

FIRST QUARTER 2024 EARNINGS CONFERENCE CALL

MAY 2024

Today's Agenda

- Business Update
 Terrie Curran, President & Chief Executive Officer
- US Commercial Launch Progress
 Martin Gilligan, Chief Commercial Officer
- First Quarter 2024 Results

 Molly Henderson, Chief Financial & Business Officer
- Closing Remarks

 Terrie Curran, President & Chief Executive Officer
- Question & Answer
 Terrie Curran, President & Chief Executive Officer
 Martin Gilligan, Chief Commercial Officer
 Molly Handerson, Chief Financial & Business Officer

Molly Henderson, Chief Financial & Business Officer Azmi Nabulsi, Chief Operating Officer



Safe harbor

This presentation contains forward-looking statements. All statements other than statements of historical facts contained in this presentation, including statements regarding our future results of operations and financial position, anticipated milestones, anticipated cash runway, expectations of generating stability data necessary to support the proposed shelf life of vonoprazan, business strategy, prospective products, product approvals, research and development costs, timing and likelihood of success, plans and objectives of management for future operations, and future results of current and anticipated products, are forward-looking statements.

In some cases, you can identify forward-looking statements by terms such as "may," "will," "should," "expect," "plan," "anticipate," "could," "intend," "target," "project," "contemplates," "believes," "estimates," "predicts," "potential" or "continue" or the negative of these terms or other similar expressions. These statements involve known and unknown risks, uncertainties and other important factors that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. These risks, uncertainties and other factors include, without limitation: our ability successfully to commercialize approved products containing vonoprazan; our new drug application for non-erosive GERD may not be approved by the FDA; our planned clinical trials of vonoprazan as an as need treatment for non-erosive GERD or for eosinophilic esophagitis may not successfully be initiated or completed; we may not be successful in expanding commercial coverage for our approved products; the inherent risks of clinical development of vonoprazan; our dependence on third parties in connection with product manufacturing, research and preclinical and clinical testing; regulatory developments in the United States and foreign countries; unexpected adverse side effects or inadequate efficacy of vonoprazan that may limit its development, regulatory approval and/or commercialization, or may result in recalls or product liability claims; our ability to obtain and maintain intellectual property protection and non-patent regulatory exclusivity for approved products containing vonoprazan; our ability to comply with our license agreement with Takeda; our ability to achieve and maintain adequate levels of coverage and reimbursement for vonoprazan; the availability of additional funds under our revenue interest financing agreement and term loan agreement; the sufficiency of our capital to fund our operations; and other risks described in our filings with the Securities and Exchange Commission (SEC), including our most recent Annual Report on Form 10-K and any subsequent filings with the SEC. You are cautioned to place undue reliance on these forward-looking statements, which speak only as of the date hereof, and we undertake no obligation to revise or update this presentation to reflect events or circumstances after the date hereof. All forward-looking statements are qualified in their entirety by this cautionary statement, which is made under the safe harbor provisions of the Private Securities Litigation Reform Act of 1995.

This presentation also contains estimates and other statistical data made by independent parties and by us relating to market size and other data about our industry. This data involves a number of assumptions and limitations, and you are cautioned not to give undue weight to such estimates. In addition, projections, assumptions, and estimates of our future performance and the future performance of the markets in which we operate are necessarily subject to a high degree of uncertainty and risk.

This presentation contains non-GAAP financial measures, which should be considered only a supplement to, and not a substitute for or superior to, GAAP measures. Refer to the Reconciliation of Non-GAAP Financial Measures to GAAP Results table on pages 17 and 18 of this presentation.



Business Update

Terrie Curran, President & Chief Executive Officer



Focused on three core goals during the early stages of launch



Communicate clinical superiority vs. a PPI



Drive brand awareness and demand



Build access for patients



Great progress made executing on VOQUEZNA launch goals

† Clinical Superiority



This medication and my [GI] saved my life. I suffered for months and had given up. Could not eat or lie down. Was in constant pain. But now I am on the way to healing. I had coffee today for first time in months.¹

Brand Awareness



43K+
prescriptions written²

17.5K+
prescriptions filled³

... ■ Patient Access



Express Scripts National Formularies



Cigna Healthcare Formularies

~72M
commercial
lives covered4

¹ Actual VOQUEZNA patient. Results not typical.

² Unique prescriptions written; IQVIA + BlinkRx as of 4/26/24.

³ IQVIA + BlinkRx as of 4/26/24.

⁴ Per MMIT formulary lookup tool as of 5/8/2024.

