



Formycon AG

The Biosimilar Experts

August 2026

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Skill, mindset and focus - our key success ingredients



Pure Play Biosimilar Company – established 2012 in Munich, Germany.

Business model contains Income from **success share and royalty streams**.



200 employees from more than 27 different countries. More than **80%** of Formycon's workforce is engaged in **R&D activities**.



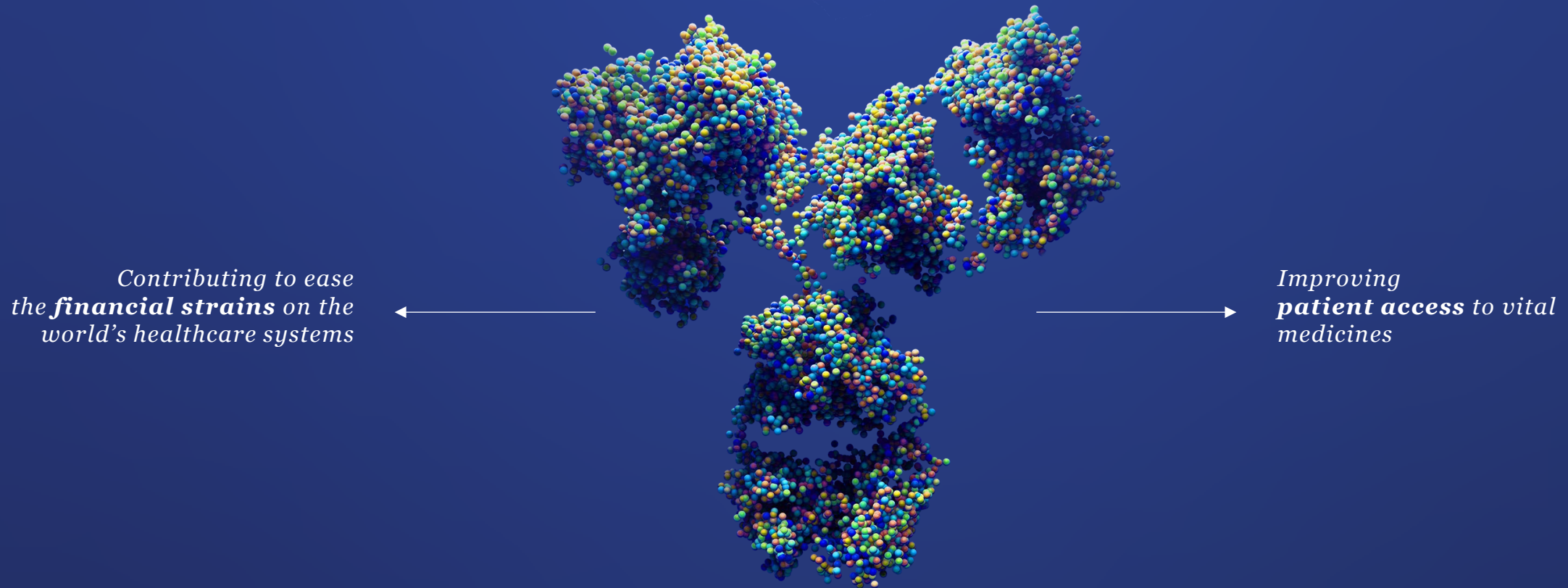
Combining high **professional expertise** in biopharmaceutical development **with agile mindset** enables Formycon to develop **multiple Biosimilar projects** in competitive timing and high quality.



Formycon's pipeline includes **three biosimilars** which are already launched in key global markets, as well as four biosimilar candidates in development.

