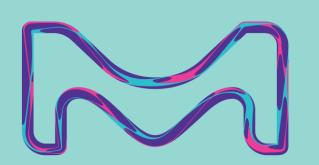
Bernstein 18th Annual Pan European strategic Decisions conference

Belén Garijo, CEO

September 22, 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

- **D1** Business overview
- **Q2** Transforming the company
- **Use 3** Healthcare Executing on the earnings phase
- Life Science Focusing on profitable growth
- **D5** Electronics Leveraging portfolio shift
- **Sustainability**
- **O7** Guidance & executive summary





Leading positions in fastest growing science & technology markets



Accelerated science & tech leadership driving growth



company innovation center parmstadt



2021 and beyond - poised for growth in a challenging environment



Acronym: VBP = volume based procurement

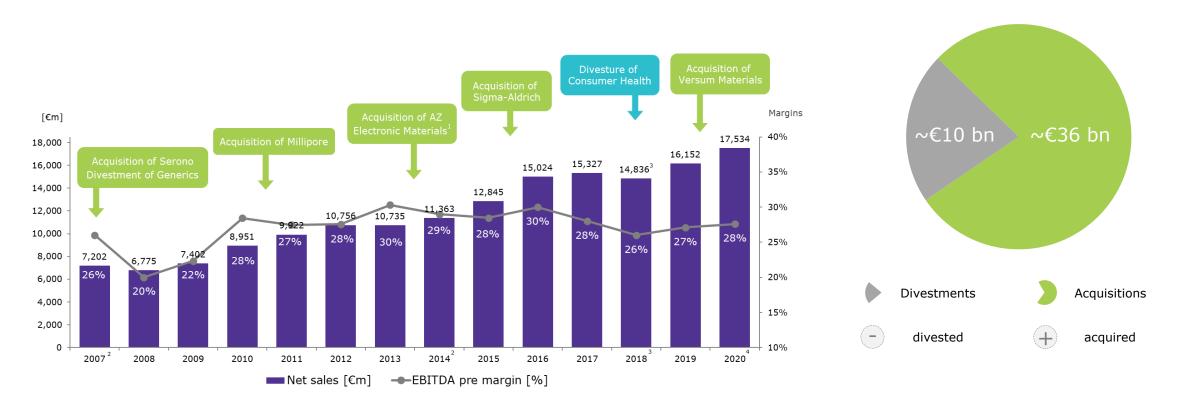




Added scale and strengthened attractiveness of portfolio

Growth trajectory

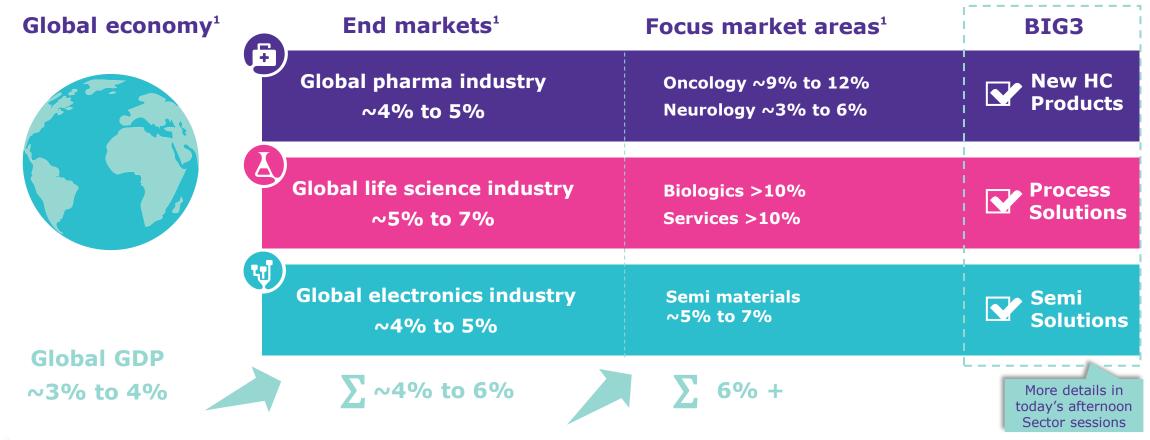
Transformation volume⁵



¹Included since 2 May 2014; ²2007 and 2014 EBITDA pre margin adjusted for comparability; ³2018 net sales reflect Consumer Health divesture (reduction of ~€1 bn net sales p.a.) ⁴2020 margin restated for €365 m patent litigation provision release; ⁵ 2007 - 2020



Well positioned to win in high-growth focus areas



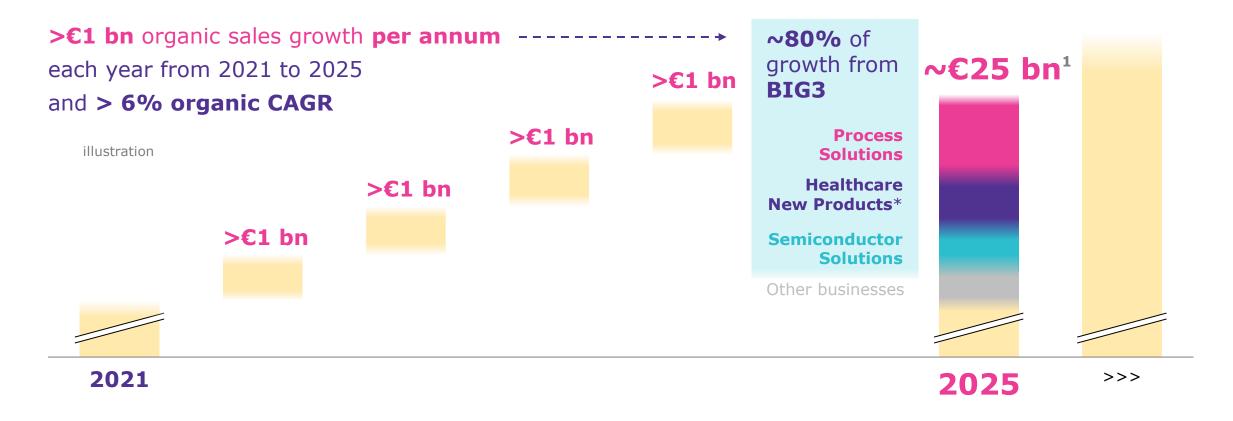


Uniquely positioned to address inevitable market trends
Diligent execution of focused & disciplined investments plan to fuel BIG3 growth



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

#25by25: delivering €5 bn+ organic additional sales by 2025





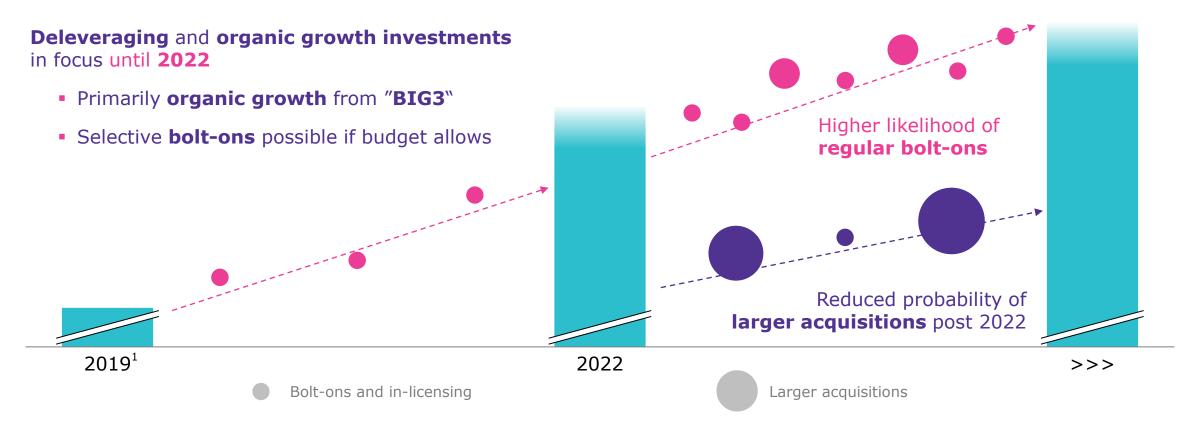
Clear commitment to efficient growth

1) including potential bolt-on M&A

*) Updated scope: New Products consist of recent launches (Bavencio[®], Mavenclad[®], Tepmetko[®] MET-exon 14) and risk-adjusted launches to come until 2025



Portfolio strategy – from transformation to evolution





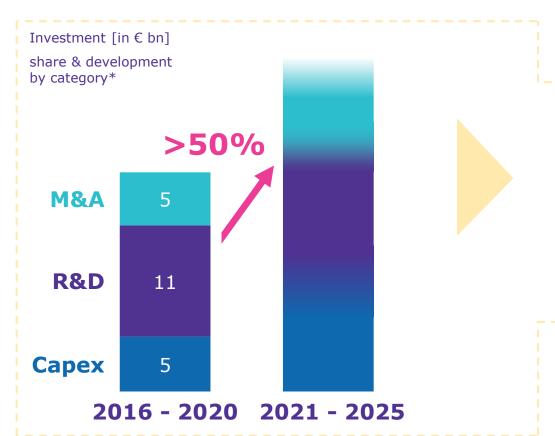
> €1bn organic sales growth per annum each year from 2021 to 2025

¹ 2019 Group sales of €16.2 bn

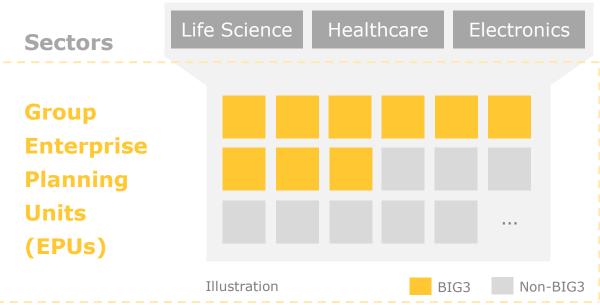


Committing to record investments - targeted @BIG3 growth engines

Leading positions in **growth markets** fuel significant **boost in investment capacity**



Targeted, **disciplined** & **focused** capital allocation across >20 enterprise planning units



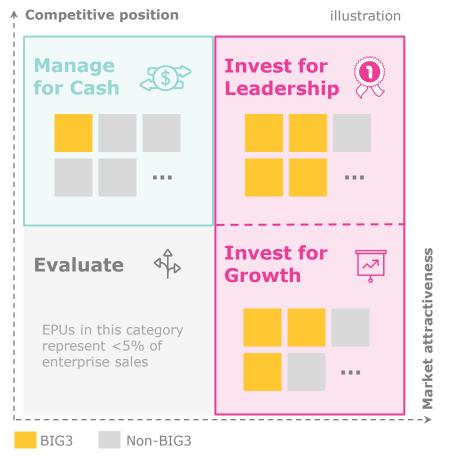
BIG3 consist of ~10 planning units with a capital allocation target of >70% between 2021 and 2025



^{* &}gt;50% statement primarily valid for R&D and CAPEX plan, future M&A purely illustrative as it is deal-dependent

Optimized capital allocation through distinct portfolio roles

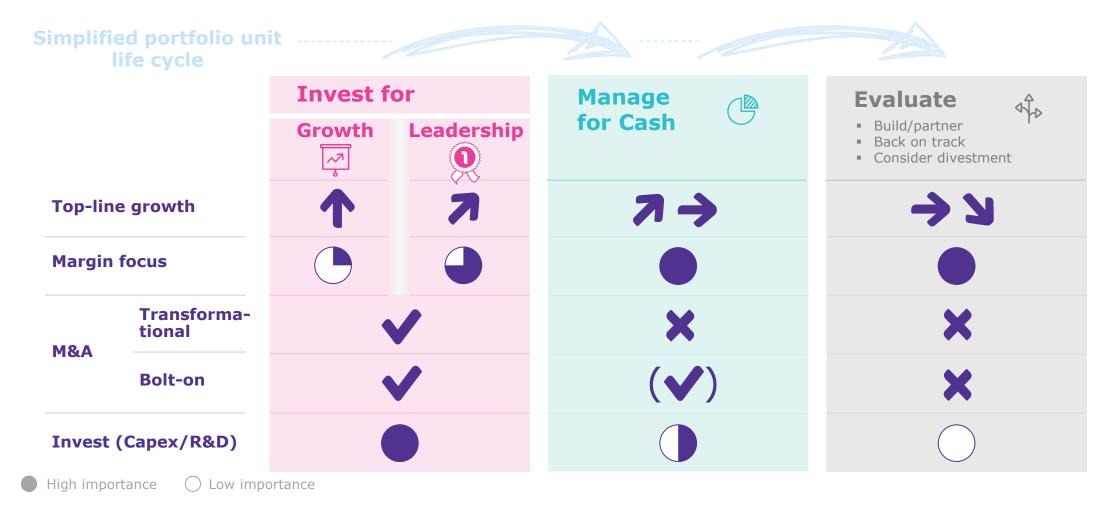
Roles determined by **thorough analysis** of **markets** and **competitive positions**



Investment focus on businesses with greatest strengths and attractive opportunities

- Segmentation in planning units allows right level of granularity in market & competitive analysis (vs. sector level or product level)
- Enterprise Portfolio Units (EPUs) with different roles drive balance between strong cash generation & targeted capital allocation
- Strong, well-positioned portfolio
- Enterprise setup allows boosted investment capacity and optionality that would be unattainable to 3 standalone businesses
- No need for further diversification (within or across sectors) or target sector ratios

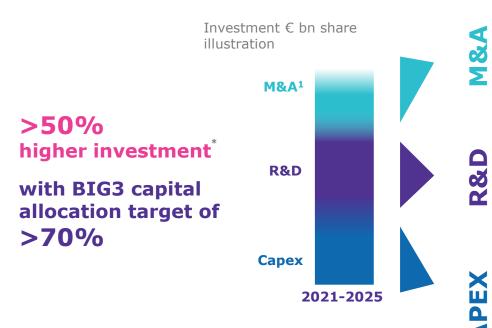
Portfolio roles provide clear guidance to resource allocation



Financial flexibility fueling investment capacity

Σ

Accelerated deleveraging to <2x net debt to EBITDA pre ratio enables unprecedented financial flexibility



Abundant growth opportunities make strategic capital **allocation** more important than ever

- Strong operating cash flow & increased debt capacity
- High single-digit € bn financial flexibility by end of 2022
- Continued higher likelihood of **bolt-on** vs. large transformational deals

Clear focus on **innovation** & further development of **pipeline productivity**

- Life Science: accelerating targeted R&D scale-up to capture trends & strengthen portfolio
- **Electronics: c**ontinuous alignment with customer technology roadmaps incl. new R&D labs
- **Healthcare:** driving "R&D productivity" to benchmark levels with focused leadership

More complex & multifaceted CAPEX decisions requiring discipline

Rich growth opportunities driven by demand step changes, e.g.

