

Injectables Inhalation Intranasal

Proprietary

Highly Purified Peptide/Protein

Biosimilar Interchangeable

Complex Generic Combination **Products**

J.P. Morgan Healthcare Conference January 2024

Forward Looking Statements

This presentation and the accompanying oral presentation contain forward-looking statements, of Amphastar Pharmaceuticals, Inc. ("Amphastar", "we". "our" and that are based on our management's current expectations and assumptions and on information currently available to management. Forward-looking statements include all statements other than statements of historical fact contained in this presentation, including, but not limited to, information concerning our business plans and objectives, potential growth opportunities, product development, regulatory approvals, market potential, efficiencies, competitive position, and industry environment, among other statements.

All statements in this presentation that are not historical are forward-looking statements, including, among other things, statements relating to our expectations regarding future financial performance and business trends, our future growth, sales and marketing of our products, market size and expansion, product portfolio, product development, the timing of FDA filings or approvals, including the DMFs of ANP, the timing of product launches, acquisitions and other matters related to our pipeline of product candidates, the timing and results of clinical trials, the prospective benefits of the acquisition of BAQSIMI®, and other future events, including potential contingent consideration amounts and terms related to the acquisition of BAQSIMI®, the anticipated benefits of BAQSIMI® to our product portfolio, Amphastar's commitment to strategically maximizing the commercial potential of BAQSIMI®, and other future events. These statements are not facts but rather are based on Amphastar's historical performance and our current expectations, estimates, and projections regarding our business, operations, and other similar or related factors. Words such as "may," "might," "would," "should," "anticipate," "predict," "potential," "continue," "expect," "intend," "plan," "project," "believe," "estimate," and other similar or related expressions are used to identify these forward-looking statements, although not all forward-looking statements contain these words. You should not place undue reliance on forward-looking statements because they involve known and unknown risks, uncertainties, and assumptions that are difficult or impossible to predict and, in some cases, beyond Amphastar's control. Actual results may differ materially from those in the forward-looking statements as a result of a number of factors, including those described in Amphastar's filings with the Securities and Exchange Commission, including in our Annual Report on Form 10-K for the year ended December 31, 2022, filed with the SEC on March 1, 2023, and in our Quarterly Report on Form 10-Q for the guarter ended September, 2023, filed with the SEC on November 8, 2023. In particular, there can be no guarantee that the acquisition of BAQSIMI® will be beneficial to our business, that any event, change or other circumstance could cause the results of the acquisition and integration of BAQSIMI® into our product portfolio to differ from Amphastar's expectation, that all or any of the contingent consideration will be payable on the terms described herein or at all, or that Amphastar can reliably predict the impact of BAQSIMI® on its financial results or financial guidance. You can locate these reports through our website at http://ir.amphastar.com and on the SEC's website at www.sec.gov. The forward-looking statements in this release speak only as of the date of the release. Amphastar undertakes no obligation to revise or update information or any forward-looking statements in presentation to reflect events or circumstances in the future, even if new information becomes available or if subsequent events cause our expectations to change

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Company Overview



Company Overview

One-Stop

Fully Integrated
 Business Model,
 R&D, Manufacturing
 and Distribution

Dual Strategies Growth Model

- Pipeline Development
- Strategic Acquisitions

Three-H Focus

- High Quality
- High Efficiency
- High Technology



Fully Integrated Business Model: One-Stop

- Extensive in-house product development capabilities
 - Technical Platforms
 - State-of-the-art instruments
 - Animal studies
 - Clinical research team