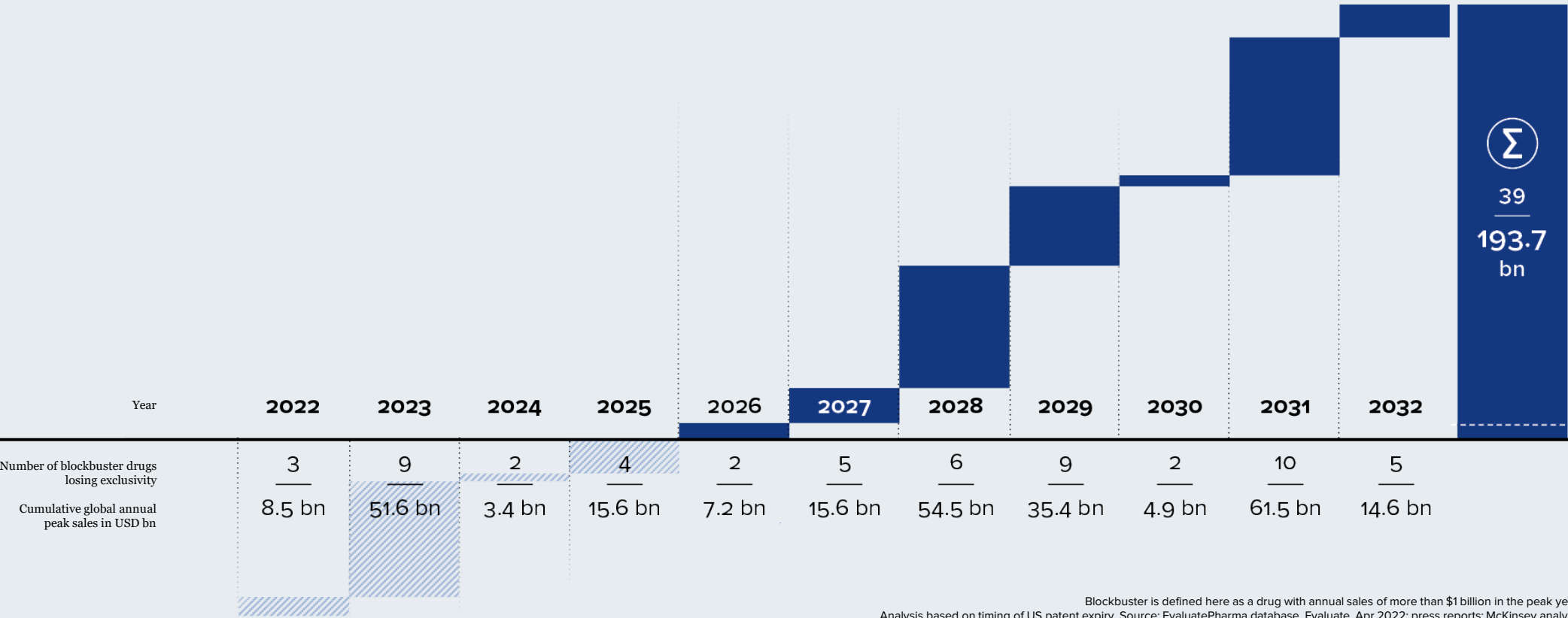
We are acting along a clear mission

Biosimilars open up enormous opportunities



Huge Biosimilar target opportunities

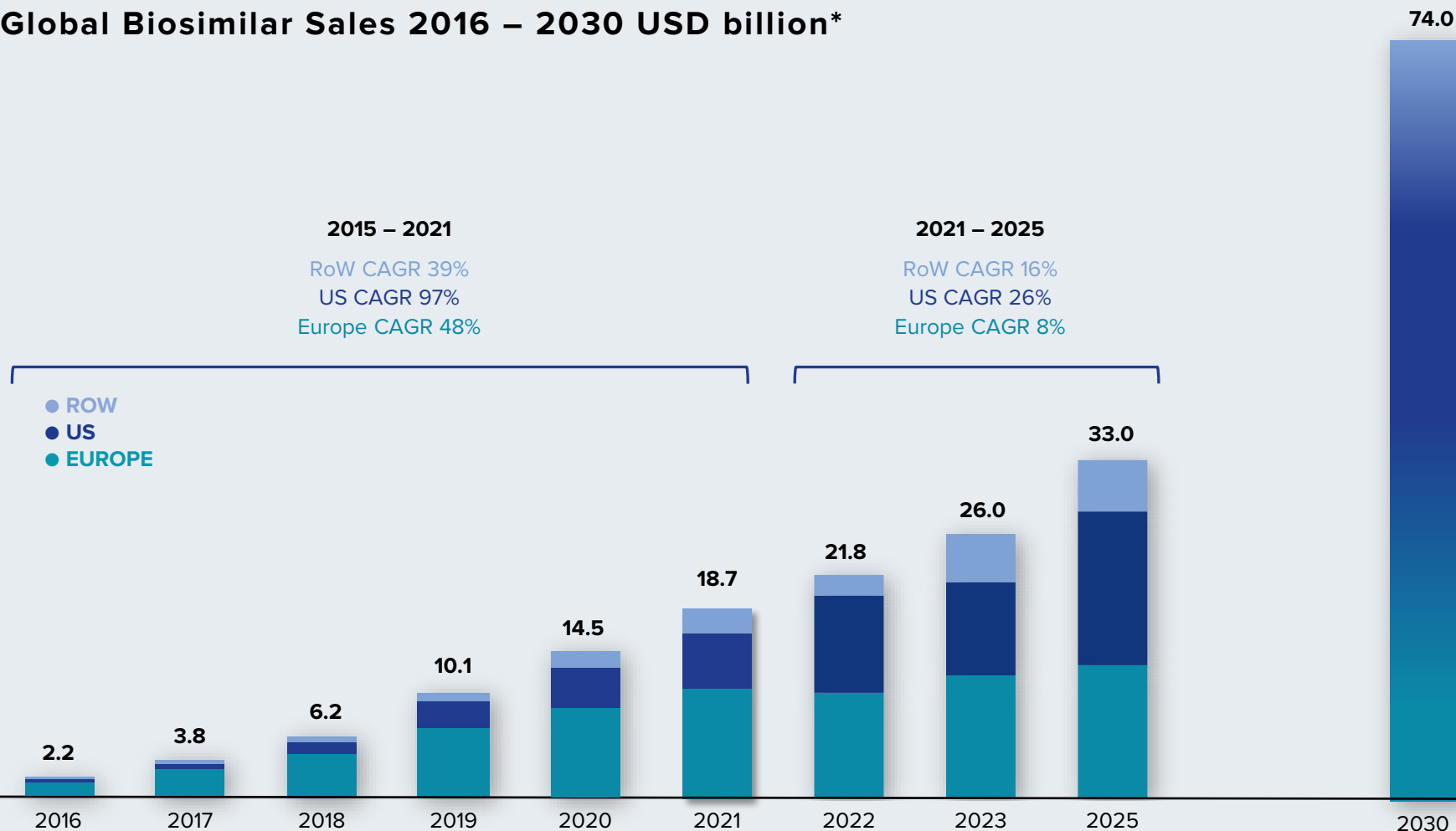
39 Blockbuster drugs with an expected global sales volume of more than 190 USD billion will lose their exclusivity in the coming years, creating huge opportunities for Biosimilars



Blockbuster is defined here as a drug with annual sales of more than \$1 billion in the peak year. Analysis based on timing of US patent expiry. Source: EvaluatePharma database, Evaluate, Apr 2022; press reports; McKinsey analysis

The Biosimilar market develops very dynamically

Global Biosimilar Sales 2016 – 2030 USD billion*



Biosimilars is the fastest growing segment in Pharma. The US market has seen the fastest growth in Biosimilars with a CAGR of 97 % from 2015 – 2021. Although projections to 2025 show a lower rate of growth, the United States is expected to stay in pole position.

* <https://www.mckinsey.com/industries/life-sciences/our-insights/three-imperatives-for-r-and-d-in-biosimilars>

#FYB4GROWTH – four strategic levers for strong competitiveness, value creation and growth



01_ Geographic Diversification

A deliberate focus on growing market opportunities beyond Europe and the U.S.



02_ Smart Portfolio

An intelligently designed and consistently executed product portfolio balancing blockbusters with niche drugs