- Process Solutions capacity & network expansion, targeted scaling of high value CDMO/CTO services across traditional & novel modalities
- Semi capacity expansion in line with customer demand & smart localization
 - Annual capex guidance: from ~€1.5 bn in 2021 to ~€2 bn by 2023



^{* &}gt;50% statement primarily valid for R&D and CAPEX plan, future M&A purely illustrative as it is deal-dependent

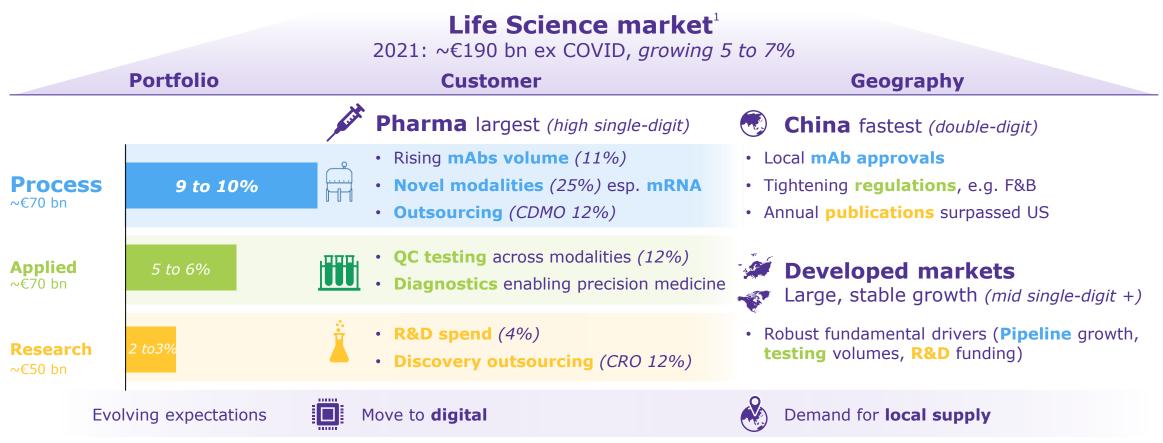
¹ M&A: Mergers and Acquisitions

Life science

Focusing on profitable growth

Life Science Market

Attractive secular drivers across segments create opportunities





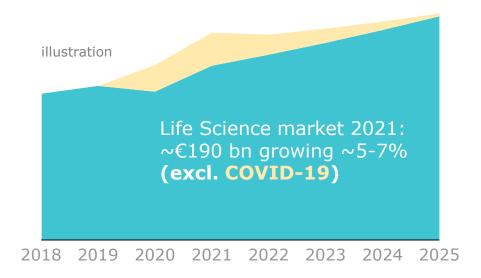
We continuously pursue key growth trends as a leading diversified life science player

¹ Company estimate based on industry forecast over 5-year horizon for the markets we operate in with all *growth rates* indicating 2021-25 CAGRs; Acronyms: mAbs = Monoclonal Antibodies; mRNA = Messenger Ribonucleic Acid; CRO = Contract Research Organization; CDMO = Contract Development Manufacturing Organization; F&B = Food & Beverage

Life Science

Driving growth and expanding leadership in Process Solutions

Robust market with diverse growth pockets, **elevated during pandemic**



- Attractive base market, on slightly higher growth trajectory post 2021
- Significant demand from COVID-19, expected to decline as pandemic recedes

Life Science will be fully equipped to capture growth opportunities with a **focus on Process Solutions**

- Major capacity and network expansion to fuel growth of key product portfolios, especially bioprocessing
- Targeted scaling of high value CDMO/CTO services across traditional and novel modalities
- Relevant and diversified portfolio in Research and Applied, not only for cash and resilience, more importantly for innovation and trend scouting
- Programmatic bolt-on M&A plus rising focus on China/APAC, innovation, and digital



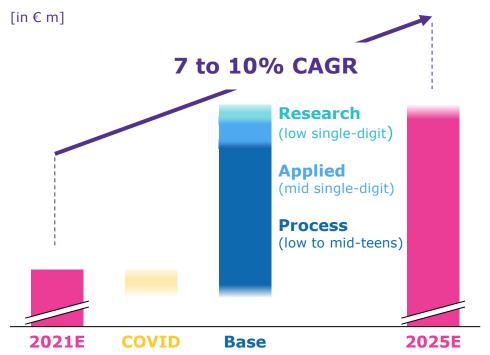
Upgrading mid-term financial ambition to **7-10% org. sales CAGR** (incl. fading COVID business)



Life Science

Strong growth and improved mid-term outlook

Life Science mid-term sales outlook



Drivers of strong mid-term outlook

- LS-wide: Above-market performance with continued positive YoY growth rates despite expected fading COVID-19 demand¹
- Process: Driving ~80% growth based on three main pillars
 - Capitalizing on strong demand for products for traditional modalities via network / capacity expansion and innovation (e.g. BioPharma 4.0)
 - Building scale in services for all modalities with a focus on mAbs, hp-APIs, ADCs, viral vectors and mRNA
 - Templating products for novel modalities
- Applied and Research: Building on strong positions with continued robust growth in line with historical trends



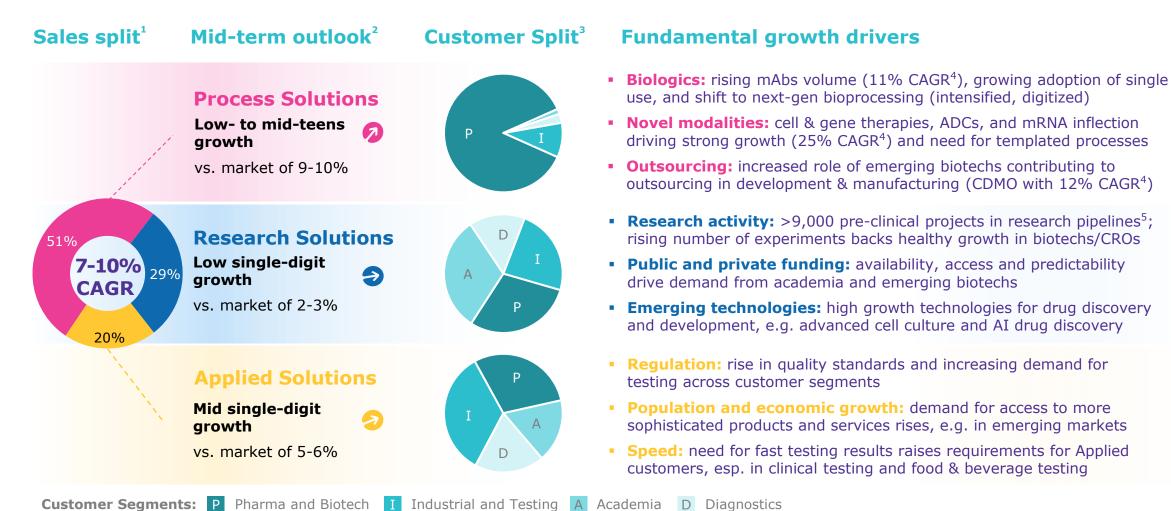
Confident to deliver above-market growth despite expected fading COVID-19 business

¹Mid-term organic sales outlook of 7 to 10% CAGR assumes fading COVID-19 related business between two scenarios: A) zero COVID-19 related business in 2025 [implied CAGR of 6 to 9%] and B) COVID-19 related business in 2025 on 2021 level [implied CAGR of 8 to 11%]



Life Science

All business units contributing to strong mid-term outlook



¹ Based on H1 2021, CAGR is organic mid-term ambition including anticipated fading of COVID contribution; ² market growth rates are excluding COVID effects; ³ indicative only; ⁴ Source: company estimate of market growth based on industry forecast over 5-year horizon; ⁵ Source: statista; Acronyms: mAbs = Monoclonal Antibodies; ADCs = Antibody Drug Conjugates; mRNA = Messenger Ribonucleis Acid, CDMO = Contract Development Manufacturing Organization; CRO = Contract Research Organization; AI = Artificial Intelligence

Life Science Strategy

Focus on strengthening the core and expanding in high-growth areas



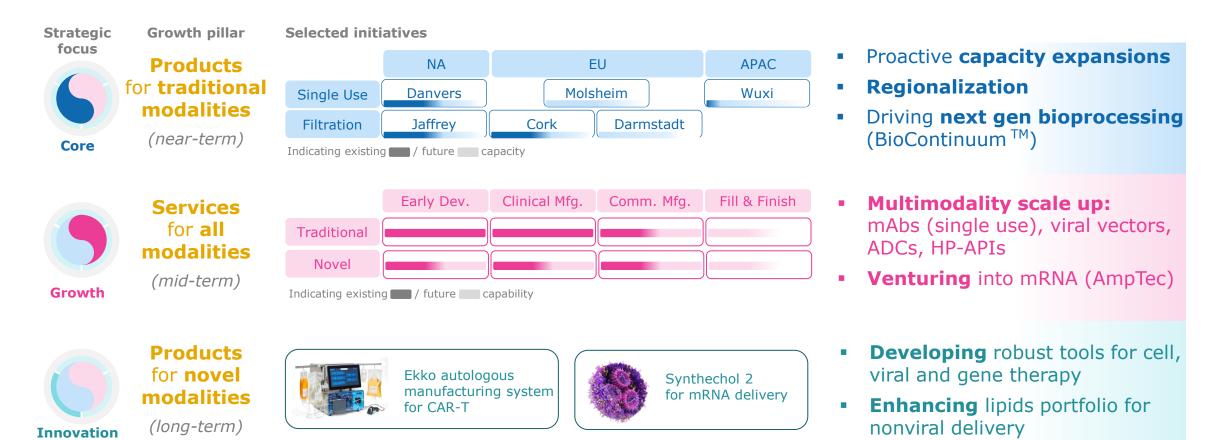


Stringent execution of strategic priorities driving sustained profitable growth; upgrading mid-term outlook to 7 to 10% org. sales CAGR (incl. fading COVID business¹)

¹Mid-term organic sales outlook of 7-10% CAGR assumes fading of COVID-19 related business between two scenarios: A) zero COVID-19 related business in 2025 [implied CAGR of 6 to 9%] and B) COVID-19 related business in 2025 on 2021 level [implied CAGR of 8 to 11%]

Process Solutions – the largest of the BIG3

Fueling the growth engine to accelerate further in a dynamic market



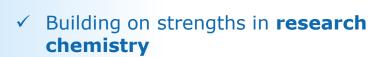


Robust supply enabling attractive base growth, services and innovation as accelerators

Acronyms: mAbs = Monoclonal Antibodies; HP-APIs = High Potency Active Pharmaceutical Ingredients; ADCs = Antibody Drug Conjugates; mRNA = Messenger Ribonucleic Acid

Leveraging strong positions for durable growth in robust markets

Selected initiatives Research Solutions



✓ Capitalizing on evolving CRO market



Strategic focus

Selected initiatives Applied Solutions

- ✓ **Lab Water** instrument platform innovation
- ✓ Expanding Pharma QC testing (biologics & novels)





- Innovating to accelerate research biology
- ✓ Expanding further in China/APAC



Growth

✓ Enhancing custom services for diagnostics



✓ Expanding further in China/APAC



Leveraging upgraded eCommerce platform



✓ Driving digital solutions





Strategic focus on sustaining core positions and capturing higher growth segments

Innovation, digital and portfolio

Key supporting pillars of strategy for value creation

Innovation

Addressing evolving market ...



Sensing unmet customer needs and trends



Building on **tech strengths** and **interdisciplinary expertise**

... to ensure differentiation



Continually **raise the bar** in core portfolios



Set the standard in rapidly evolving segments

Sustaining leading positions and capturing new growth areas

Digital

Internal



Data & AI driven decisions



Infrastructure & capabilities

External



Digital engagement



Go-To-Market



Digital solutions

Drive business value and address evolving customer expectations

Resource allocation

Focus on organic growth...



Capex increasing $\sim 2.5x^1$



Target R&D ratio ~5% of sales

...paired with targeted bolt-on acquisitions



Technologies



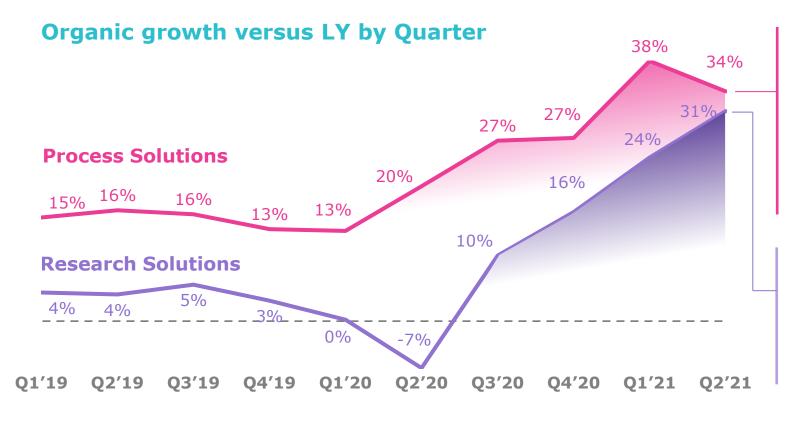
Capabilities

Maximizing returns through strategic capital allocation

¹Comparing average capex spend p.a. over the period 2016-20 with the period 2021-25



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





нealthcare Executing on the earnings phase

Healthcare

Well positioned in a dynamic environment, focused leadership in R&D

Existing market trends unchanged...