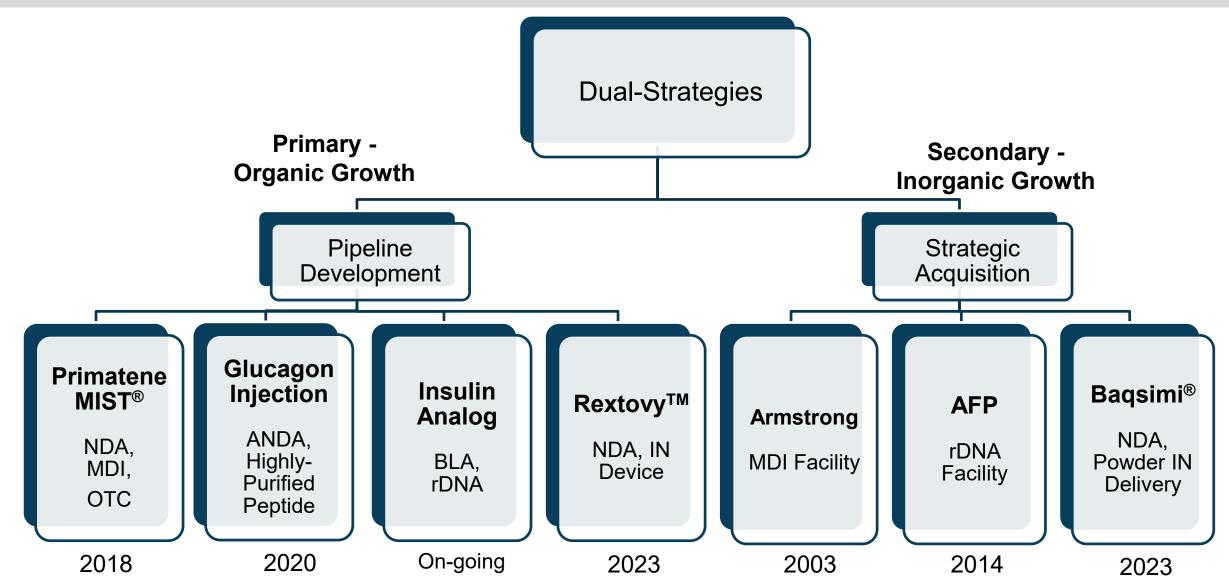
- Fully integrated back end manufacturing capabilities
 - API and key materials
 - Device and key components
- Complete front end integration
 - Marketing
 - Distribution



 Control over quality and compliance throughout the product development and manufacturing cycle



Dual-Strategies Growth Model





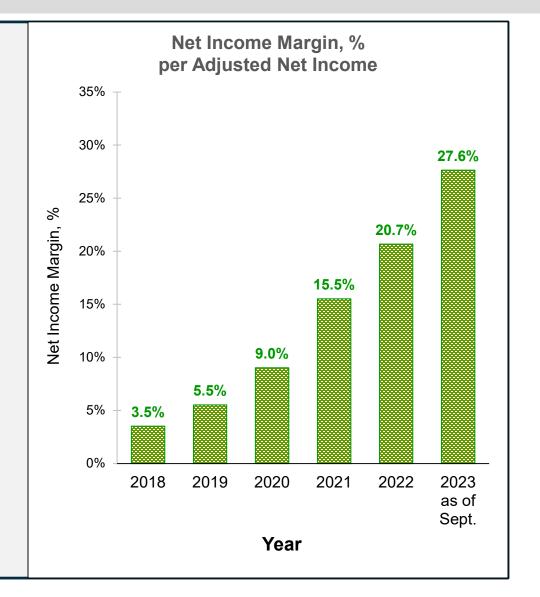
Three-H Focus

Amphastar's Management team operates the Company to:

- Insist on High Quality
- Emphasize High Efficiency and
- Rely on High Technology to Develop Pipelines

The 3-H focus results in high net income margin

(\$Million or Specified)	2018	2019	2020	2021	2022	YTD 2023-09
Revenue, x	295	322	350	438	499	466
Net Income (GAAP)	-5.7	48.9	1.4	62.1	91.4	101.4
Net Income, Adjusted, y	10.4	17.8	31.6	68.0	103.2	128.8
Net Income Margin, Adjusted, $=y/x$, %	3.5%	5.5%	9.0%	15.5%	20.7%	27.6%