03_ Excellence and Innovation

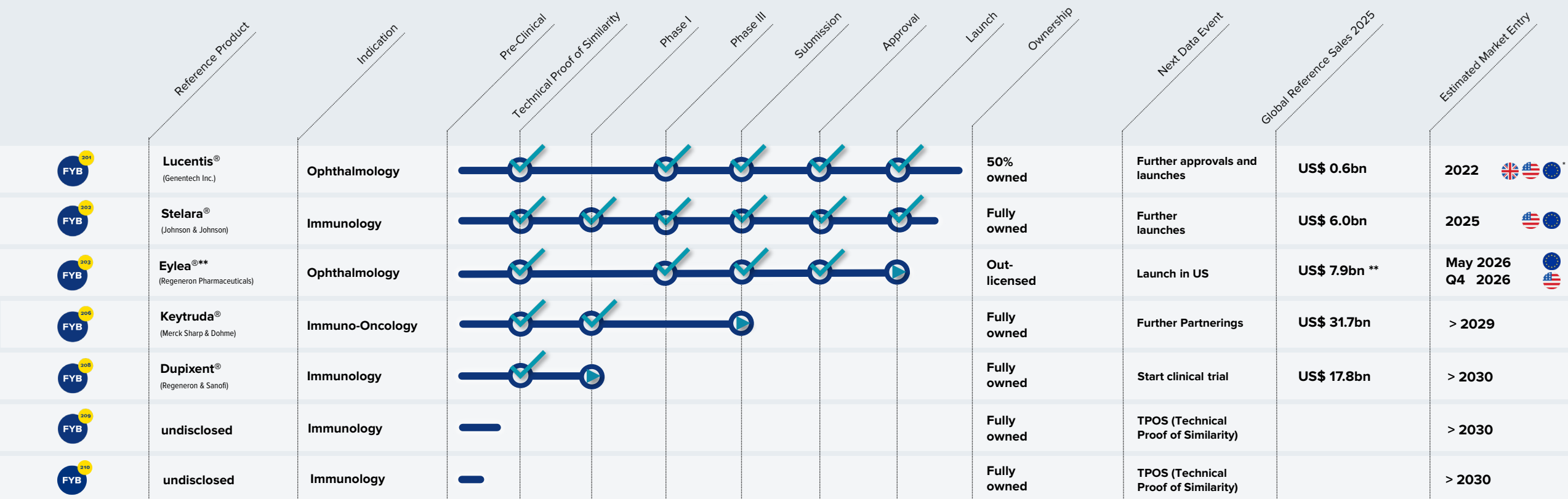
Excellence in science and execution create innovative solutions and lead to first-mover opportunities



04_ Lean Development

Cost efficient fit-for-future program combined with streamlined regulatory framework make developments leaner and faster

Diversified portfolio of commercial, late and mid stage programs



ongoing completed

* and further Regions
** Eylea® 2mg + 8mg (High-Dose) combined

#FYB4GROWTH – Good Momentum for 2026 with Key Milestones Already Achieved



Geographic Diversification

FYB201

- Brazil (BIOMM) ✓

FYB202

- Everex (LATAM) ✓

FYB206

- Lotus (APAC) ✓
- *Planned*: LATAM and further territories

FYB208

- Start of partnering activities



Smart Portfolio/ Progress to Market

FYB201

- Cimerli® Re-entry in U.S. ✓
- Nufymco® (Zydus) reimbursement code granted and phased US-launch started across multiple distribution channels ✓

FYB203

- Launched in EU ✓
- US-launch in Q4/2026

Portfolio-additions in H2 2026



Excellence & Innovation

FYB202

- Autoinjector introduced in the EU ✓

FYB206

- Clinical development successfully completed ✓
- Preparing for submission

FYB208

- Scaling up manufacturing at cost-competitive setting
- Detailing clinical strategy with agencies on Phase III waiver



