

Safe Harbor Statement

This presentation contains forward-looking statements that are based on the current expectations and beliefs of Amgen. All statements, other than statements of historical fact, are statements that could be deemed forward-looking statements, including any statements on the outcome, benefits and synergies of collaborations, or potential collaborations, with any other company (including BeiGene, Ltd., Kyowa-Kirin Co., Ltd., or any collaboration to manufacture therapeutic antibodies against COVID-19), the performance of Otezla® (apremilast) (including anticipated Otezla sales growth and the timing of non-GAAP EPS accretion), the Five Prime Therapeutics, Inc. acquisition, the Teneobio, Inc. acquisition, the ChemoCentryx, Inc. acquisition, or the proposed acquisition of Horizon Therapeutics plc, as well as estimates of revenues, operating margins, capital expenditures, cash, other financial metrics, expected legal, arbitration, political, regulatory or clinical results or practices, customer and prescriber patterns or practices, reimbursement activities and outcomes, effects of pandemics or other widespread health problems such as the ongoing COVID-19 pandemic on our business, outcomes, progress, and other such estimates and results. Forward-looking statements involve significant risks and uncertainties, including those discussed below and more fully described in the Securities and Exchange Commission reports filed by Amgen, including our most recent annual report on Form 10-K and any subsequent periodic reports on Form 10-Q and current reports on Form 8-K. Unless otherwise noted, Amgen is providing this information as of the date of this presentation and does not undertake any obligation to update any forward-looking statements contained in this document as a result of new information, future events or otherwise.

No forward-looking statement can be guaranteed and actual results may differ materially from those we project. Our results may be affected by our ability to successfully market both new and existing products domestically and internationally, clinical and regulatory developments involving current and future products, sales growth of recently launched products, competition from other products including biosimilars, difficulties or delays in manufacturing our products and global economic conditions. In addition, sales of our products are affected by pricing pressure, political and public scrutiny and reimbursement policies imposed by third-party payers, including governments, private insurance plans and managed care providers and may be affected by regulatory, clinical and guideline developments and domestic and international trends toward managed care and healthcare cost containment. Furthermore, our research, testing, pricing, marketing and other operations are subject to extensive regulation by domestic and foreign government regulatory authorities. We or others could identify safety, side effects or manufacturing problems with our products, including our devices, after they are on the market. Our business may be impacted by government investigations, litigation and product liability claims. In addition, our business may be impacted by the adoption of new tax legislation or exposure to additional tax liabilities. If we fail to meet the compliance obligations in the corporate integrity agreement between us and the U.S. government, we could become subject to significant sanctions. Further, while we routinely obtain patents for our products and technology, the protection offered by our patents and patent applications may be challenged, invalidated or circumvented by our competitors, or we may fail to prevail in present and future intellectual property litigation. We perform a substantial amount of our commercial manufacturing activities at a few key facilities, including in Puerto Rico, and also depend on third parties for a portion of our manufacturing activities, and limits on supply may constrain sales of certain of our current products and product candidate development. An outbreak of disease or similar public health threat, such as COVID-19, and the public and governmental effort to mitigate against the spread of such disease, could have a significant adverse effect on the supply of materials for our manufacturing activities, the distribution of our products, the commercialization of our product candidates, and our clinical trial operations, and any such events may have a material adverse effect on our product development, product sales, business and results of operations. We rely on collaborations with third parties for the development of some of our product candidates and for the commercialization and sales of some of our commercial products. In addition, we compete with other companies with respect to many of our marketed products as well as for the discovery and development of new products. Discovery or identification of new product candidates or development of new indications for existing products cannot be guaranteed and movement from concept to product is uncertain; consequently, there can be no quarantee that any particular product candidate or development of a new indication for an existing product will be successful and become a commercial product. Further, some raw materials, medical devices and component parts for our products are supplied by sole third-party suppliers. Certain of our distributors, customers and payers have substantial purchasing leverage in their dealings with us. The discovery of significant problems with a product similar to one of our products that implicate an entire class of products could have a material adverse effect on sales of the affected products and on our business and results of operations. Our efforts to collaborate with or acquire other companies, products or technology, and to integrate the operations of companies or to support the products or technology we have acquired, may not be successful. A breakdown, cyberattack or information security breach could compromise the confidentiality, integrity and availability of our systems and our data. Our stock price is volatile and may be affected by a number of events. Our business and operations may be negatively affected by the failure, or perceived failure, of achieving our environmental, social and governance objectives. The effects of global climate change and related natural disasters could negatively affect our business and operations. Global economic conditions may magnify certain risks that affect our business performance could affect or limit the ability of our Board of Directors to declare a dividend or our ability to pay a dividend or repurchase our common stock. We may not be able to access the capital and credit markets on terms that are favorable to us, or at all.