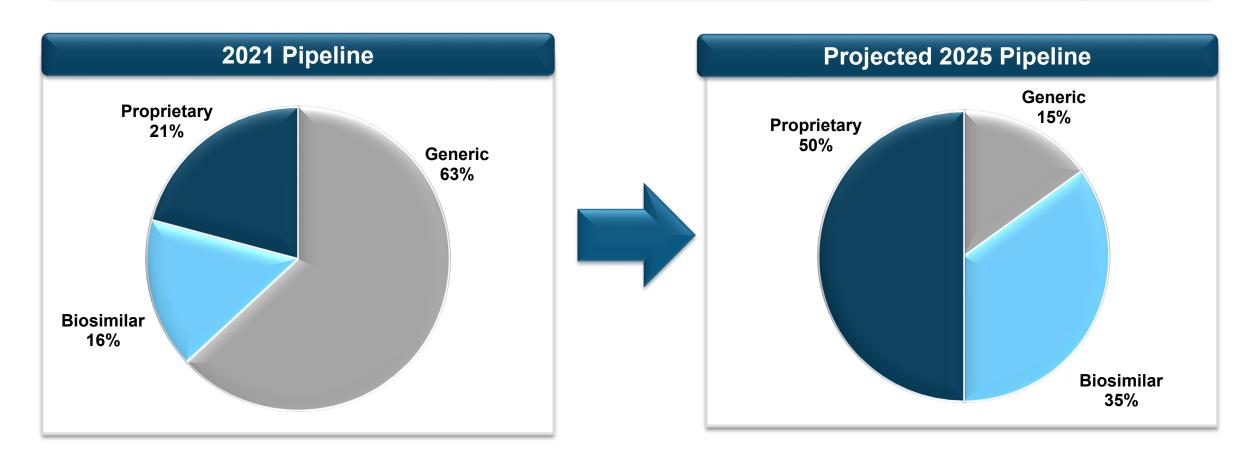


Leveraging Strategic Vision & Core Strengths



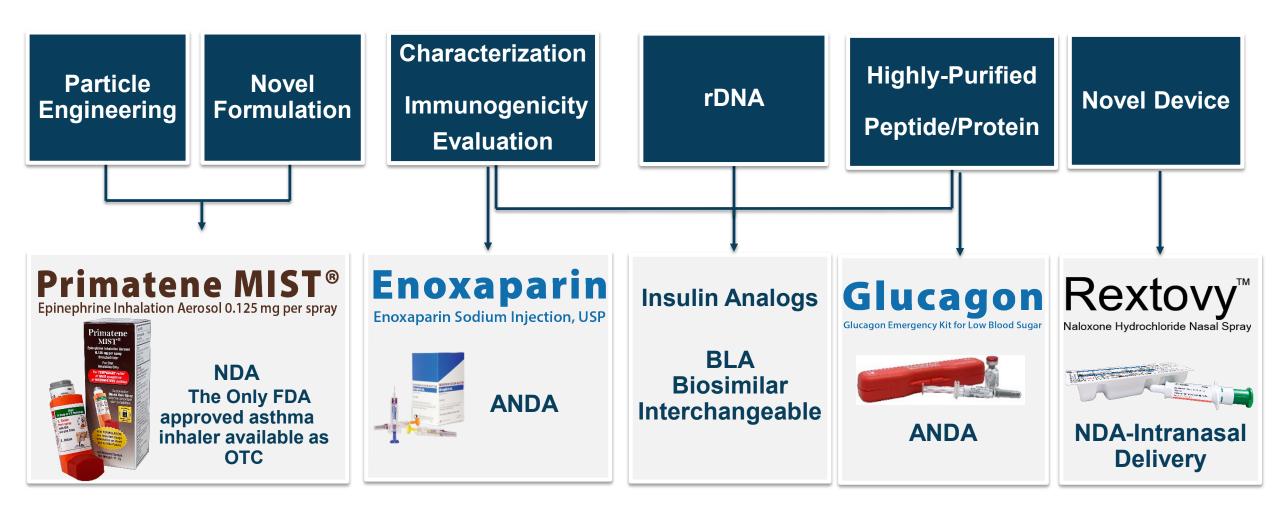
Pipeline Evolution

Amphastar's pipeline is projected to advance with a greater focus on proprietary and biosimilar products