- High growth in largest TA Oncology
- Precision medicines to further increase share
- Cost pressure and pricing volatility growing

... well positioned to grow further

- Long-term Oncology strategy with diversified clinical portfolio
- Innovative models of treatment personalization, e.g. in Fertility
- Diversified geographical sales footprint, lower exposure to potential U.S. pricing reforms
- Global while local 18 manufacturing sites and 4 R&D hubs across the globe
- Broad spectrum of payer types due to TA and geographical variety

Focused leadership in R&D driven by recent launches & pipeline



- ✓ From correlated to distinct uncorrelated risks
- ✓ Variety of pathways supported by new modalities e.g. ADCs, Oral ATRi, TIGIT...



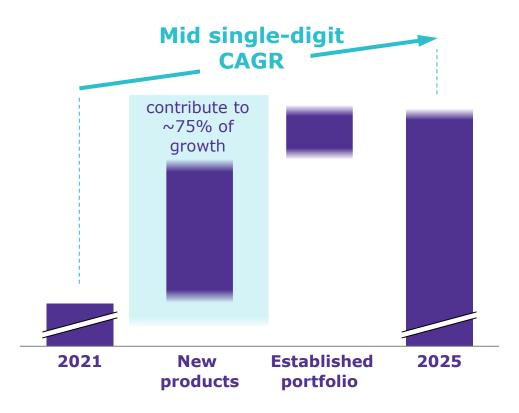


Introducing mid-term financial ambition of mid single-digit org. sales CAGR



Healthcare

Growth driven primarily by innovation, augmenting a solid established portfolio foundation



Risk-adjusted illustration

Positioned to accelerate science & technology leadership



Profitable sales growth above global pharmaceutical market (4.6% across TAs until 2025)1



New products

- Evobrutinib (BTKi) to strengthen our leadership position in Multiple Sclerosis
- Multiple assets within our targeted Oncology TA focus
- Committed to drive our recent launches Bavencio, Mavenclad[®] and Tepmetko[®]



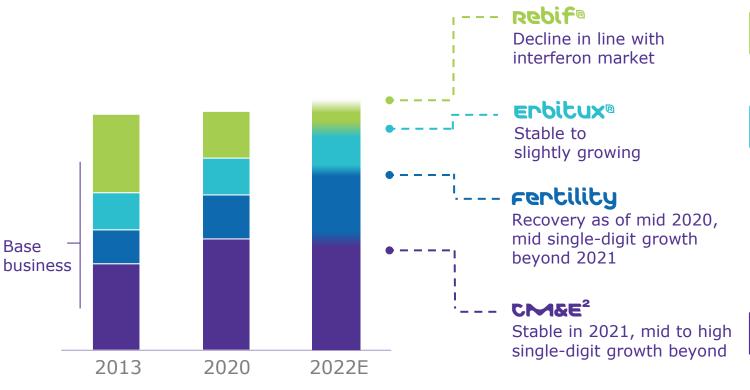
Sustainable long-term growth for Healthcare beyond 2025

¹ Company estimates of mid-term growth outlook based on industry forecasts and reports from public research institutes (e.g. IQVIA Global Medicine Trend Report from April '21)

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)

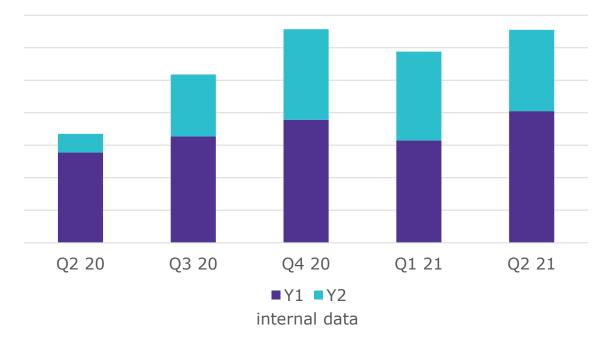
Healthcare: Mavenclad® US

Q2 Y1 patients growing QoQ with expanded prescriber base



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





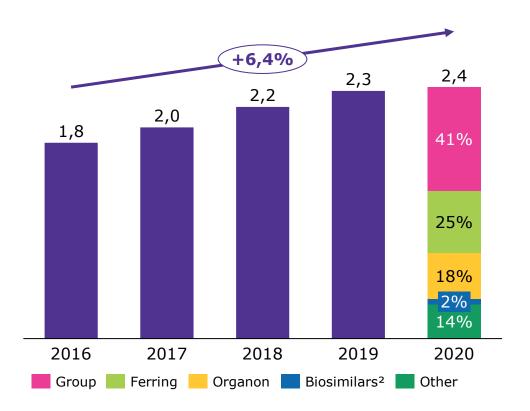
- New patients growing QoQ both in terms of volume and % of total patients
- New prescribers growing QoQ by 23%
- **Total patient growth in H2 2021** to be driven by:
 - Higher volume of H2 2020 return patients
 - Continued new patient growth



Healthcare

Strengthening our leadership in Fertility

Global Gonadotropin market in € bn¹



Strategies to expand our leadership position

GONAL-f The world's most prescribed r-hFSH treatment³

- Growing prevalence of infertility driven by delayed parenthood and lifestyle habits
- Increasing awareness and access to treatment

Pergoveris: Only recombinant FSH + LH product in the market

- Doubling sales and market share since 2017
- Strong growth potential by geographical expansion

Innovations & digital solutions (e.g. Philips partnership)

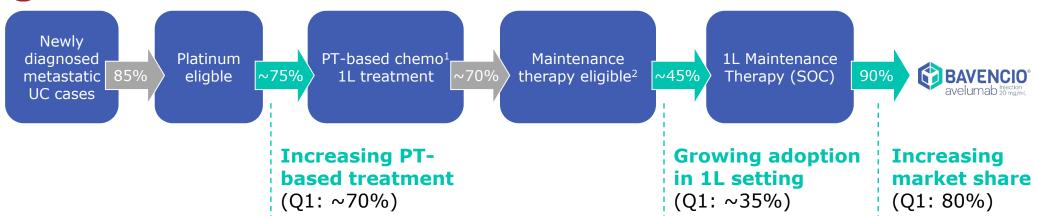
- Better treatment insights & improved convenience
- Contributing to higher live birth rates



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:



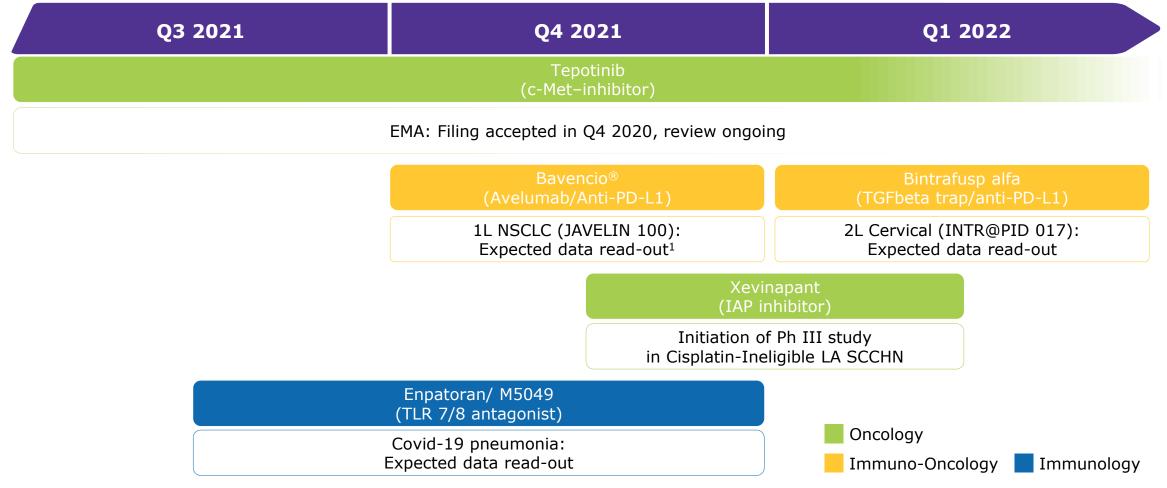


Europe & Japan – Recently approved, encouraging uptake:

- 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



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, ¹Clinical timelines are event-driven and may be subject to change

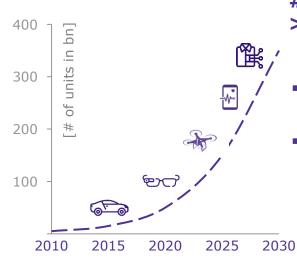




Electronics industry has clear roadmaps with vast market potential and substantial need for materials innovation

More applications than ever before to fuel **Semiconductor growth**

12% Automotive
7% Server / Storage / Communication infrastructure (5G)
6% Consumer (wearables/IoT)
5% Industrial/Medical/Military/Aviation
4% PC/Computing
4% Mobile phones / tablets

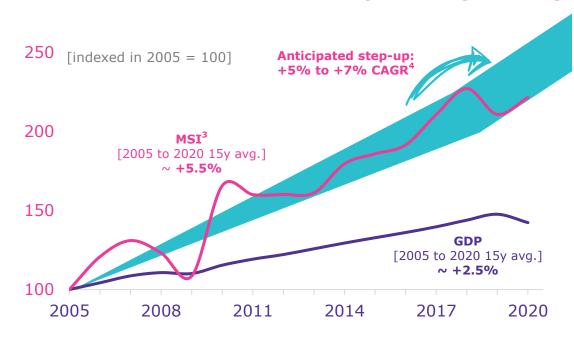


of AIoT¹ devices >350 bn in 2030

- More innovative chips required
- Value-share of chips per device increasing from:

~18% in 2020 to **>30%** in 2025²

Growth expected to accelerate... ...with reduced upward cyclicality

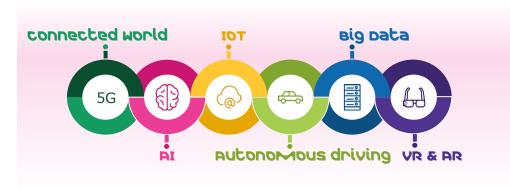


- Diverse end applications & higher capital discipline expected to reduce future cyclicality
- Semiconductor Materials show lower cyclicality than
 Semiconductor market

1) AIoT = Artificial Intelligence + Internet of Things; 2) McClean, Prismark Semiconductor and packaging report Q4 2020; 3) Million Square Inch of silicon wafers; 4) Group estimate based on industry forecasts

Building on leading positions in semiconductor materials

External trends drive strong demand



Tech trends inevitably drive exponential data growth requiring semiconductors and displays in virtually all data applications

- Customers & governments significantly accelerating semi capacity expansions
- Semiconductor materials market expected to accelerate growth based on broader demand

Electronics well positioned to capture it

- One of the strongest portfolios providing solutions necessary to enable industry growth
- Enabling breakthrough technologies, in high value areas of wafer processing & display innovation
- Pioneering in high-throughput experimentation and Big Data & AI projects with key customers
- Shift to executing growth accelerating investment and innovation in sync with customer plans



Further upgrading mid-term financial ambition to 3 to 6% org. sales CAGR

Abbreviations: AI = Artificial Intelligence; IoT= Internet of Things; VR = Virtual Reality; AR = Augmented Reality

Shifting gears from transformation to growth execution

From building a platform...