The directors of Amgen accept responsibility for the information contained in this presentation. To the best of the knowledge and belief of the directors (who have taken all reasonable care to ensure that such is the case), the information contained in this presentation is in accordance with the facts and does not omit anything likely to affect the import of such information.



AGENDA

1	Introduction	Arvind Sood		
2	Overview & Strategic Rationale	Bob Bradway		
3	R&D Overview	Dave Reese		
4	Commercial Overview	Murdo Gordon		
5	Financial Overview Peter Griffith			
6	Q&A	Bob Bradway, Peter Griffith, Murdo Gordon and Dave Reese		

TRANSACTION HAS COMPELLING STRATEGIC AND FINANCIAL RATIONALE

Complementary fit

 Strengthens Amgen's portfolio of first-in-class / best-in-class innovative therapeutics by adding complementary Horizon medicines addressing the needs of patients suffering from rare or orphan diseases

Growth acceleration

 Leverages Amgen's decades of commercial and medical leadership in inflammation and nephrology and global scale to maximize growth potential of Horizon products

Global capabilities

 Amgen's R&D and biologics manufacturing capabilities add value to Horizon's portfolio

Attractive financial profile

- Robust combined cash flow enables sustained investment in innovation and growing dividend
- Accelerates revenue growth; accretive to non-GAAP earnings from 2024

Substantial value creation for shareholders of both companies



HORIZON THERAPEUTICS OVERVIEW AND KEY DEAL TERMS



- Amongst fastest-growing biotechnology companies
- Focused on rare, autoimmune and severe inflammatory diseases
- ~2,000 global employees passionately focused on the needs of patients
- FY 2021 revenue of \$3.2B (47% YoY growth)
- Attractive biologic franchises include:







Key Deal Terms

- \$116.50 per share in cash (~20% premium to December 9 closing price)
- Transaction equity value of \$27.8B
- Closing expected in first half of 2023, subject to receipt of Horizon shareholder approval and customary regulatory approvals

Substantial value creation for shareholders of both companies

HORIZON'S PORTFOLIO OF FIRST- AND BEST-IN-CLASS BIOLOGIC THERAPIES ADDRESS GRIEVOUS ILLNESS



Thyroid eye disease (TED)

- Rare autoimmune condition associated with Graves disease results in eye bulging, double vision, inflammation and pain
- Monoclonal antibody targets insulin-like growth factor 1 receptor, which blocks signaling through the TSHR and IGF-1R complex





Chronic refractory gout

- Debilitating disease driven by elevated serum uric acid levels; patients seen by rheumatologists
- Pegylated uricase enzyme that facilitates rapid depletion of serum uric acid, leading to resolution of tophi and other gout-associated complications





Neuromyelitis Optica Spectrum Disorder (NMOSD)

- Severe autoimmune condition that can lead to blindness, paralysis and death
- Humanized, anti-CD19 B-cell depleting monoclonal antibody that suppresses B-cell related autoimmune activity





HORIZON'S PIPELINE HAS MULTIPLE OPPORTUNITIES IN INFLAMMATION

Program	Indication	Ph 1	Ph 2	Ph 3
	Myasthenia Gravis			
UPLIZNA®	IgG4-Related Disease			
	Systemic Lupus Erythematosus (SLE)			
	Alopecia Areata (AA)			
Daxdilimab	Discoid Lupus Erythematosus (DLE)			
	Lupus Nephritis (LN)			
	Dermatomyositis (DM)			
	Sjogren's Syndrome			
Dagadalihan	Rheumatoid Arthritis			
Dazodalibep	Kidney Transplant Rejection			
	Focal Segmental Glomerulosclerosis (FSGS)			
HZN-825	Diffuse Cutaneous Systemic Sclerosis (dcSSc)			
ΠZIN-023	Idiopathic Pulmonary Fibrosis (IPF)			
HZN-1116	Autoimmune Diseases			
				4N4CEN!

AMGEN'S GLOBAL R&D AND BIOLOGICS CAPABILITIES WILL ADD VALUE TO HORIZON'S PORTFOLIO



ACQUISITION CAPITALIZES ON AMGEN'S DECADES OF LEADERSHIP IN INFLAMMATION AND NEPHROLOGY

















ustekinumab

(biosimilar to STELARA®)

aflibercept (biosimilar to EYLEA®) eculizumab

(biosimilar to SOLIRIS®)

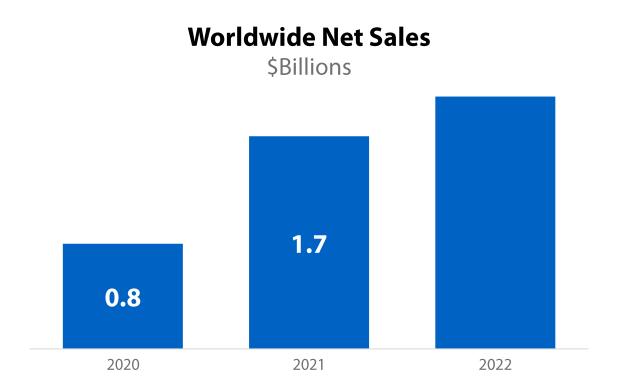
Combined portfolio growth will be strengthened by Amgen's global presence and commercial and medical capabilities in inflammation and nephrology

STELARA* is a registered trademark of Janssen Pharmaceutica NV; EYLEA* is a registered trademark of Regeneron Pharmaceuticals, Inc.; SOLIRIS* is a registered trademark of Alexion Pharmaceuticals, Inc.