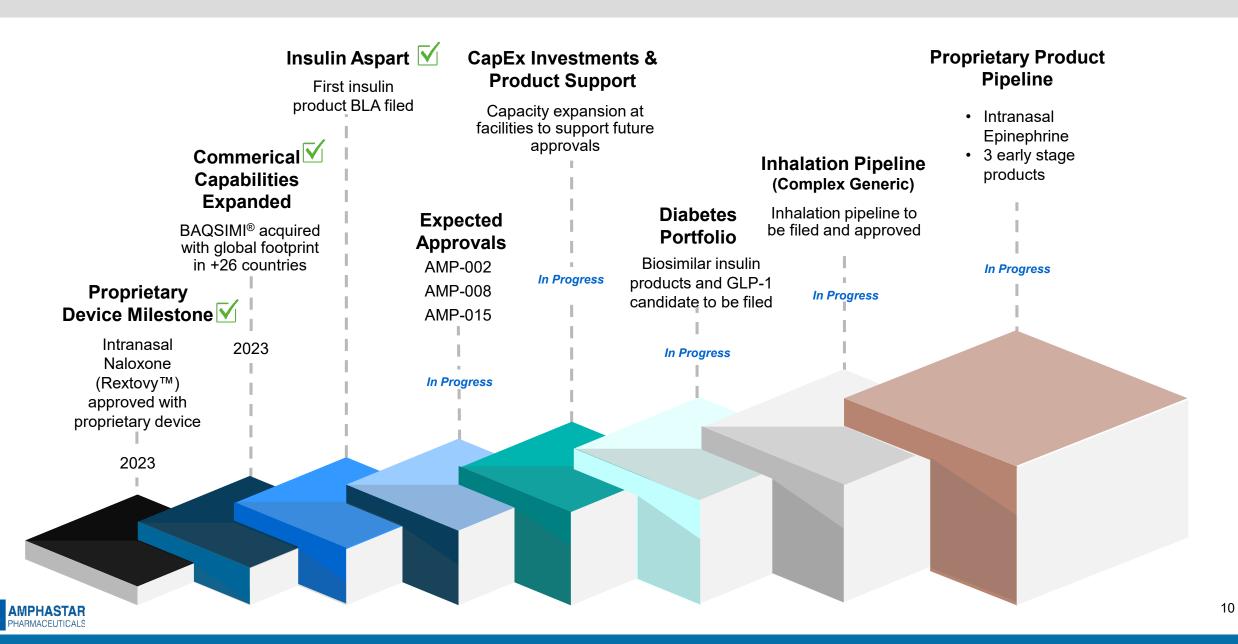


Technical Platforms





Strategic Shift Toward Proprietary & Biosimilars Drugs



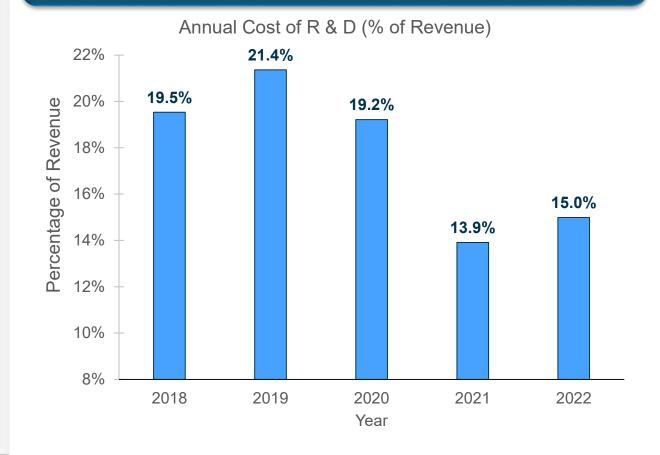
R&D and Pipeline



Focused on R&D Investment

- Strategic focus to make substantial R&D investments to expand our product portfolio
- Diverse pipeline development with flexibility and scalability for sourcing API, starting material, and research under our vertically – integrated platform
- Emphasis and investment in R&D differentiates us from our competitors as our focus is on the long-term growth of our company
- R&D from API, early stage, and clinical trials and from laboratory to scale-up

Self-funded R&D investment of approximately \$330 million in the recent 5 years





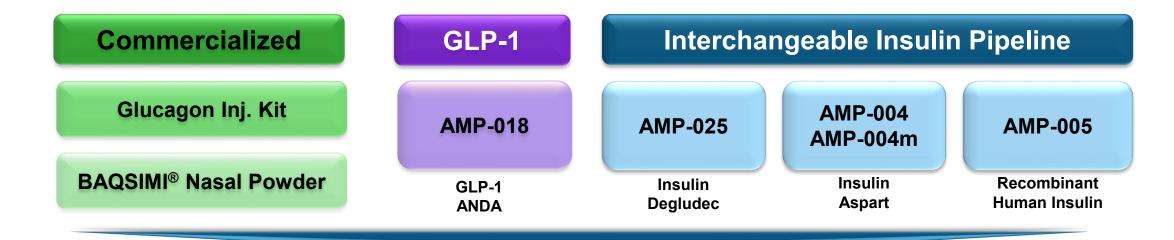
Pipeline – ANDAs with Technical Barriers