Successful "Bright Future" transformation

- Superior business composition
 - ~10 equally sized, differentiated tech platforms
 - Wider customer base than ever before
 - Balanced presence in all major customer hubs
- Clear focus on Semi & innovation

Significantly higher Versum synergies

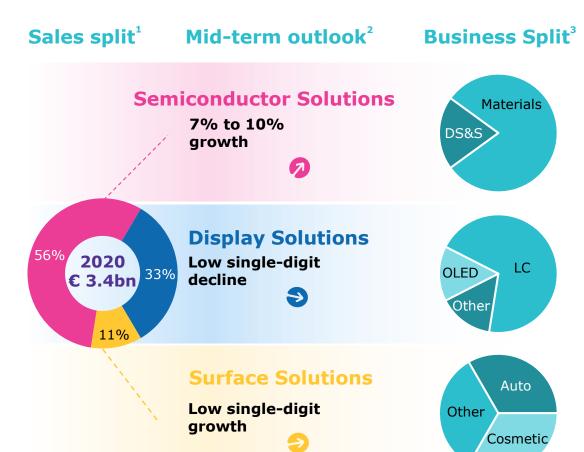
- Accelerated capex investment plan
 - Smart localization
 - Global presence & local proximity
- Display & Surface well managed according to Enterprise Planning Unit role



- Semiconductor Solutions: to grow 200 to 300bps²
 above faster underlying market of 5% to 7%
- **Display Solutions:** low-single digit decline in the coming years expected to return to growth until 2025
- Surface Solution: low single-digit growth post COVID-19 turnaround
 - 1) illustrative split by semiconductor solutions technology platform
 - 2) "Basis points"



Portfolio refocus drives mid-term guidance upgrade to 3 to 6% CAGR



Fundamental growth drivers

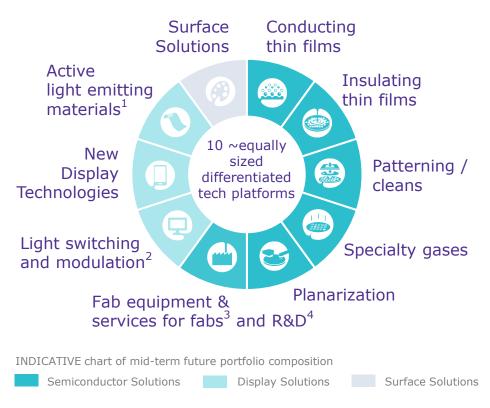
- Continued market growth due to technological advances (Artificial intelligence, 5G, Big Data and cloud, Internet of Things) serving customers in Logic, Memory, Packaging and others
- 5 to 7% market growth⁴
- 200 to 300bps above-market growth from share gains & better portfolio (incl. 100 to 150bps 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



¹ Based on FY 2020, CAGR is organic mid-term ambition; ²growth rates are organic CAGRs; ³indicative only ⁴Source: Group estimate based on industry forecasts; 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

Electronics: Successful transformation, ready for accelerated growth

An integrated electronics innovation leader with one of strongest industry portfolios



- 1) Incl. materials for OLED displays 2) Incl. liquid crystal displays
- 3) Delivery Systems & Services 4) Intermolecular

Delivering on promises and above

- Bright Future program key deliverables & savings
- €3.3 bn sales with ~90% of sales focused on electronics
- Successful integration of Versum
- Attractive, stable EBITDA pre margins
- Display & Surface well managed according to Enterprise Planning Unit role
- Upgraded mid-term guidance of 3% to 6% sales CAGR

Everything in place to gain further speed as vital growth engine for the Company

"Level Up" - Shifting gears from transformation to growth execution

Level up **scale**

- Capacity investments synchronized to customers' expansion plans
- Ability to tackle industry challenges & supply reliability
- Continue localizing footprint close to customers in Korea, Taiwan, China, and U.S.

Enabling ambition of 3 to 6% org. CAGR 2021 to 2025+



Level up **tech**

- Further sharpen focus on profitable innovation, addressing key inflection points
- Address sustainable innovation
- Expand R&D and keep one of the highest R&D rates in the industry

Level up people & capabilities

- Attractive employer for NextGen talents
- Drive safety, quality & sustainability
- Drive data analytics & digital transformation

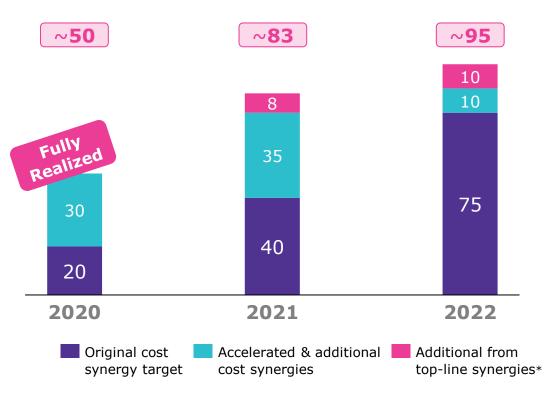
Level up **portfolio**

- Commitment to relevant portfolio breadth in high growth areas
- Diligent bolt-on acquisitions
- Targeted expansion of portfolio



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



Group

Accelerating positive impact on environment and society

Positive impact through our products

- Provide access to medicine
- Enable cutting-edge technologies
- Support greener production

Stabilize production and minimize risks

- Actively manage ESG risks
- Training on policies & standards
- Manage and secure supply chain

Minimize negative impact through our operations

- Achieve climate neutrality
- Manage water efficiency
- Reduce waste

Our sustainability goals

Dedicated to human progress



In 2030, we will advance human progress for >1 billion people through sustainable science & technology

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

Creating sustainable value chains



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

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

Reducing our ecological footprint



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

- Climate change & emissions
- Water & resource intensity



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



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

⁴Sites > 70.000 m³/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)



Group

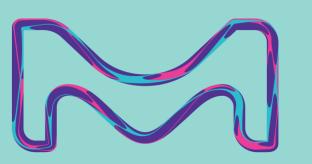
Clear set of tasks to enable our 2030 ESG goals

Tasks 🗒	Development In place	Results and next steps ©
Analysis of requirements: Strategy, business, regulation, stakeholders		Ongoing due to developing field of ESG stakeholder expectations and regulation
Build effective data platform for internal steering		Feed in internal data, external data, industry benchmarks and modelling projects in 2021
Develop ESG KPIs for reporting		Developing & evaluating ESG KPIs for steering, milestones, decide on reporting in 2021
Develop SBV tool¹ to measure product sustainability value		SBV tool developed and tested in up to 10 cases by 2021
Link ESG ² to board compensation with 20% sustainability factor from 2022		Approved by AGM, qualitatively included, quantitative factor to be developed in 2021
Further incorporate ESG ² in R&D, Controlling, M&A and Supply Chain		ESG-Framework M&A/Capex in 2021 , progress in supply chain, controlling, R&D
Decide on dedicated investments and initiatives to achieve targets		Business strategies and priorities under development, budgeting & decisions in 2021

¹Sustainable Business Value: Dive in deeper and read the research article on the <u>SBV method</u>; ²ESG: Environmental, Social, Governance







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%

Executive SUMMary

4 key priorities to deliver on our science & tech ambitions

Focusing on **Sustainability**

Mobilizing for Efficient growth



Driving
Culture & Leadership





xibneqqa





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

- Organic: +18% to +21%
- 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

- Organic: +6% to +8%
- 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

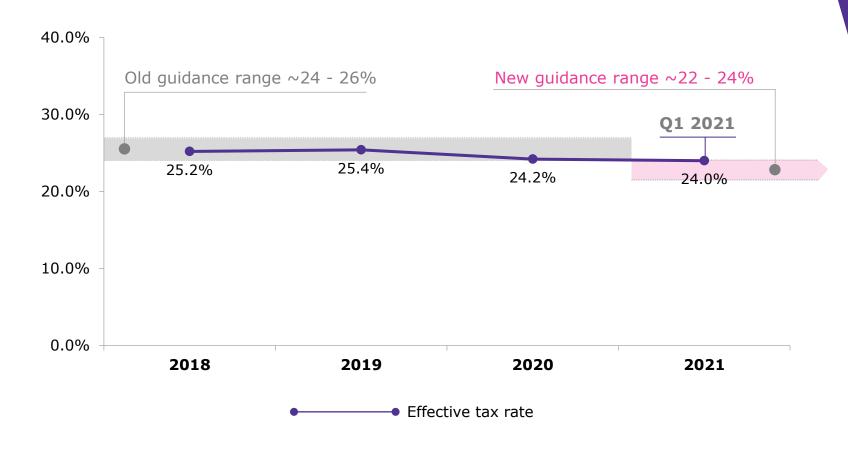
Further financial details

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

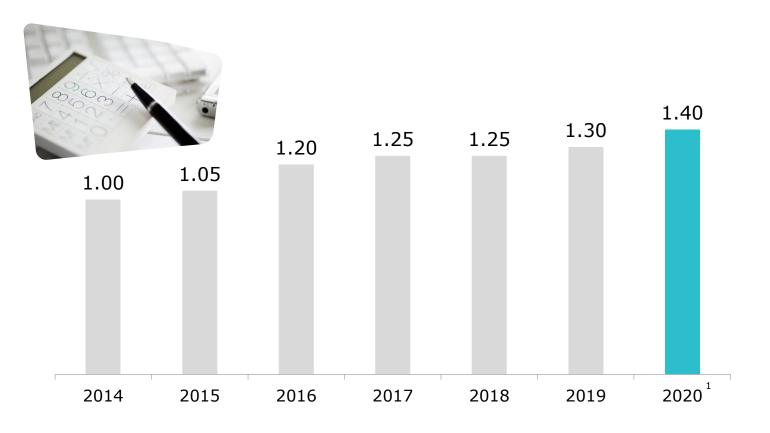
- Deleveraged to ~2x net debt/EBITDA pre already in 2020
- M&A on hold until 2022; only smaller deals to be realized if budget available
- Annual capex guidance: from ~€1.5 bn in 2021 to ~€2 bn by 2023 and increased focus on organic investment
- Dividend policy mirrors
 sustainable earnings trend

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

Group's ownership structure

Shareholders of Merck KGaA, Darmstadt, Germany

- Shareholders hold ~30% of Company'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 Company's shareholders:

- Approving financial statements of the Group
- Working together with Executive Board, receiving reporting on progress regarding business and financial development at the Company, 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) holds ~70% of Group's total capital
- These 70% of the total capital are not entitled to vote at Group's 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 E. Merck KG, Darmstadt, Germany
- 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

ariable	40-50%	 Reflecting the long-term strategy for Group's growth and (from 2022) sustainability ambition 4 years performance cycle: 3 years target achievement + 1 year holding period Based on virtual Company 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
>	25-35%	 Three-years average profit after tax of the Group, multiplied with individual permille rate From 2021 reduced individual performance factor of 0.8-1.2 can increase (bonus) or decrease (malus) the amount based on a set of criteria, 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 €) Mandatory personal investment in Group Shares amounting to one third of the net payment of the profit sharing (4 year holding period)
	6-9% 0-3%	Pension Entitlements Defined contribution Additional Benefits Mainly contributions to insurance policies, personal security expenses, company car
Basic	15-20%	 Basic Compensation Fixed and non-performance related compensation Paid in 12 equal monthly installments 1.4 million € for the chairman / up to 1.1 million € for the members of the executive board 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, we are part of FTSE4Good Index, measuring the performance of companies with strong ESG practices (top 15).



In 2020, we have once more achieved **prime status** by **ISS Oekom**.



In 2019, our 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.



We 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 we 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).

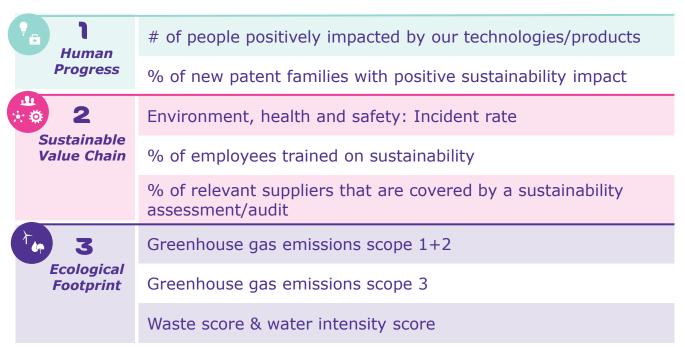
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Group

Expanding KPIs to monitor and steer sustainability comprehensively

Target Exemplary KPIs to be implemented in 2021





- KPI set will evolve over time
- External reporting requirements: 140+ ESG indicators



Our sustainability ambition is integrated in the Executive Board compensation system

From 2021 onwards

Explicit reflection of the new targets on qualitative level (profit sharing modifier)

In addition, from 2022 onwards

Quantitative sustainability factor in the long-term incentive plan for the Executive Board

Set of sustainability KPIs will build the underlying basis for the development of the sustainability factor



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

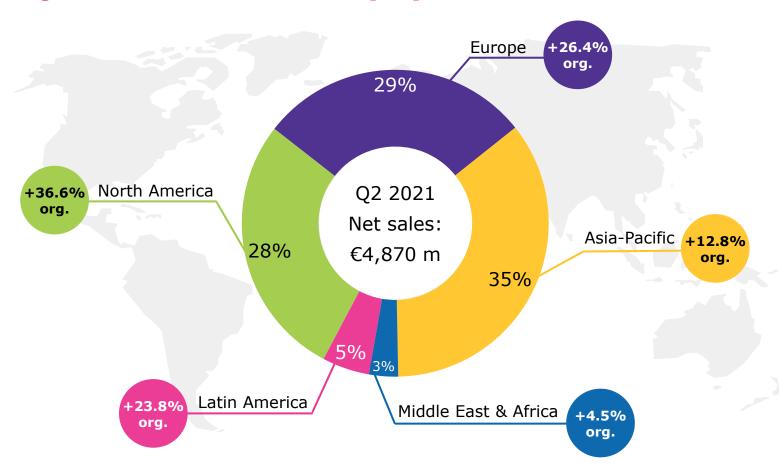
- Supporting profitable growth strategy
- IRR > WACC
- EPS pre accretive
- Maintain investment grade rating



- 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

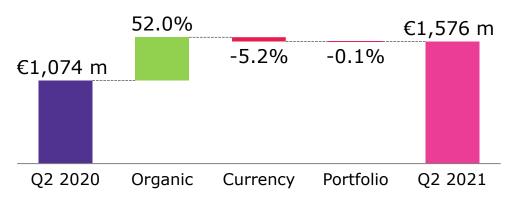


Strong Life Science demand and robust Healthcare recovery drive 23% sales and 52% EBITDA pre organic growth

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



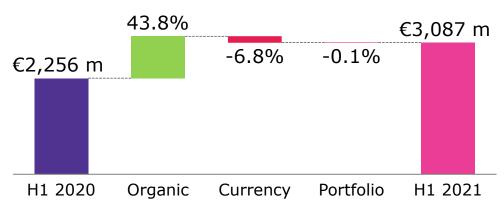
- Organic EBITDA pre increases by more than 50% and more than twice as fast as sales
- Strong uptake in Life Science and Healthcare gross profit paired with continued cost discipline in all sectors vs. soft Q2 2020
- FX burden of -5% across various currencies with largest negative impact from USD and 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 1.625% 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