Approaching potential Non-Erosive GERD launch and planned development

Prepared for an immediate launch into Non-Erosive GERD Daily Dosing

July 19, 2024 Action Date ~15M Dx & Tx Adults¹ Similar Patient Journey

Same Target Writers

Planned
Development
in 2024

As Needed treatment of heartburn associated with **Non-Erosive GERD**

- » Positive Phase 2 results
- » Planning to initiate Phase 3 trial in 2024

Treatment of **Eosinophilic Esophagitis** (**EoE**) for adult & pediatric use

» Planning to initiate Phase 2 trial in 2024





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US Commercial Launch Progress

Martin Gilligan, Chief Commercial Officer



The commercial launch of VOQUEZNA continues to build momentum





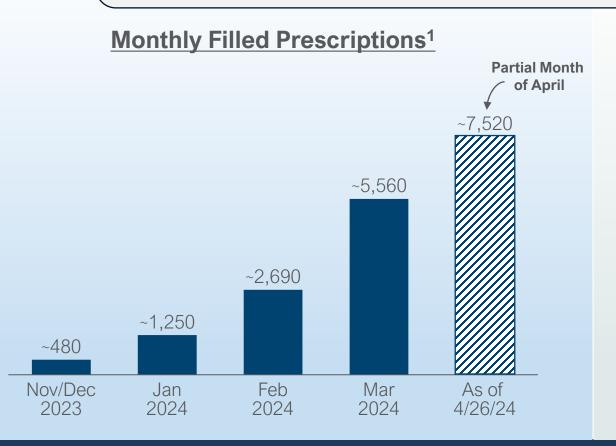


Demonstrating consistent monthly growth in key launch metrics

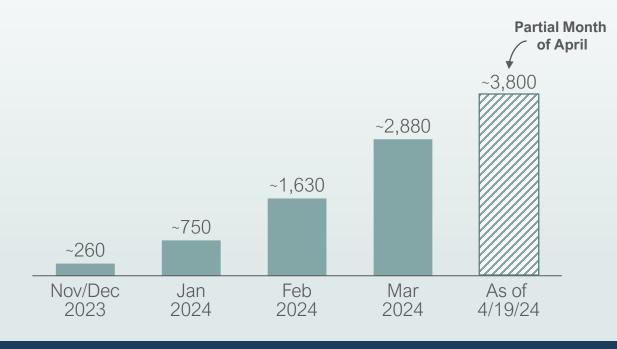






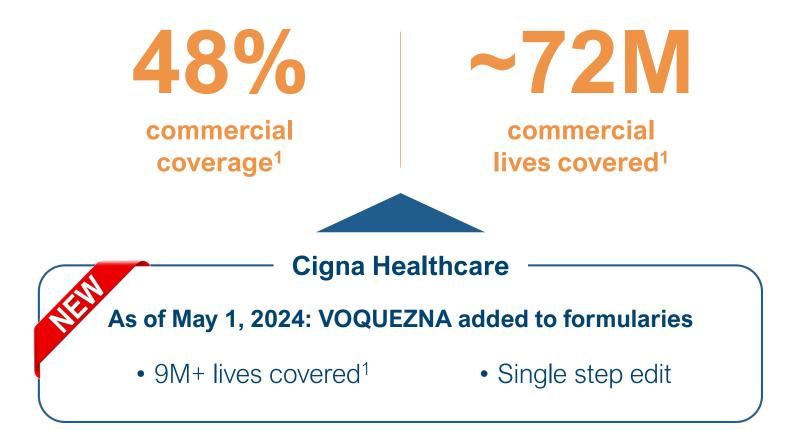


Cumulative Unique Writers²



Building expanded commercial coverage with large payers





Additional coverage anticipated throughout 2024







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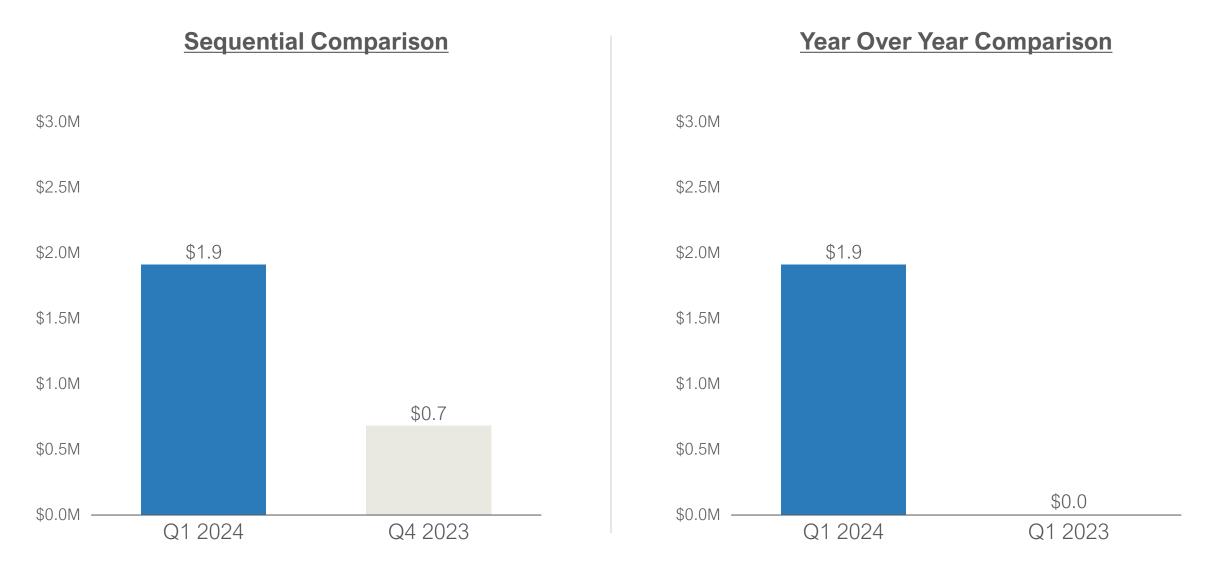