Generic Pipeline, 8 Candidates

ANDA Type	Product Code	Current Stage	*IQVIA Sales
	AMP-002	GDUFA Q2 2023, no FDA Action and no pending requests as of 1/04/24	+\$500 Million
Injectable AMP-015 (Teriparatide)		CRL received and responded; GDUFA Q1 2024	+\$600 Million
	AMP-018	GLP-1: Planned filing 2024	+\$2 Billion
	AMP-008	CRL received, and responded; GDUFA Q2 2024	-
	AMP-007	Filed, GDUFA expected Q4 2024	
Inhalation AMP-016		Stability/clinical trials	+\$6 Billion
	AMP-017	Planned filing 2024	
	AMP-022	Development	



Diabetes Portfolio



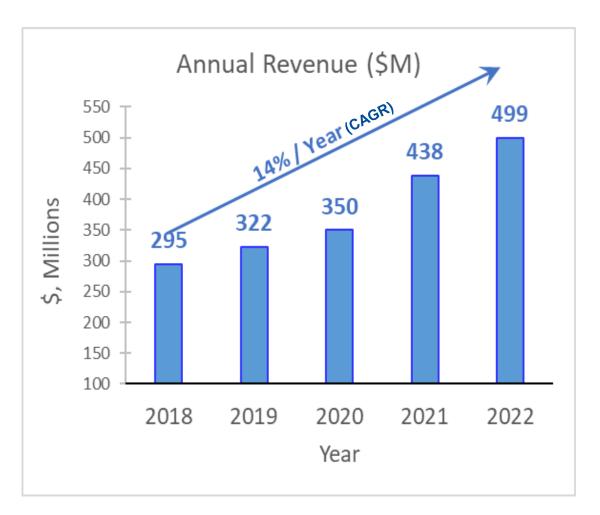
- BAQSIMI[®], the first and only FDA approved glucagon nasal powder
- First and only FDA approved generic Glucagon injection kit
- GLP-1 ANDA development
- Insulin Pipeline:
 - Covers the full spectrum of the insulin from rapid to long acting
 - AMP-004 BLA filed
 - \$8 Billion in IQVIA sales as of September 30, 2023, ~70 million of units of both pens and vials