[€m]	H1 2020	H1 2021	Δ
Net sales	8,489	9,501	11.9%
EBITDA pre	2,256	3,087	36.9%
Margin (in % of net sales)	26.57%	32.49%	5.919%
EPS pre	2.80	4.42	57.9%
Operating cash flow	1,019	2,104	106.6%
[€m]	Dec. 31, 2020	June 30, 2021	Δ
[€m] Net financial debt	Dec. 31, 2020 -10,758	June 30, 2021 -10,141	Δ -5.7%
Net financial debt	-10,758	-10,141	-5.7%

- 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 2020		Q2 2021	
	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

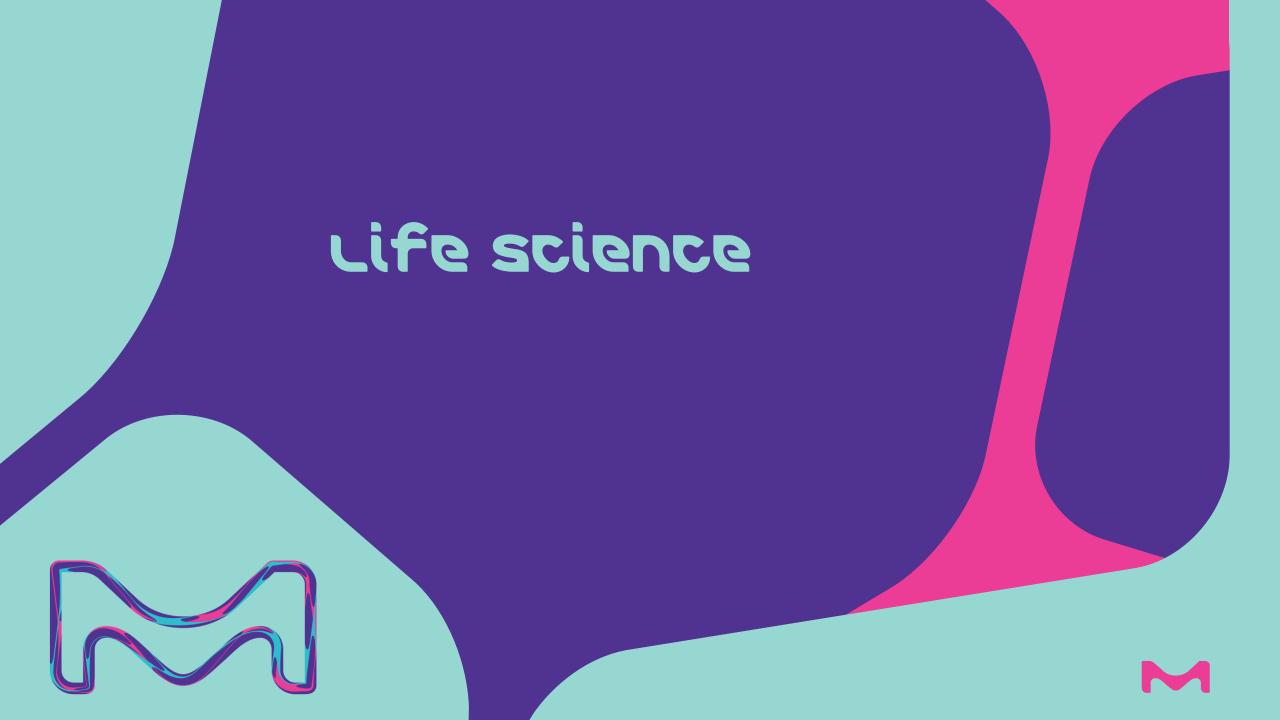
[€m]	H1 2020		H1 20	021
	Adjustments	thereof D&A	Adjustments	thereof D&A
Healthcare	-12	2	21	3
Life Science	-4	0	8	0
Electronics	165	112	27	7
Corporate & Other	25	0	103	1
Total	174	114	159	11





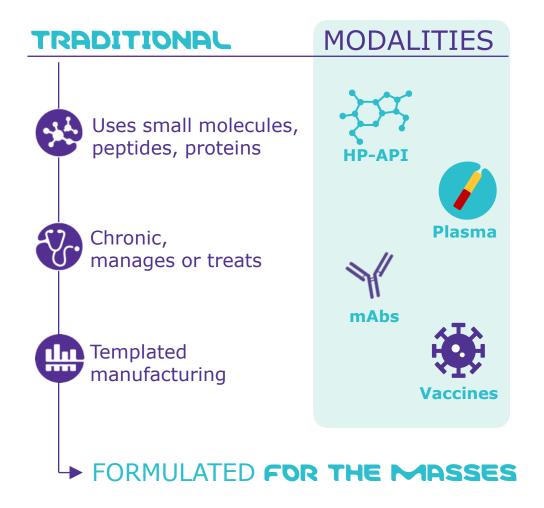
Financial Calendar

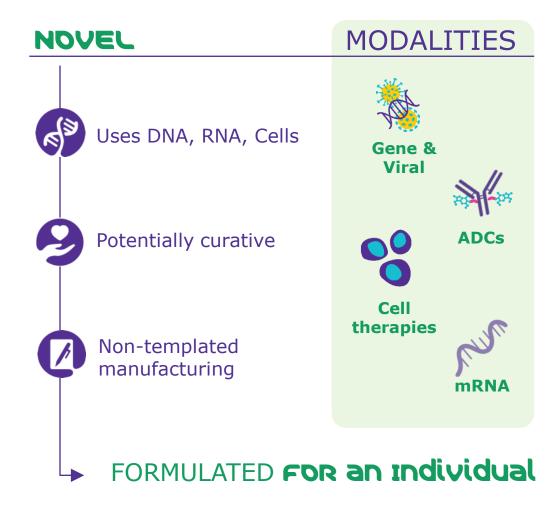
Date	Event
August 5, 2021	Q2 2021 Earnings release
September 9, 2021	Virtual Capital Markets Day
November 11, 2021	Q3 2021 Earnings release
April 22, 2022	Annual General Meeting
August 4, 2022	Q2 2022 Earnings release



Process Solutions: Therapies are evolving from treatments to cures

Advancing traditional is critical as novel modalities develop



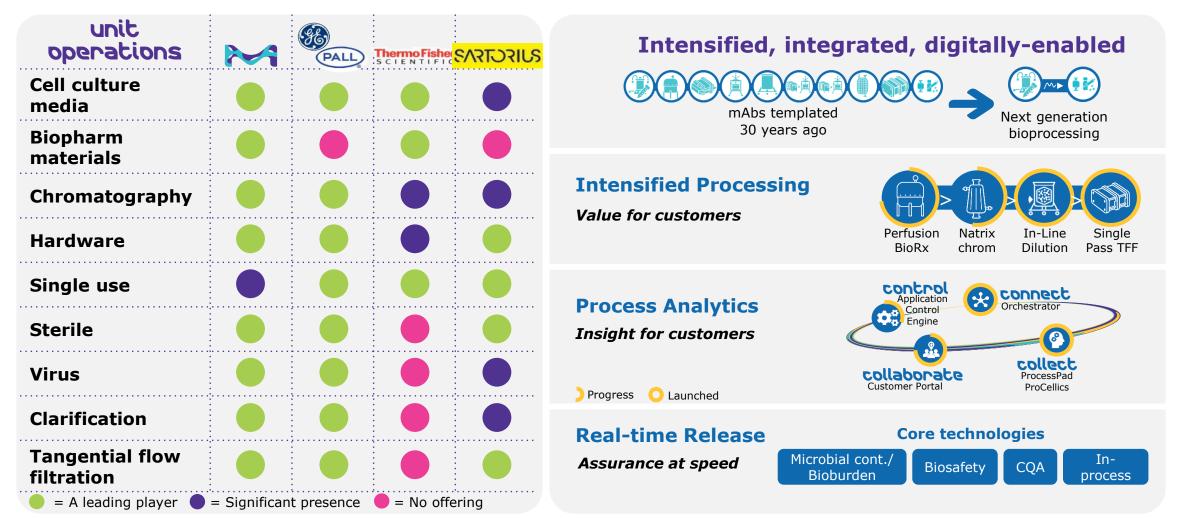


Acronyms: HP-API = highly potent active pharmaceutical ingredient; mAbs = monoclonal antibodies; DNA = deoxyribonucleic acid; (m)RNA = (messenger) ribonucleic acid; ADC = antibody drug conjugate



Process Solutions

Moving from today's technology to BioPharma 4.0



Sources: press releases, company reports, and internal assessments



Process Solutions

COVID demands align with existing strengths

COVID-19 Outlook

Type & Global Doses

Implications



Nucleic Acid Vaccines

- 4.8 Billion Gov. contracted
- **4.3 Billion** 2021 manuf.
- Leveraging **human** factory
- Emerging manuf. process
- Lipids are critical

Viral & Protein Vaccines

- 6.8 Billion Gov. contracted
- **8.7 Billion** 2021 manuf.
- Protective immune response
- Multiple templates
- Leveraging Single Use



mAb Therapeutics

- **2.5 Million** Gov. contracted **13.5 Million** 2021 manuf.
- Bind & block virus entry
- Universal templates
- Leading position for 8/9 unit ops

Product offering

Capabilities in all unit operations



CCM &



Plasmid

linearization²



Chrom



IVT &

capping





& Formulation²

Filtration Encapsulation















Cell expansion



Chrom



Filtration¹



Fill & **Finish**



SU Cons¹





CCM & BPM¹



Cell expansion



Chrom



Filtration¹



Fill & **Finish**



SU Cons¹

1) Used across manufacturing unit operations; 2) Lacking GMP enzymes and nucleotides products; Source: press releases, company reports, internal assessments; Acronyms: CCM = cell culture medium, BPM = biopharma materials, IVT = in vitro transcription, SU Cons = single use consumables



Process Solutions: Strategic direction

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



Modality









Revenue driver



Near term



Mid term



Long term



Offering



Products

Filtration

Technology



Services



Products



Portfolio



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



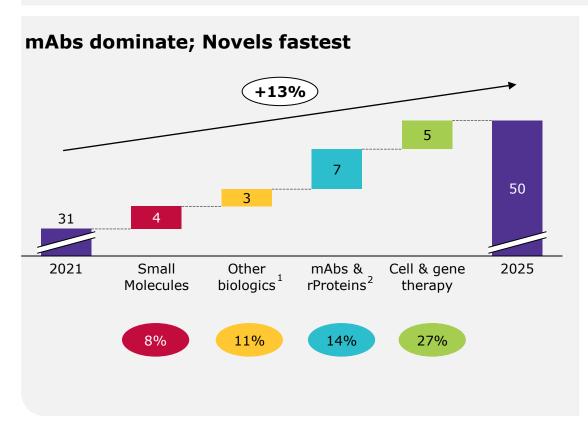
- Templated viral manufacturing consumables and reagents
- ✓ Autologous manufacturing system

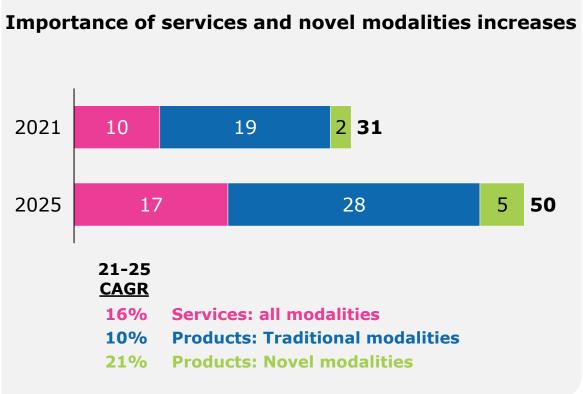


Process Solutions

Opportunities in services to accelerate double-digit growth

Accessible Market (€ bn)





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 Acronyms: mAbs = monoclonal antibodies, rProteins = recombinant proteins, ADCs = antibody drug conjugates



Next-generation bioprocessing on the cards



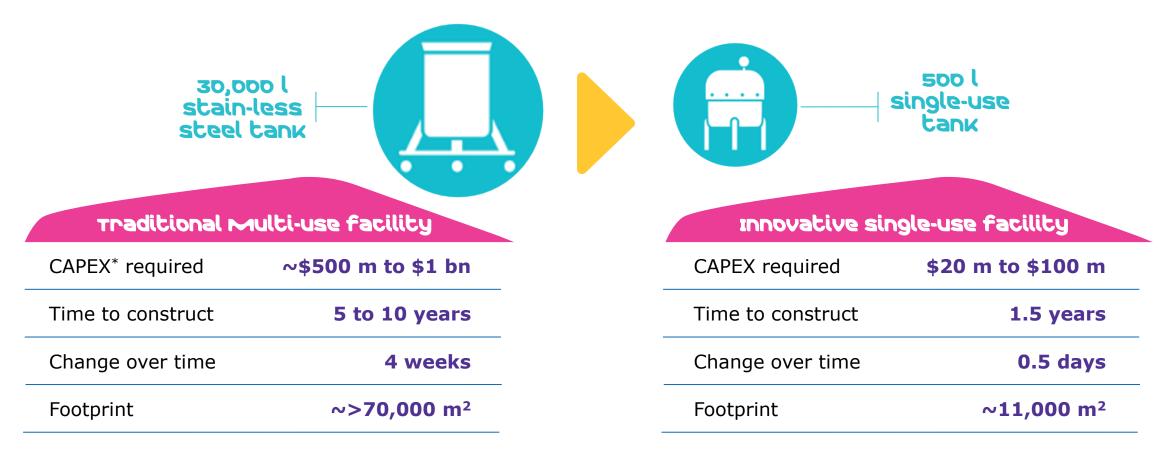


Continuous bioprocessing will ...