First Quarter 2024 Results

Molly Henderson, Chief Financial & Business Officer

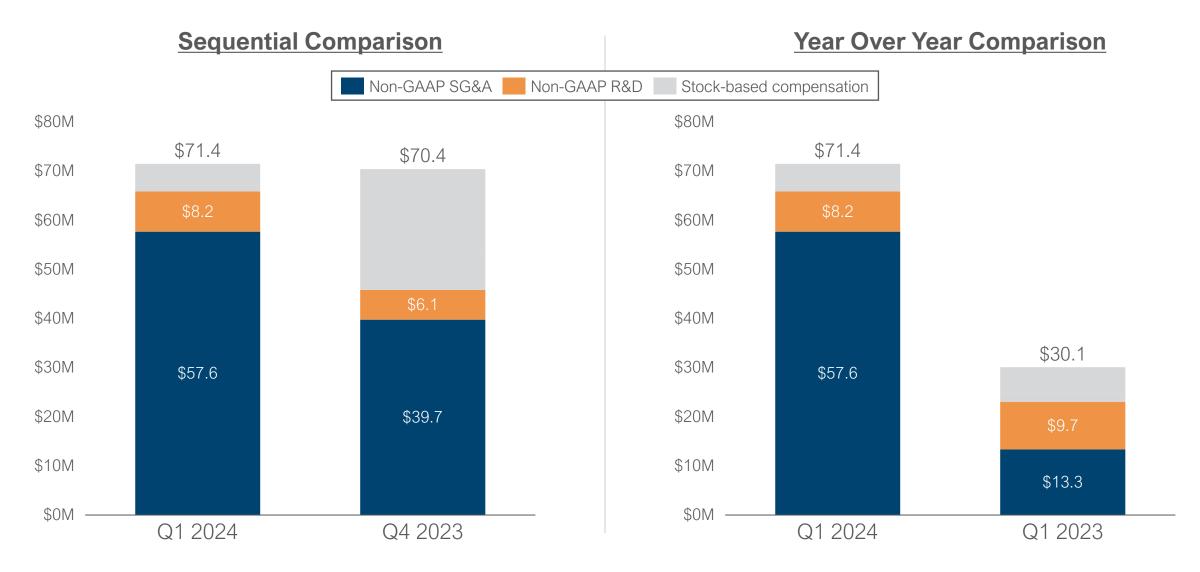


Key financial measures: Net Revenues





Key financial measures: Operating Expenses





Key financial measures: Adjusted Net Loss and Cash

Reconciliation of GAAP to Non-GAAP Adjusted Net Loss

(in thousands, except per share amounts)

<u>Q1 2024</u>	Q4 2023
(\$82,852)	(\$79,569)
\$5,626	\$24,583
\$11,956	\$8,462
\$474	\$566
(\$64,796)	(\$45,957)
(\$1.11)	(\$0.80)
	(\$82,852) \$5,626 \$11,956 \$474 (\$64,796)

\$322.2M

in cash and cash equivalents as of March 31, 2024

We believe this balance and other anticipated capital¹ will be sufficient to fund operations through the end of 2026



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Closing Remarks

Terrie Curran, President & Chief Executive Officer



Delivering on the VOQUEZNA launch strategy according to plan

4Q '23

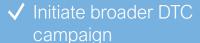
- ✓ Product in market
- ✓ Sales force onboarding
- ✓ Limited access/coverage
- ✓ HCP & consumer digital campaign launch

1Q '24



- ✓ Erosive GERD formulary reviews begin
- ✓ Ramp up of educational speaker programs

2Q '24



- + Growth in commercial access/coverage
- Increasing prescription
 volume as access is secured

2H '24



+ Accelerating prescription volume due to broader addressable population



Question & Answer

Terrie Curran, President & Chief Executive Officer Martin Gilligan, Chief Commercial Officer Molly Henderson, Chief Financial & Business Officer Azmi Nabulsi, Chief Operating Officer