TEPEZZA® IS A HIGHLY EFFECTIVE THERAPY FOR PATIENTS WITH THYROID EYE DISEASE





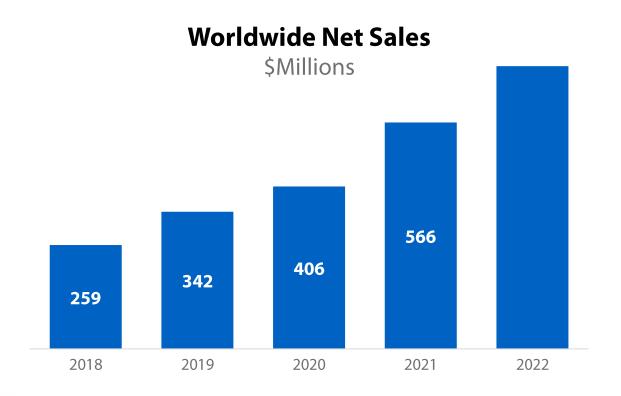
- YTD Q3 '22 Sales of \$1.5B (37% YoY)
- Currently approved in U.S. for treatment of thyroid eye disease (TED)
- OPTIC-J trial underway to support potential Japan approval
- Chronic / low Clinical Activity Score (CAS)
 TED data expected in 2023

TEPEZZA's growth will be strengthened by Amgen's global presence and commercial and medical capabilities in inflammation



KRYSTEXXA® CONTINUES TO REACH MORE PATIENTS WITH CHRONIC REFRACTORY GOUT



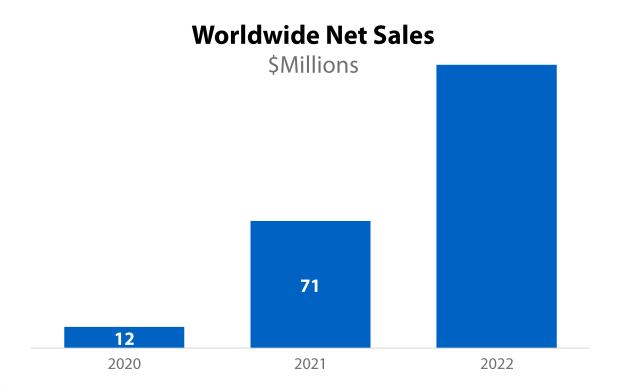


- YTD Q3 '22 Sales of \$0.5B (27% YoY)
- Currently approved in U.S. for treatment of chronic refractory gout
- Launched KRYSTEXXA + methotrexate campaign in 2022, following FDA approval of expanded label
- Opportunity for continued penetration of currently indicated patient population

KRYSTEXXA's uptake will benefit from Amgen's 20-year history with rheumatologists







- YTD Q3 '22 Sales of \$113M (148% YoY)
- Currently approved in U.S. and Europe for NMOSD
- Opportunity for expansion into Europe and other ex-U.S. markets
- Opportunity in additional indications
 - Lifecycle management studies underway in Myasthenia Gravis (MG) and IgG4-Related Disease

Note: 2020 and 2021 include \$12M and \$11M of Viela Bio, Inc. sales, respectively (UPLIZNA acquired by Horizon on 3/15/21)

Amgen's commercial and medical capabilities will accelerate awareness, adoption and global expansion of UPLIZNA

AMGEN HAS THE CAPABILITIES TO DRIVE GROWTH OF HORIZON'S PORTFOLIO



Global Scale and Reach in ~100 Countries;

Decades of Experience In Biologics, Inflammation and Nephrology



Established History With Rheumatologists and Other Inflammation Specialists



Sales Force, Access, Supply Chain, Medical and **Patient Support Capabilities**

Accelerating Growth Products and Launches



TRANSACTION HAS STRONG FINANCIAL PROFILE

Combined free cash flow (~\$10 billion in 12 months ending Q3 2022*) enables: Sustained investment in innovation Continued dividend growth Timely de-leveraging Near-Term Accretion Accelerates revenue growth; accretive to non-GAAP earnings from 2024 Annual pre-tax efficiencies of at least \$500 million by the end of the third fiscal year following transaction close

Strong Credit Profile

- Goal is to maintain a strong investment grade credit profile with leverage inline with current levels by the end of 2025
- Expect to retire > \$10 billion of debt through 2025