- 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





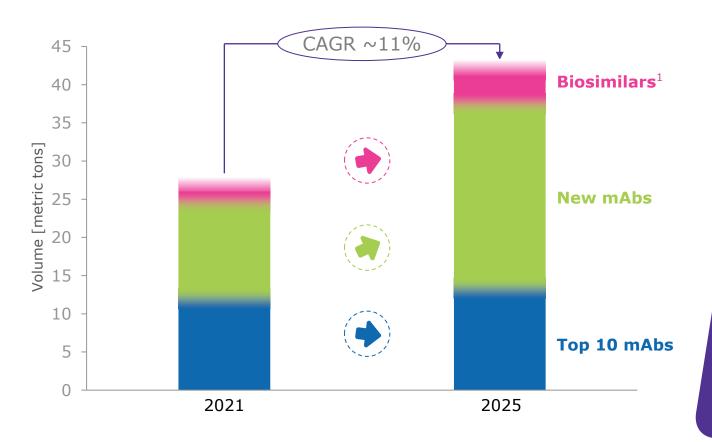
Strong demand for single-use technologies and Process Solutions' broad offering was and will remain a key source of growth for Life Science



Process Solutions

mAbs market democratization will drive diversification, change & variability

mAb volume projections 2021 to 2025



market development

- mAbs market will grow ~11% CAGR
- Top 10 originator mAbs represent ~45% (11mT) of volume today; volume grows but share declines to ~35% (14mT) in 2025
- Biosimilars continue to gain share

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



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

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

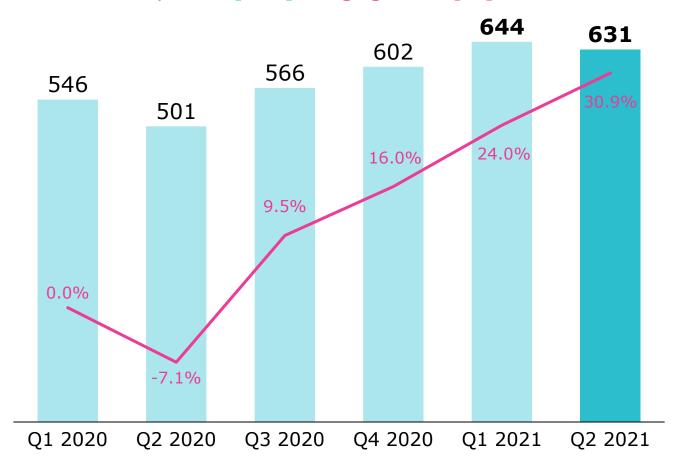


- Continued double-digit growth in the core business paired with rising COVID-19 contributions (mainly vaccine related)
- BioP as main growth driver, formulation growing fastest, services also strong
- Growth starting to moderate as expected amid rising comps (Q2 2020 with initial COVID-19 contributions)
- 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 [%]

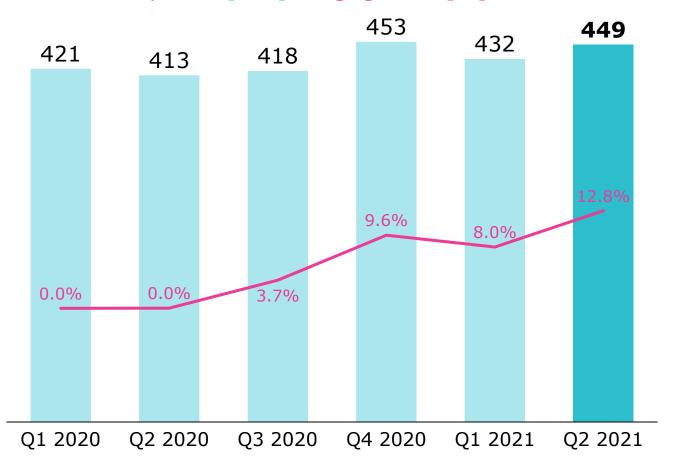


- Strong double-digit growth, accelerating further on soft comps (Q2-20 with biggest impact from lockdowns)
- Ongoing core business recovery and catchup amid rising lab activity
- Diagnostics related COVID-19 tailwinds continue albeit slowing as expected
- North America as fastest growing region, followed 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 [%]

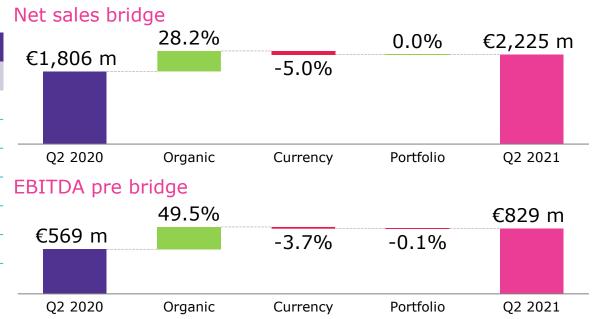


- Growth accelerating to double-digits as recovery continues amid still easy comps (H1-20 with flat growth due to lockdowns)
- Core business as main driver with broadbased performance across business lines
- COVID-19-related sales are negligible
- North America as fastest growing region, followed by 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





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



Round 1		Round 2	Rou	nd 3	Round 4
2019			2020		DATE 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 op.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

J...a. , t. a.c. ca...co. 12

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 inhibitor
Non-small cell lung cancer,
METex14 skipping⁹

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

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

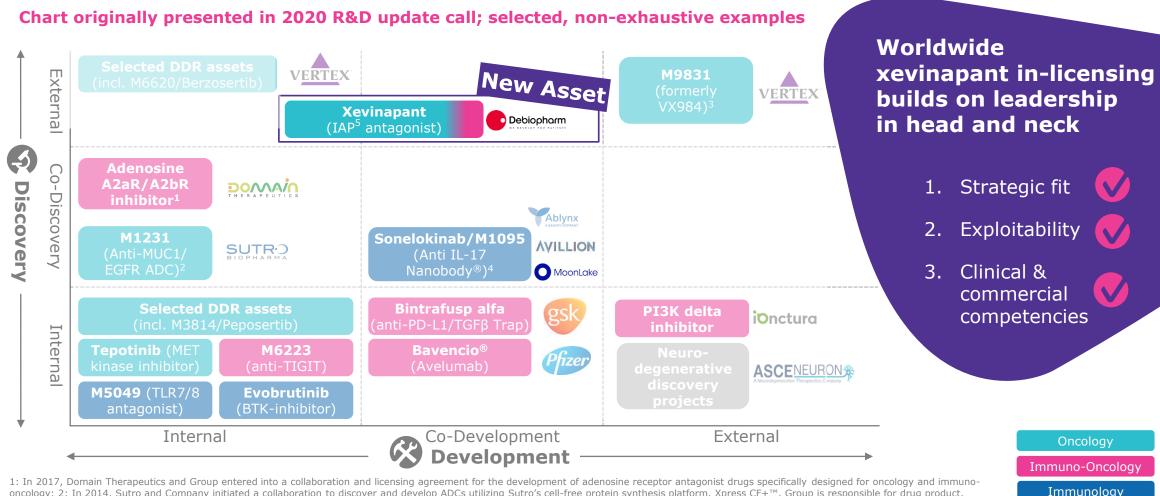
Unless noted otherwise, clinical programs conducted in collaboration with external partners are not shown unless Company has co-ownership of data. In such case the indication is shown in *Italics*. Pipeline products are under clinical investigation and have not been proven to be safe and effective. There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



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. ఄ On March 01, 2021, Group announced a worldwide in-licensing agreement with Debiopharm, Switzerland, for the development and commercialization of xevinapant (Debio 1143). ९ 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.

Xevinapant

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



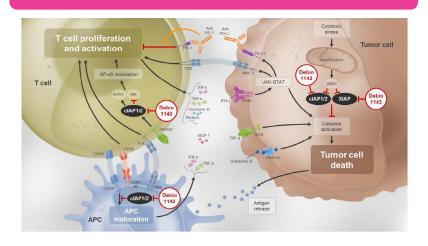
1: In 2017, Domain Therapeutics and Group entered into a collaboration and licensing agreement for the development of adenosine receptor antagonist drugs specifically designed for oncology and immuno-oncology; 2: In 2014, Sutro and Company initiated a collaboration to discover and develop ADCs utilizing Sutro's cell-free protein synthesis platform, Xpress CF+™. Group 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

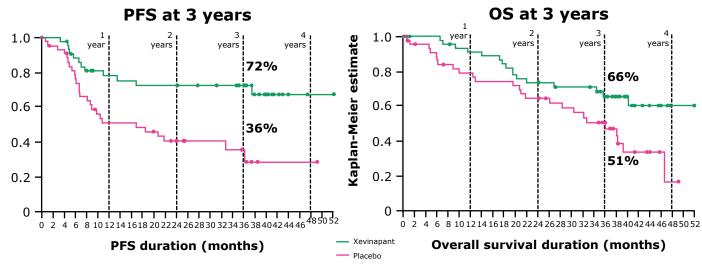
Mode of Action¹



- 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	Group 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	Company to pay industry-typical sales royalty to Debiopharm	

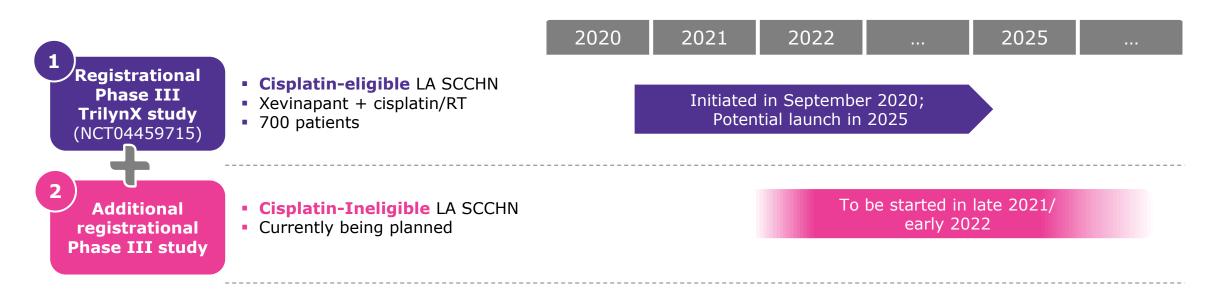
¹ thereof up to ~€300 m for focus H&N indications)



 $^{^{\}rm 2}$ 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



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
	Mavenclad®	23	100% (p = 0.99) ^b
DMT treated patients	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%
Не	Healthy subjects		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

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



EXPLOTE 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



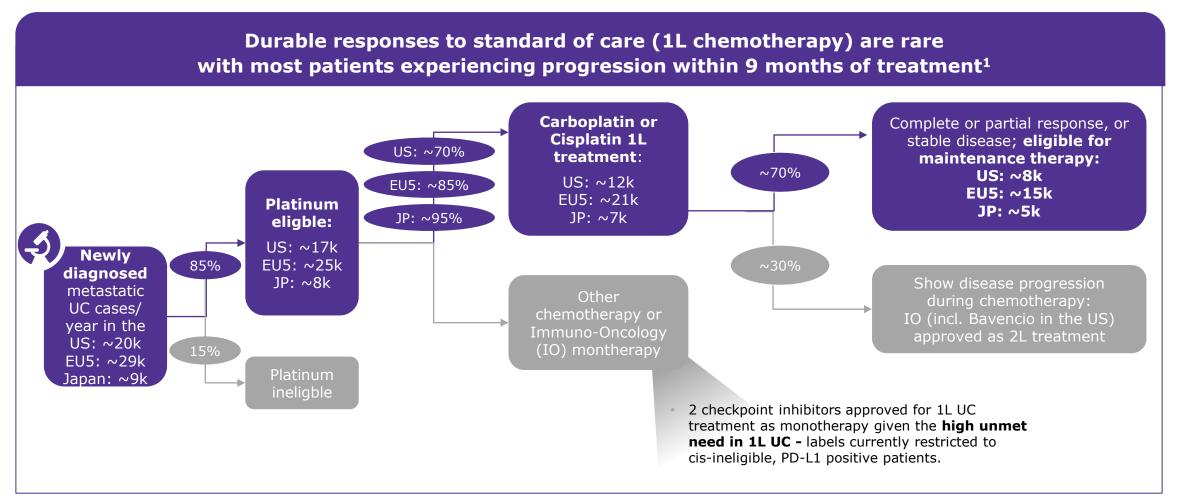




1: Piotrowska et al., "Landscape of Acquired Resistance to Osimertinib in EGFR -Mutant NSCLC and Clinical Validation of Combined EGFR and RET Inhibition with Osimertinib and BLU-667 for Acquired RET Fusion", AACR Cancer Discovery 2018; 2: Wu et al., "Long term outcomes to tepotinib plus gefitinib in 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)



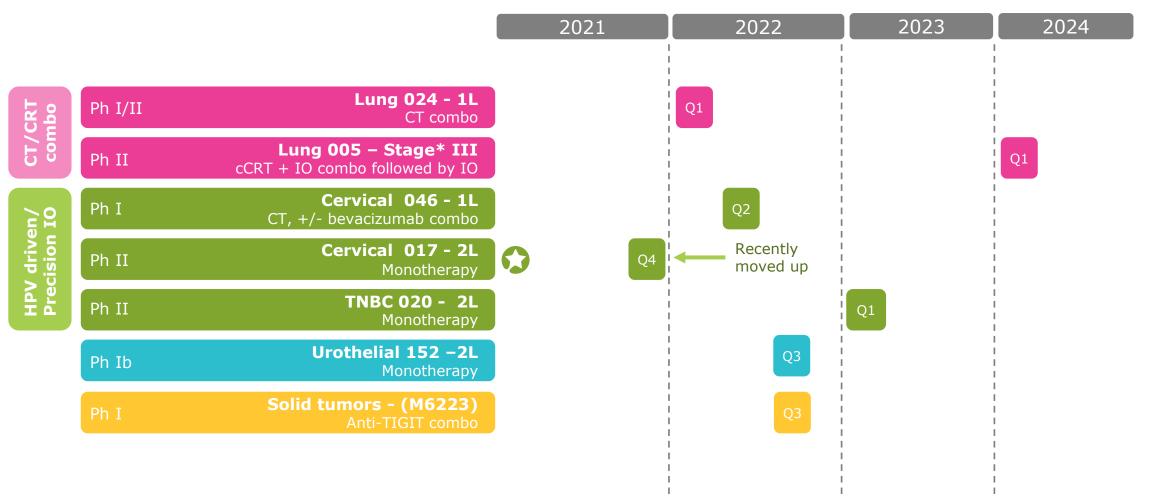




Bintrafusp alfa

INTR@PID Program: Upcoming Readouts





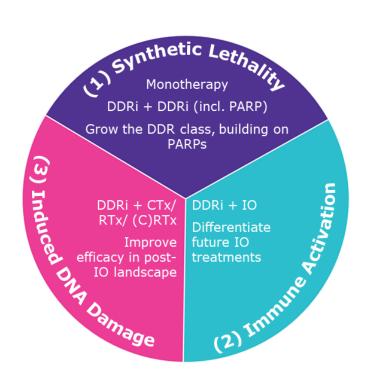
Acronyms: BTC = Biliary Tract Cancer; CT = Chemotherapy; EMT = Epithelial-mesenchymal transition; HPV = Human papillomavirus; NSCLC = Non-small Cell Lung Cancer; RT = Radiation therapy; TNBC = Triple-Negative Breast Cancer; * unresectable; *All clinical timelines are event-driven and may be subject to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