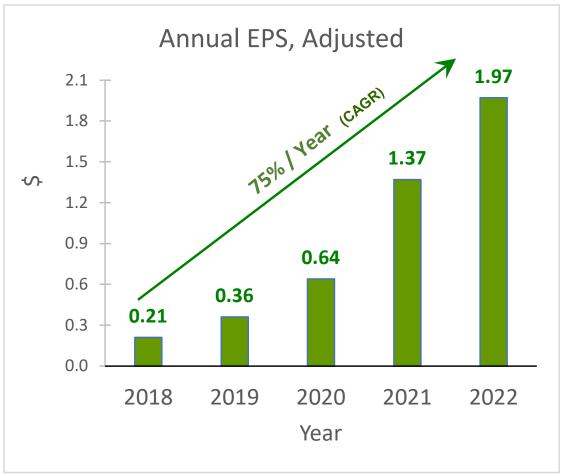


Sales and Marketing



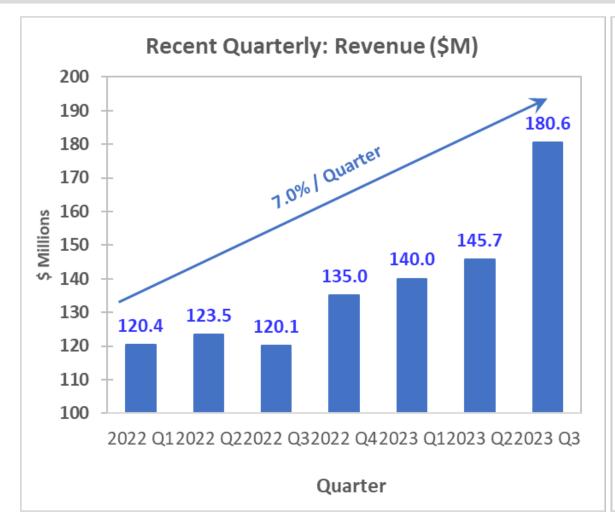
Sales and EPS Trend

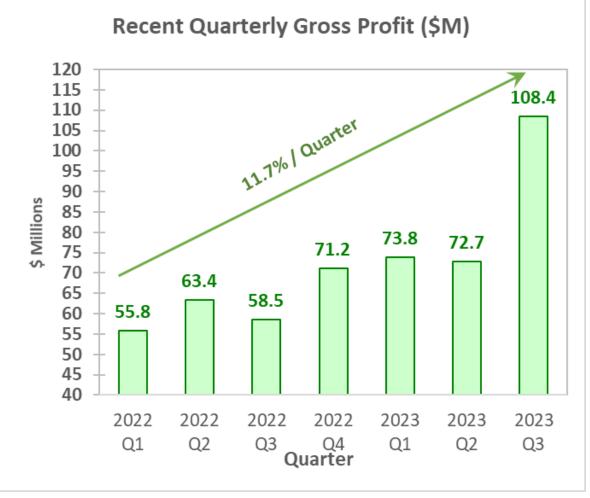






Recent Quarter Trend: Sales & Gross Profit

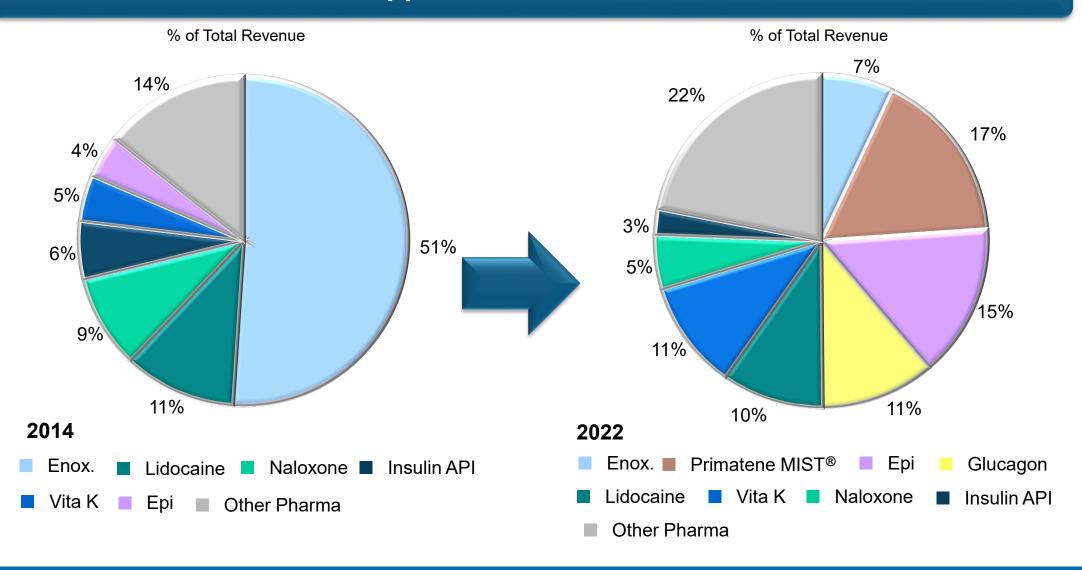






Existing Products Provide Strong Base

New Launches have Supported Diversification of the Revenue Base

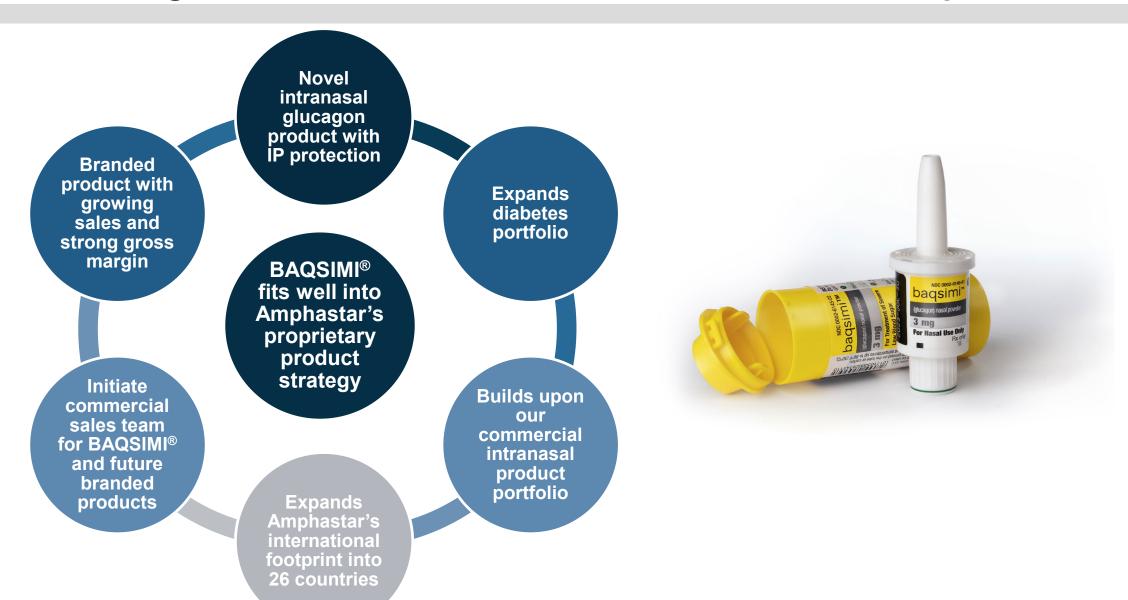




Key Proprietary Products



BAQSIMI® Strategic Rationale: A Transformative Transaction for Amphastar





BAQSIMI® Patient Impact

Glucagon is underutilized:

The American Diabetes Association (ADA) recommends that patients at increased risk for Level 2 hypoglycemia be prescribed glucagon¹

Amphastar will focus on BAQSIMI® to better serve patients

Approximately 7 million people are treated with insulin and only about 0.7 million (~10%)² of these patients currently utilize glucagon

BAQSIMI[®] is currently a category leader for ease in patient use:

Simple nasal administration:

Currently the only non-injection glucagon approved by the FDA, passively absorbed in the nose, provide lower barrier for administration than injection

Ready-to-use with no reconstitution or priming required

Portability for Consumers:

Smaller product size than other glucagon products, and wider temperature storage range than other glucagon injection product.



BAQSIMI® Forecast

Sales

- Projected to reach \$145 million to \$155 million annualized in 2023