Lean Development

Transformation

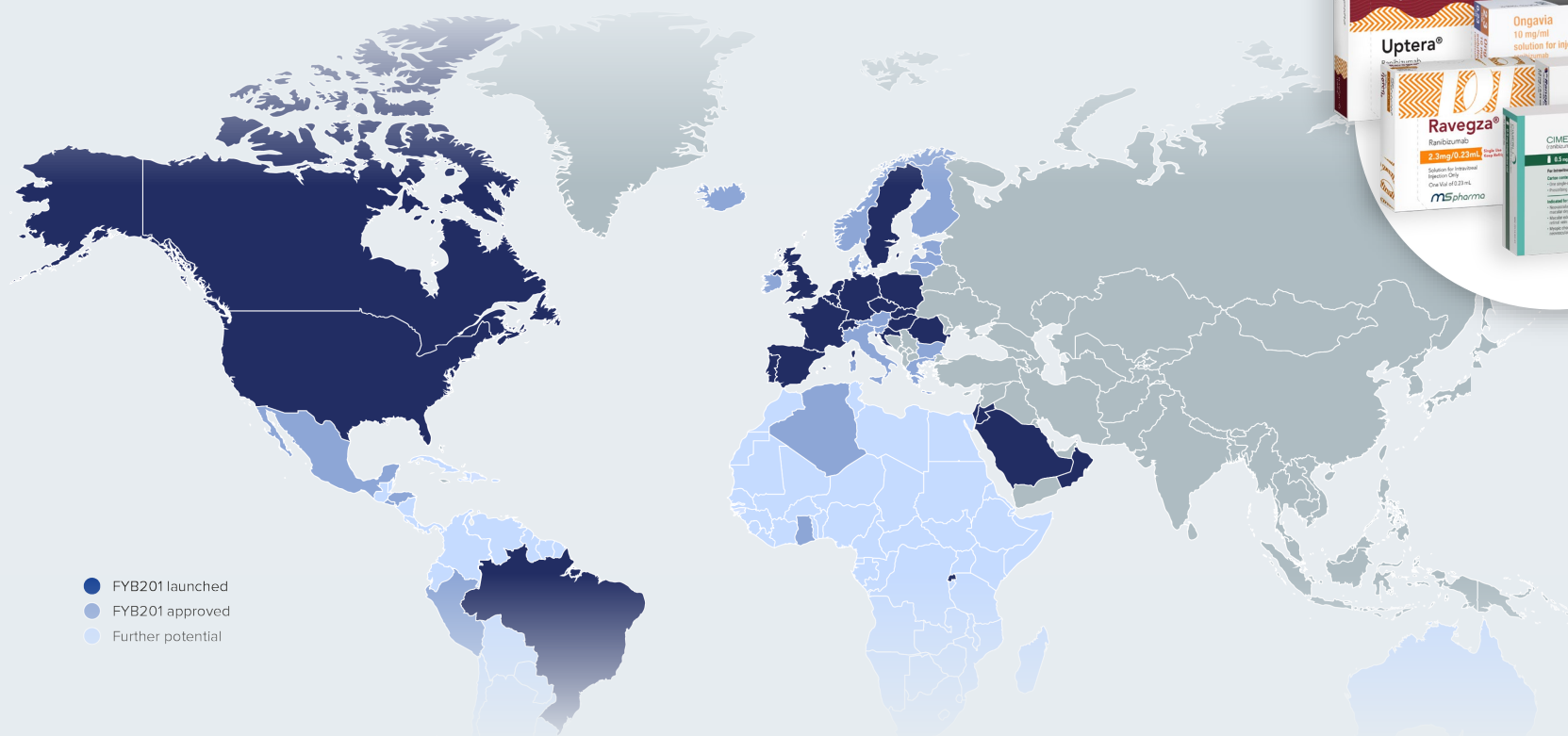
- FIT4FUTURE: Timelines and costs for new development of programs reduced by 30% ✓
- Cost-competitive CDMO partnership with OneSource Specialty Pharma (India) in place ✓
- Further streamline regulatory approaches and intensify deployment of AI



