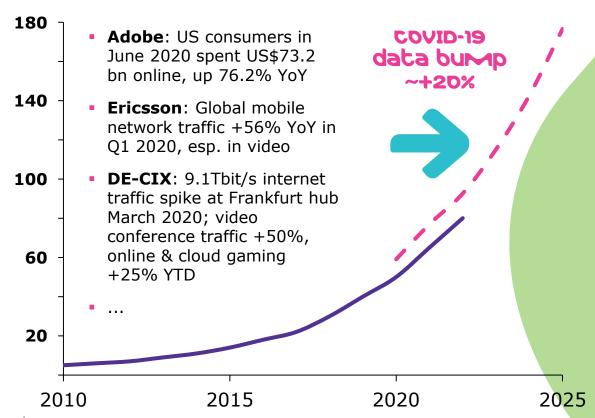
Semiconductor Solutions supports growth trend as part of "the company behind the companies, advancing digital living"

1) <u>alliedmarketresearch.com</u>, Prismark 2020, CAGR 2021-2026; 2) <u>fortunebusinessinsights.com</u>, <u>post-gazette.com</u>, CAGR 2018-2026; 3) <u>mordorintelligence.com</u>, <u>computerweekly.com</u>, CAGR 2020-2025; 4) <u>mordorintelligence.com</u>, Prismark 2020; CAGR 2020-2025; 5G = 5th-generation cellular wireless; IoT = Internet of Things 5) <u>mordorintelligence.com</u>, autonomous car market value CAGR 2020-2025

Semiconductor Solutions

COVID-19 has vaulted the "digital transformation" by ~5 years¹

Expected COVID-19 impact on global datasphere² [zetabytes]



¹Source: McKinsey May 2020 "The COVID-19 recovery will be digital: A plan for the first 90 days"; ²Source: Seagate, IDC April 2020, Merck KGaA, Darmstadt, Germany

- Work-from-home/stay-at-home economy
 - Significant increase in video conferences, online shopping, online gaming, streaming
- Change in consumers' and enterprises' digital behavior expected to be long-lasting
- Need for more, faster & more reliable data processing, storage and bandwidth
 - Acceleration of semiconductor demand

semiconductor solutions stands ready to support increased demand



covid-19 impact on data growth expected to be positive

Expanding the limits of how small you can go

Pattern collapse



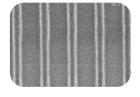


AZ® rinse materials



As lines get narrower and closer together in advanced chip generation, lines tend to "stick" due to surface tension.

Lithography limitation



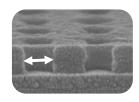


Directed self-assembly (DSA)



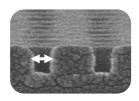
Block copolymer can generate small lines or contact holes by selfassembly. This allows miniaturization without expensive new equipment.

Wide features





AZ® shrink materials



Shrink materials "shrink" the gap between lines and, hence, allow the manufacture of narrower features otherwise not possible.

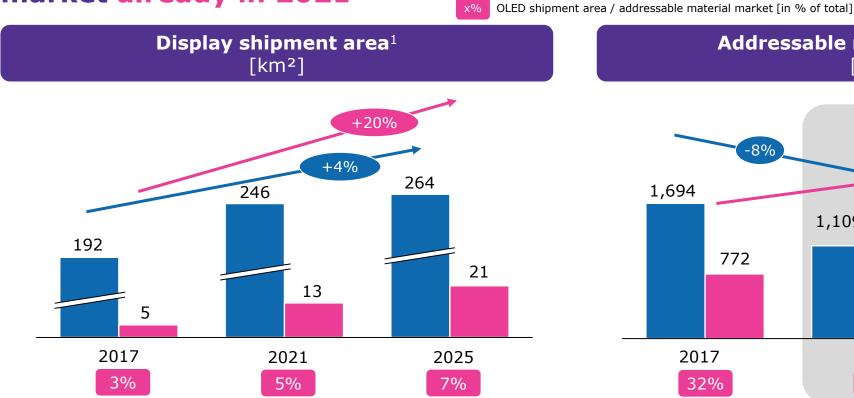


Company delivers highly innovative solutions for complex customer problems



Display Solutions - OLED material market to exceed LC material

market already in 2021

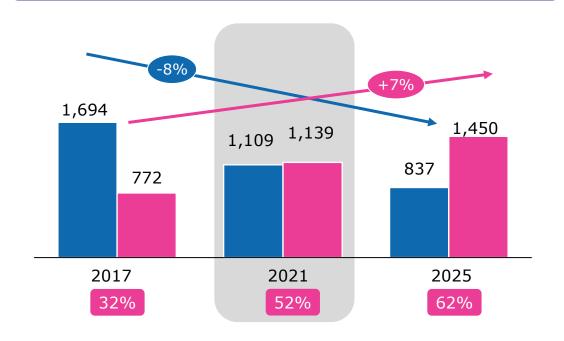




 OLED growing faster than LCD, but LCD to command 90+% area share for foreseeable future

Addressable material market² [€m]

Liquid Crystals



- Material value per OLED display higher than in LCD
- OLED material market to exceed LC material market by 2021, but market split between many more players



OLED

Electronics Q2: Strong performance in Semi, recovery in Surface, and stabilizing Display Solutions drive double-digit organic sales growth

Electronics P&L

[€m]	IFRS		Pr	·e
	Q2 2020	Q2 2021	Q2 2020	Q2 2021
Net sales	814	857	814	857
M&S [*]	-134	-137	-131	-136
Admin	-44	-30	-36	-28
R&D	-68	-67	-69	-66
EBIT	-30	118	101	129
EBITDA	219	252	-	-
EBITDA pre	238	258	238	258
(in % of net sales)	29.3%	30.1%	29.3%	30.1%



Comments

- Semiconductor Solutions: record quarter in terms of sales; 12% organic
 growth across all businesses with Semi Materials growing even faster
 than strong DS&S
- Display Solutions: down -1% organically as LC decline was nearly fully offset by growth in remaining portfolio primarily strong OLED
- Surface Solutions: delivers 41% organic growth over pandemic-impacted Q2 2020; visible recovery across all end markets

- M&S up 4%, largely driven by higher logistic costs, while admin and R&D are declining
- All P&L lines continue to reflect diligent cost management amid Bright Future transformation and Versum integration synergies
- EBITDA pre (+14% org.) continues to exceed sales growth, but burdened by -6% FX headwinds



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