- Net Economic Benefit (sales minus expenses) will be booked until Amphastar takes over distribution from Lilly, on a country by country basis throughout 2024
- Projected to reach peak of \$250 million to \$275 million
- Adjusted EPS⁽¹⁾
 - Project \$0.12 to \$0.18 incremental adjusted EPS in 2023
 - Project \$2.00 to \$2.50 incremental adjusted EPS at peak
 - (1) Adjusted EPS is a non-GAAP financial measure. Reconciliation to the nearest GAAP measure is unavailable
 without unreasonable efforts. Refer to the section titled "Non-GAAP Financial Measures" for an explanation of nonGAAP financial measures.





Primatene MIST®

- Primatene MIST[®], a proprietary and patent-protected over-thecounter epinephrine inhalation product
- The only FDA approved asthma inhaler available OTC, launched Dec 2018
- Multiple scientific articles were published in support of Primatene
 MIST®
- Intensive cardiovascular studies >40,000 data points
- US Adult asthma patients: 20 million per CDC*
- 17% of Amphastar sales in 2022



Highlights and Catalysts



Growth Drivers and Upcoming Milestones

Key Growth Drivers in 2024

- BAQSIMI®
 - Increased promotion
- Glucagon Injection Kit
 - Increased market opportunity
 - Increased capacity approved
- Primatene MIST[®]
 - Advertising campaign

- Expected approvals
 - AMP-002 CRL responded; GDUFA Q2 2023
 - AMP-008 CRL responded; GDUFA Q2 2024
 - AMP-015 (Teriparatide) CRL responded;
 GDUFA Q1 2024

Milestones

Recent Filings

- AMP-007 Filed
- AMP-004 Filed

Planned Filings in 2024

- Intranasal Naloxone Rx to OTC switch
- AMP-018 ANDA GLP-1
- AMP-004m BLA
- AMP-017 ANDA