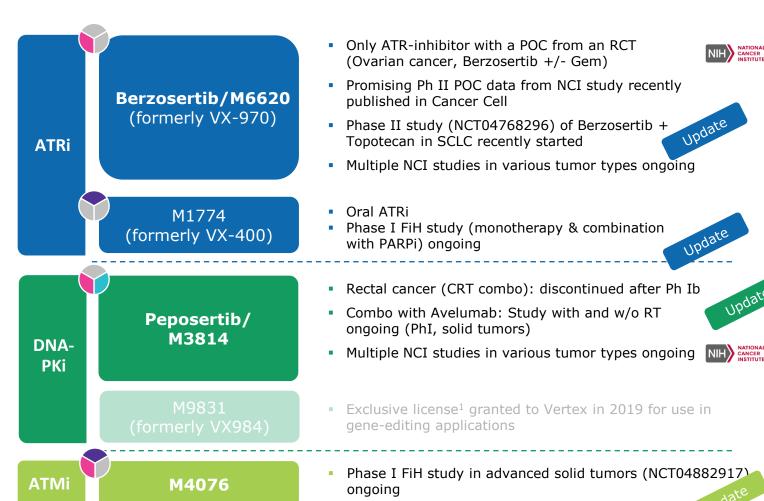


DNA Damage Response (DDR)

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

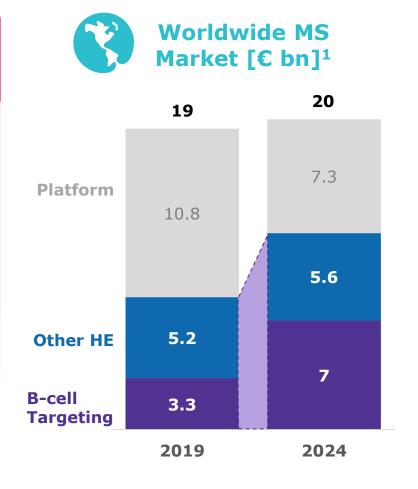


We pioneered BTKi development for MS with Evobrutinib

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

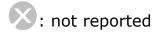
Evobrutinib stands out amongst BTK inhibitors under development

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)
I 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

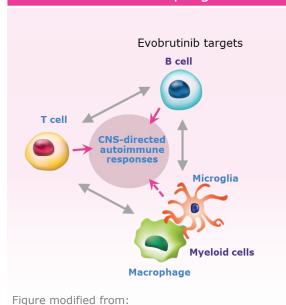




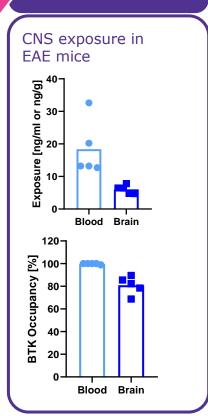
Evobrutinib targets inflammation and progression

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

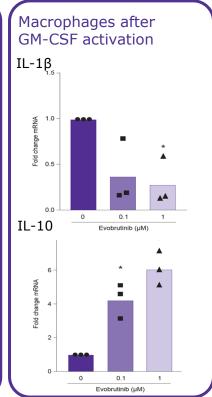
MOA on B-cells and macrophage/microglia cells is an innovative oral approach to MS therapy with potential to impact compartmentalized inflammation and progression



Evobrutinib is a CNS penetrant BTKi²

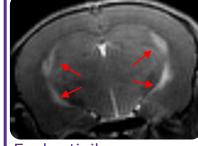


Evobrutinib shifts macrophage phenotypes from M1 pro- to M2 anti-inflammatory ³

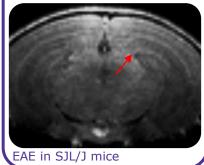


Evobrutinib reduces CNS B cells and leptomeningeal inflammation⁴





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}



Li R, et al. 2018.

^{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

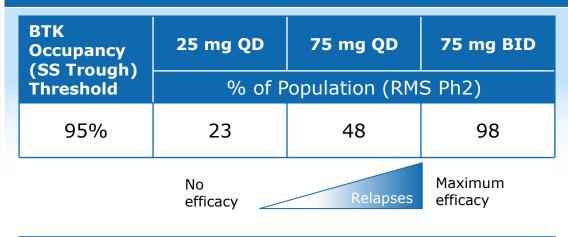
Evobrutinib is optimally dosed to offer best-in-class BTK inhibition Optimized dose selection & targeted covalent binding results in sustained BTK inhibition that is necessary for robust efficacy

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



		Selectivity	action in CNS
втк	Evobrutinib		
Covalent binding			
втк	Tolebrutinib Multiple other kinases	•	
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		

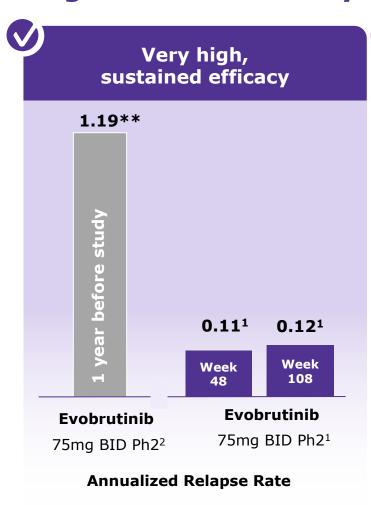


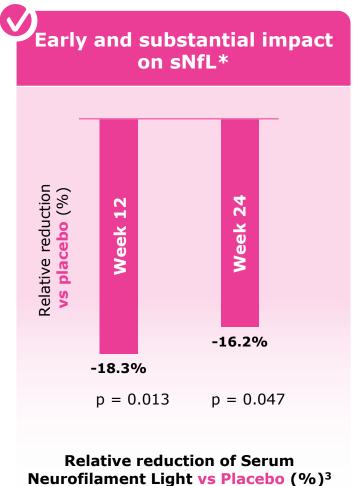
Selectivity* Sustained



Evobrutinib holds unmatched Long-Term Data among BTKi class in MS Best-in-disease efficacy & favorable safety over 2 years in

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

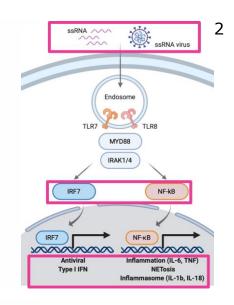


M5049 (TLR7/8 antagonist)

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



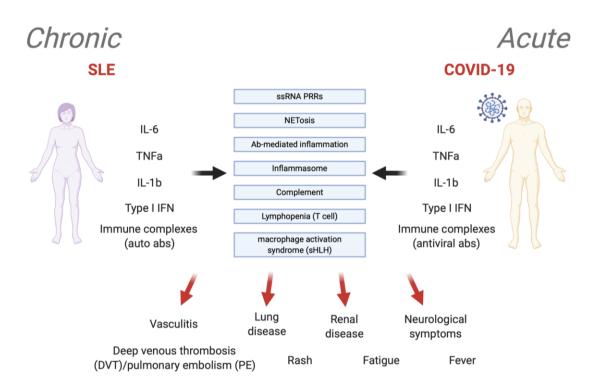
Preliminary Phase I data warrant further investigation as a potential treatment for autoimmune diseases including SLE

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

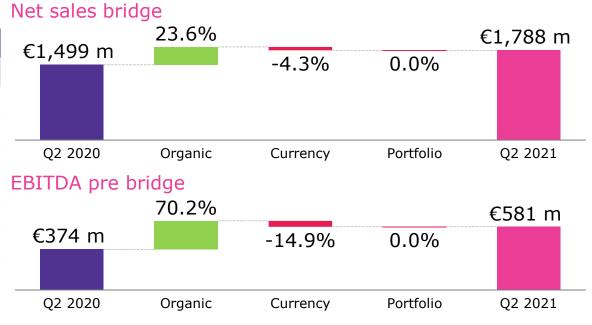


^{1:} Illustration created in-house; Acronyms: SLE = Systemic lupus erythematosus

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

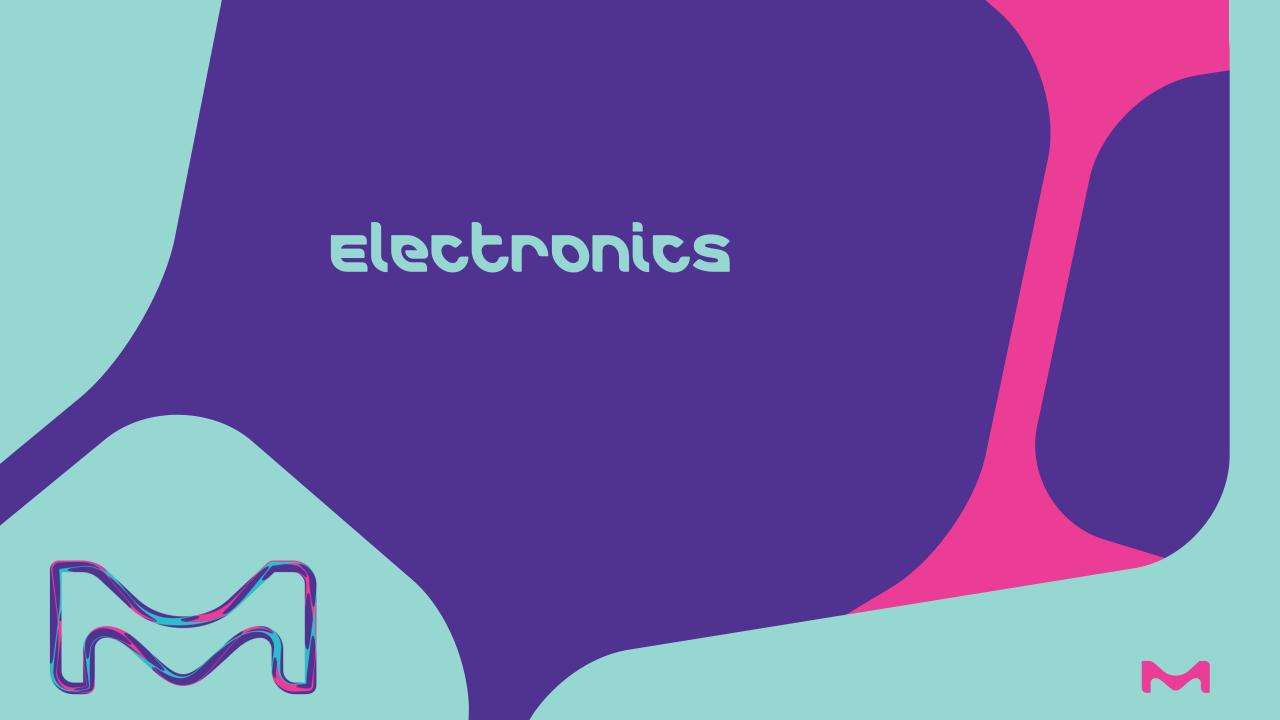
- Mavenclad® grows +102 % organically to €157 m, amid first signs of recovery of dynamic market and low comps; Rebif® declines -9%
- Oncology up +49%; Bavencio[®] grows +206% fueled by UC 1L launch in key markets; Erbitux[®] up +36% supported by Eli Lilly supply agreement
- Base business up +15%, driven 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.