Lucentis® Biosimilar FYB201 – Strong Presence across the World



**FYB201/ranibizumab so far
launched in 24 Countries**



US, DE **SANDOZ**

US **zydus**

UK, EU, Canada **teva**

MENA **MSpharma**

Sub-Sahara **BIO USAWA**

Brazil **BIOMM**

CIMERLI

Epruvy®

Nufymco®

Raniviso

Ranopto™

Ongavia

Ravegza®

Uptera®

BioUcenta™

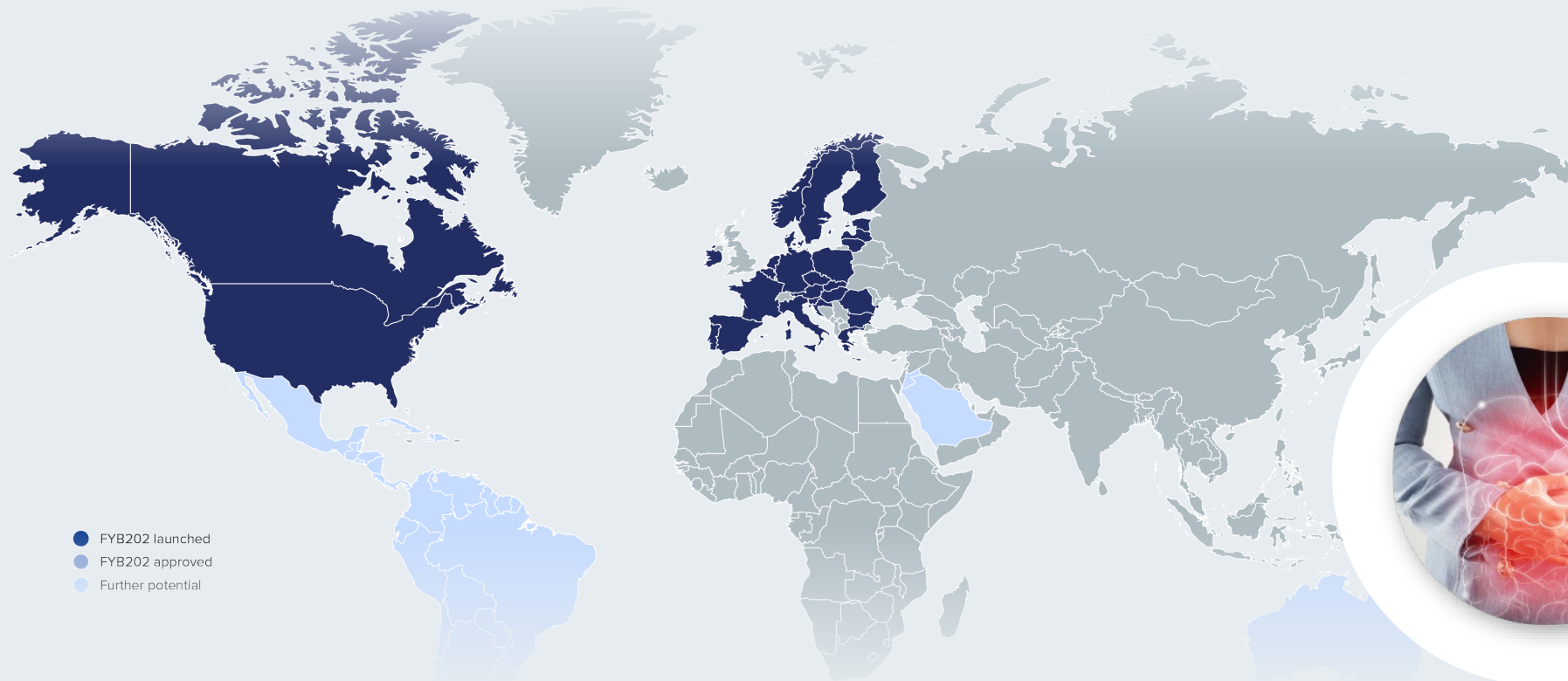
Raniviso

Lucentis® is a registered trademark of Genentech Inc.; Raniviso® is a registered trademark of Bioeq AG; Ongavia® is a registered trademark of Teva Pharmaceutical Industries Ltd.; Cimerli® is a registered trademark of Coherus BioSciences, Inc.; Ranopto™ is a trademark of Teva Canada Limited; Ravegza® and Uptera® are registered trademarks of MS Pharma; BioUcenta™ is a trademark of Bio Usawa (not approved yet)

Stelara® Biosimilar FYB202 – push for growth with Otulfi®



**FYB202/ustekinumab launched
in the US, Europe and Canada**



Global		
DE		
MENA		
Parts of LATAM-Region		

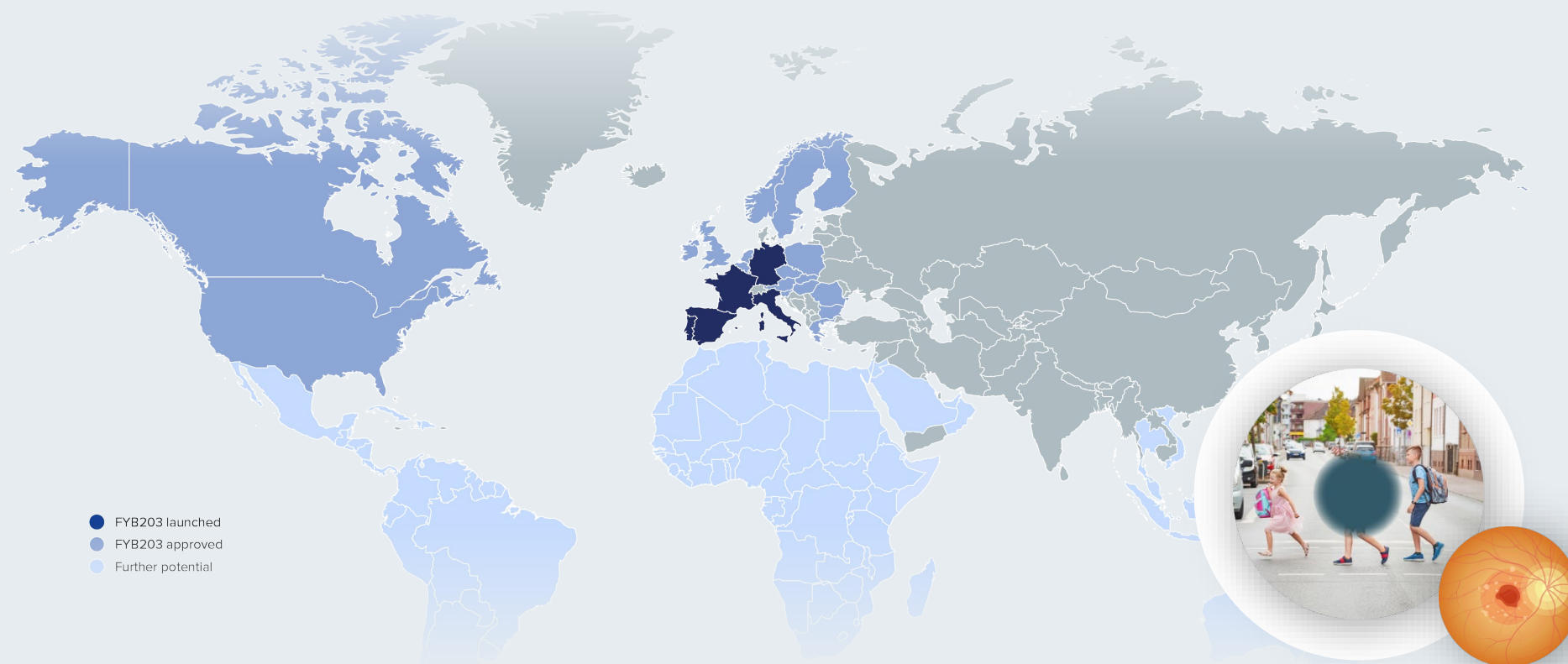


Stelara® is a registered trademark of Johnson & Johnson
Otulfi® is a registered trademark of Fresenius Kabi Deutschland GmbH in selected countries; Fymkina® is a registered trademark of Formycon AG

Eylea® Biosimilar FYB203 – launched in EU and will be launched in the US in Q4 2026



FYB203 (aflibercept) is approved in the US, EU, and UK



US, Canada		[	AHZANTIVE®
EU			
EU		[	Baiaama®
Italy			
MENA		[	Fovlya®
APAC			
LATAM			
Australia			



Eylea® is a registered trademark of Regeneron Pharmaceuticals Inc.
AHZANTIVE® and Baiaama® are registered trademarks of Klinge Biopharma GmbH; Fovlya® is a registered trademark of MS Pharma

FYB206 – Keytruda® Biosimilar – Striking several important partnerships



FYB206

Targeted Reference Indications

Immuno-oncology: Melanoma (black skin cancer), non-small cell Lung Cancer, classical Hodgkin's Lymphoma and other Tumor Diseases

Target Market 2025

USD 31.7 billion

Project Rights

100% of project and commercialization rights

Achievements and next important Milestones

- First commercialization partnership concluded with MS Pharma for the MENA region, for US and Canada with Zydus and for parts of APAC region with Lotus, further regional partnerships expected in due time
- Patient enrollment for clinical development completed (Last Patient-In) → **Dahlia PK Study was successfully completed in July 2026**
- Streamlined clinical strategy aligned with FDA in Q1/2025
- Market Launch in the United States and the EU after loss of exclusivity of the reference drug – expected after 2029

US, Canada



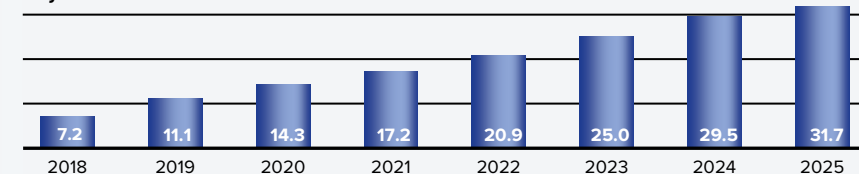
MENA



APAC



Keytruda® Sales in USD billion



*www.custommarketinsights.com/report/keytruda-market/
Keytruda® is a registered trademark of Merck Sharp & Dohme LLC

FYB208 – Dupixent® Biosimilar Candidate successfully achieved TPoS



FYB208

Targeted Reference Indications

Immunology: Moderate to severe atopic dermatitis, severe asthma, chronic rhinosinusitis with nasal polyps, chronic obstructive pulmonary disease (COPD)