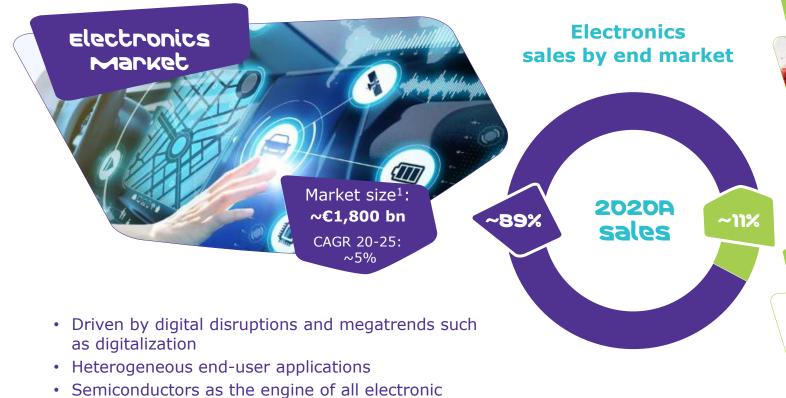
¹ Volume Based Procurement

^{*} Marketing and selling expenses

Totals may not add up due to rounding



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

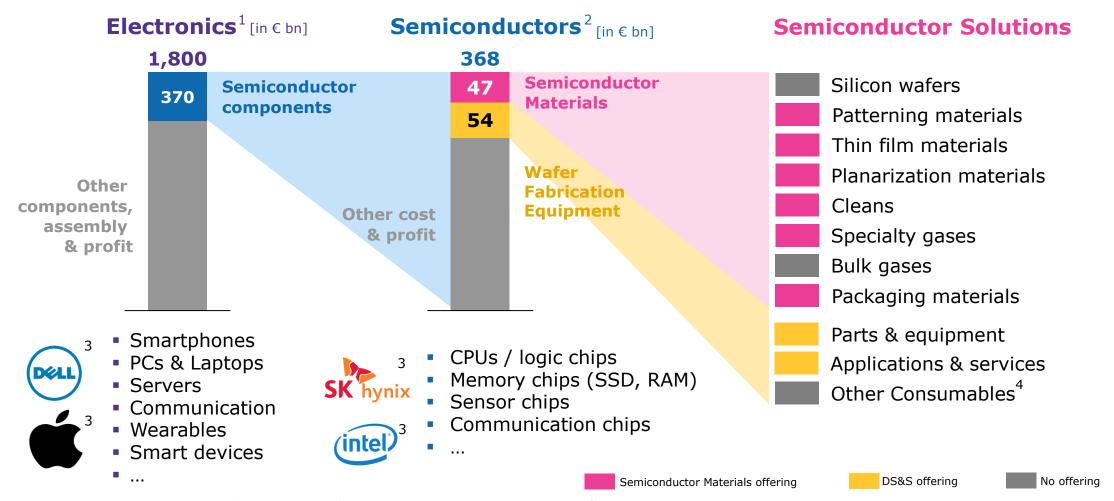
- Driven by world GDP growth
 - Rising living standards and higher disposable income

Market size: ~€400 bn

¹Prismark 2021

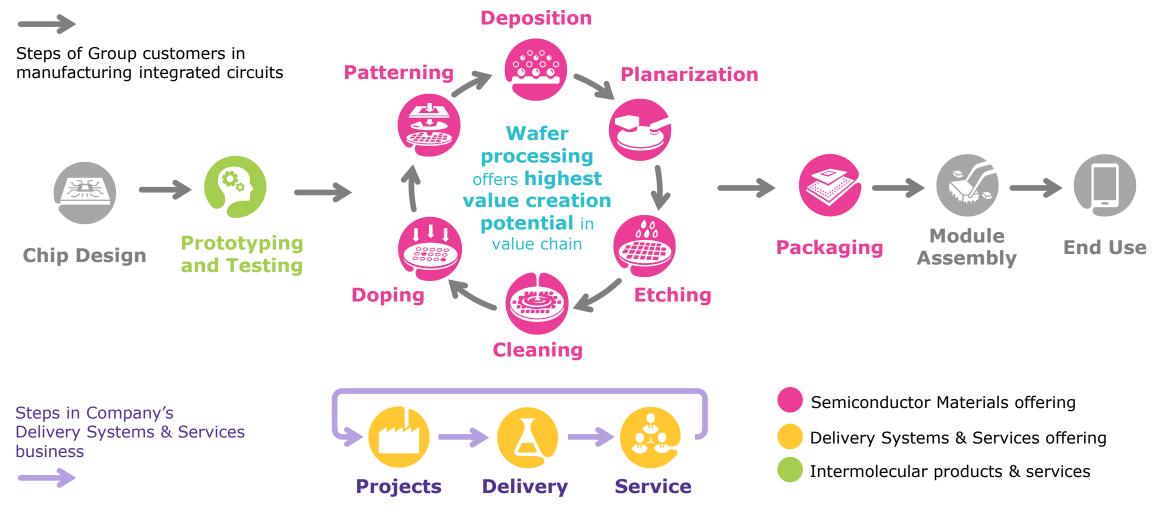
systems

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



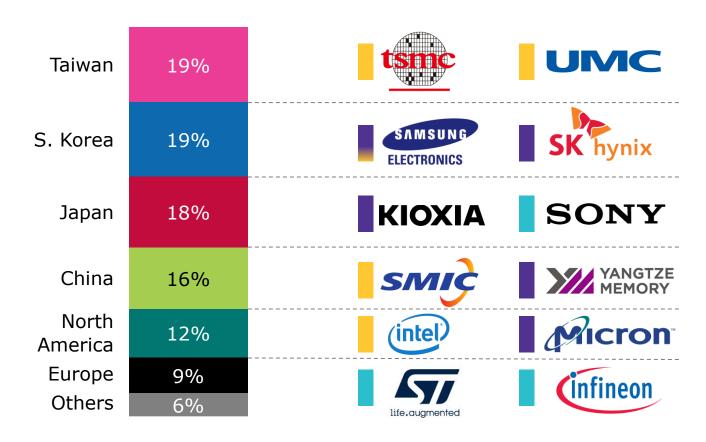
llustrative Industry P&Ls based on Sources: ¹Prismark 2021, ²Prismark 2021 & WSTS/SIA & SEMI Q1 2020; ³Representative player in the industry, non-exclusive list, not based on any underlying criteria; ⁴e.g. Filters, Pads, etc.; CPU = Central Processing Unit; RAM = Random Access Memory; SSD = Solid State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



customers

supplying all top 10 chip makers and virtually all of the top 100^3

Focus on other chips

Semiconductor Solutions has

Focus on logic chips

Focus on memory chips

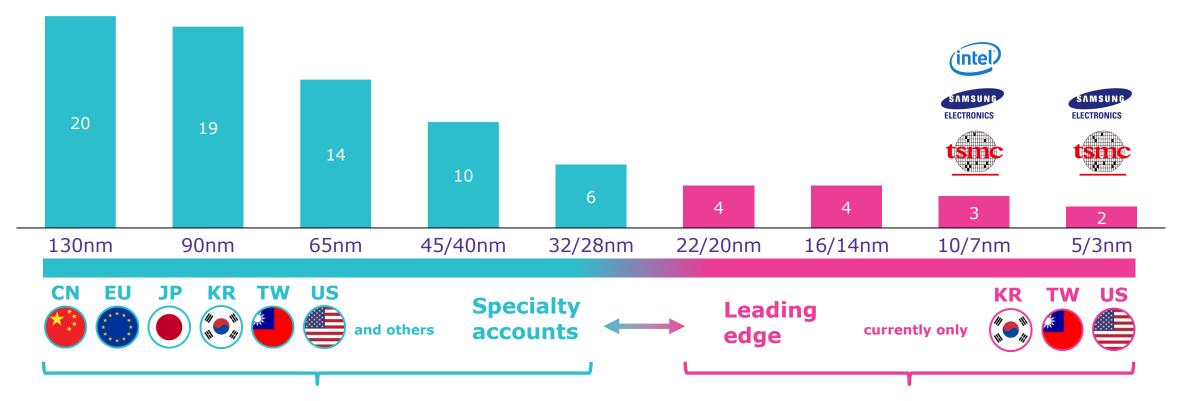
lerlying criteria ³Based on H1 2020 Sales

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

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

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

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

Data created worldwide

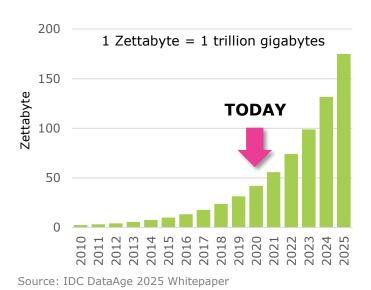
is growing +30% annually

All segments of data application are affected by global data growth

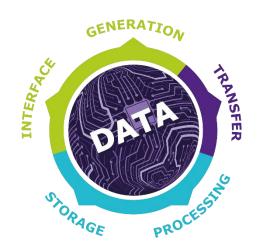
Technology trends strongly impact relevance of data application segments

Technology market growth - examples

Size of global data sphere









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

"Level Up" Scale - Expanding in synch with customer demand

A partner of choice, in step for higher demand

Semiconductor industry CAPEX p.a.¹ [bn€] Investing in lock-step with customers' 130 130 115 investments 90 57 (+approx. 2-year ramp-up time) 2018 2022 2026 2028 2016 **Electronics CAPEX** illustrative [majority in Semi] **Further** investment depending on customer Versum Materials demands 2021 2024 2018 2026 2028

1) McClean March 2021, Company announcements; FX rate: 1.18 \$/€; 2) incl. IT

Smart localization, investing close to customers

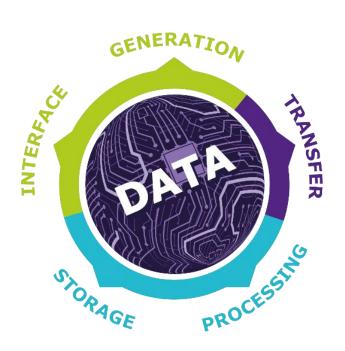
Share of semi			
material market ¹	Key investments		
Taiwan ~25%	 Additional large site ensuring supply for key customers Further expand R&D footprint 		
Korea ~25%	 Expand strong footprint, even higher customer proximity 		
China ~15%	 Become most localized multi- national ready for growth Expand production and R&D 		
USA ← ~10%	 Expand capacity for global & local supply Enable new sites of key customers 		

¹⁾ Source: Linx Consulting – only relevant segments, 2020; 2) Aggregated spend 2021-25, total spend: ~2.1bn€, thereof ~11% (~250m€) in Germany, also to fund globatograpments.

"Level Up" Tech – delivering today and enabling tomorrow

Strong portfolio of silicon & optical technologies empowering today's applications

Solutions empowering customers to make smaller, faster, more energy efficient & more sustainable devices in various applications across the entire data sphere



Data Generation

Sensors, MEMs

Data Transfer

Network chips, interconnects, antenna

Data Processing

Logic chips: CPUs, GPUs

Data Storage

Memory Chip: DRAM, 3D NAND

Data Interface

Displays: TV and mobile (LC, OLED)

And enabling tomorrow's leading-edge technologies

Selected future applications

Next-Gen image sensors

Smart Antennas

Transition to nanosheets

3D NAND with >300 layers

Foldable displays

Our contribution

Leverage semiconductor & display knowledge for new & optimized sensors

New LC, enabling beam forming, replacing bulky mechanical antennas

Novel etchants & dopants allowing higher scaling & lower power consumption

ALD¹ materials that help minimize cross talk between memory cells

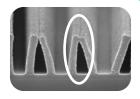
Broad set of materials from encapsulation to the OLED stack

1) Atomic Layer Deposition



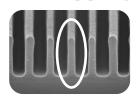
Expanding the limits of how small you can go

Pattern collapse



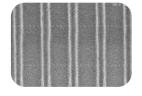


AZ® rinse materials



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

Lithography limitation



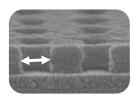


Directed self-assembly (DSA)



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

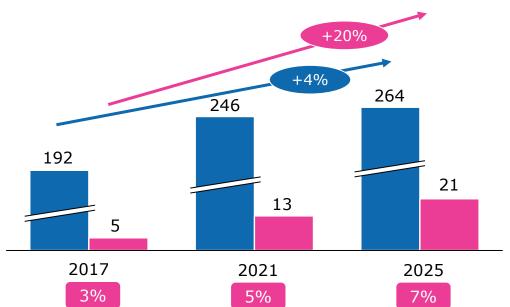


Display Solutions - OLED material market to exceed LC material

OLED shipment area / addressable material market [in % of total]

market already in 2021



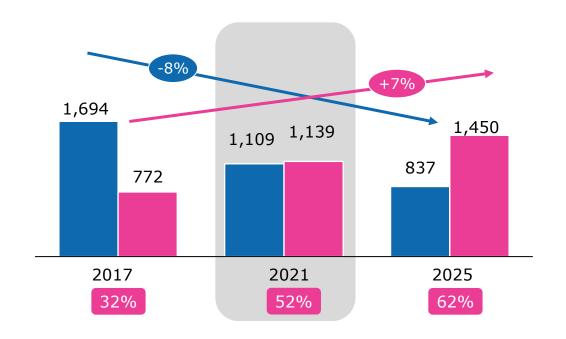


Continued growth across all technologies

 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

Strategic roadmap materializing

—Measures for a bright future ——



Darmstadt

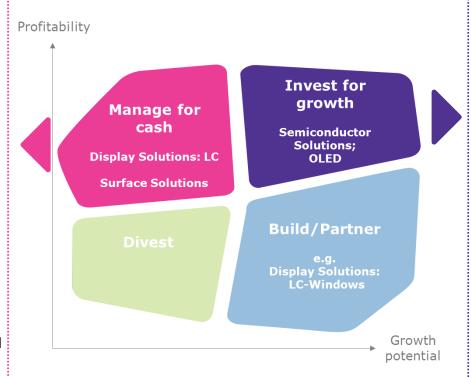
- In Darmstadt focus on R&D and production
- Immediate bottom line contribution from 2019 onwards
- Reduce the number of FTEs by ~15%
 = ~400 FTEs



 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

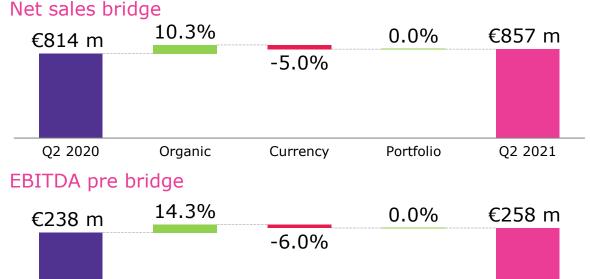


Electronics Q2: Strong performance in Semi, recovery in Surface, and stabilizing Display Solutions drive double-digit organic sales growth

Q2 2020

Electronics P&L

[€m]	IFRS		Pre	
	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

Currency

Portfolio

- 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

Organic



Q2 2021

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