Target Market 2025

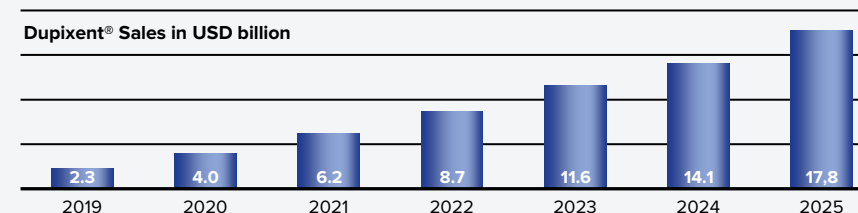
USD 17.8 billion

Project Rights

100% of project and commercialization rights

Achievements and next important Milestones

- Technical Proof of Similarity (TPoS) achieved in Q3 2025
- Manufacturing, Development of Clinical (PK) Study Design / Start of early Clinical Development Activities
- Start of Partnering Activities in 2026



Dupixent® is a registered trademark of Sanofi Biotechnology

2026 Outlook – Guidance Confirmed



Guidance 2026	Revenue 	EBITDA 	Adjusted EBITDA 	Working Capital 	Guidance 2026
	60 to 70 € million	0 to 10 € million	5 to 15 € million	20 to 30 € million	- FY 2026 guidance unchanged across all key financial KPIs - H1 revenue growth supported by increasing milestone payments and royalties - FYB206 development milestones and FYB202 royalties expected to be the principal revenue contributors in H2 - Continued FYB202 market penetration and acceleration remains important for full-year revenue performance - Focus on disciplined execution of development and commercial activities - Going Concern confirmed
H1 2026	Revenue	EBITDA	Adjusted EBITDA	Working Capital	
	25.8 € million	-3.4 € million	-6.9 € million	51.2 € million	
YE 2025	Revenue	EBITDA	Adjusted EBITDA	Working Capital	
	44.5 € million	-3.6 € million	-2.3 € million	70.1 € million	

Formycon – stable Anchor Investors

Bond:

- ISIN / WKN: NO0013586024 / A4DFJH
Volume of 70m EUR / Term is 4 years (final payback in July 2029)

Stock market:

Market Segment: Frankfurt Stock Exchange
Regulated Market (Prime Standard)

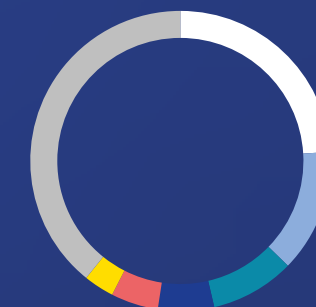
- **ISIN / WKN:** DE000A1EWVY8 / A1EWVY
- **Uplisted to Prime Standard on Nov. 12, 2024**
- **Registered capital:** € 17,672,927
Shares outstanding: 17,672,927 (w/o par value)
- **Market price / Market capitalization:** ~ € 350 million

Trading volume / Average share liquidity:

- **Q1/2026:** 29,741 shares/day
- **Q1/2025:** 82,491 shares/day

Shareholder Structure

- ~ 24 % Santo Holding (Deutschland) GmbH
- ~ 13 % Wpart GmbH, Wen.Co Invest GmbH, Peter Wendeln
- ~ 9 % Gedeon Richter
- ~ 6 % Active Ownership
- ~ 5 % Detlef & Ursula Spruth
- ~ 3 % Stefan R.
- ~ 40 % Free Float**



**per definition of Deutsche Börse

Research coverage:

- | | | | |
|--------------------|------------|------------------------|-------------------|
| – Berenberg | <i>Buy</i> | – mwb Research | <i>Buy</i> |
| – First Berlin | <i>Buy</i> | – Oddo BHF | <i>Neutral</i> |
| – HC Wainwright | <i>Buy</i> | – Royal Bank of Canada | <i>Outperform</i> |
| – Jefferies | <i>Buy</i> | | |
| – Kepler Cheuvreux | <i>Buy</i> | | |

Fully focused Pure-Play Biosimilar Developer



WE CREATED
a strong Platform with
track record



WE HAVE all ingredients to
successfully fulfill our
mission



WE ACT in a highly
attractive market



WE ARE entering the next
stage of the Formycon
Growth Story

Formycon AG



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