Alkermes°

First Quarter 2024 Financial Results & Business Update

May 1, 2024

Forward-Looking Statements and Non-GAAP Financial Information

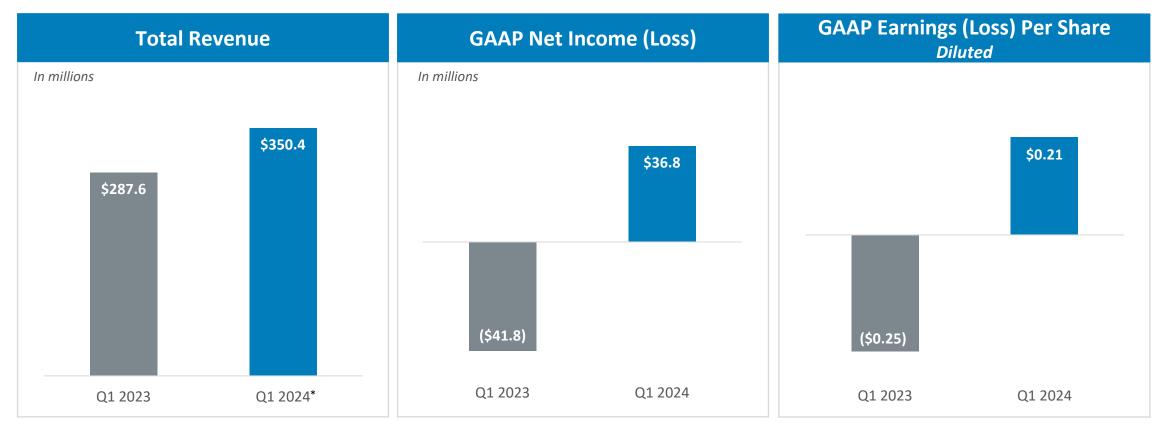
Certain statements set forth in this presentation constitute "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, including, but not limited to, statements concerning: Alkermes plc's (the "Company") expectations with respect to its current and future financial, commercial and operating performance, business plans or prospects, including its expected cash and revenue generation, expectations of profitability and potential return of capital to shareholders; the potential therapeutic and commercial value of the Company's marketed products and development candidates; the Company's expectations regarding plans and timelines for further clinical development activities, including study design and timelines and presentation of clinical data for ALKS 2680; and the Company's plans to advance and expand its neuroscience pipeline. The Company cautions that forward-looking statements are inherently uncertain. Actual performance and results may differ materially from those expressed or implied in the forward-looking statements due to various risks, assumptions and uncertainties. These risks, assumptions and uncertainties include, among others: whether the Company is able to sustain profitability; the unfavorable outcome of arbitration or litigation, including so-called "Paragraph IV" litigation or other patent litigation which may lead to competition from generic drug manufacturers, or other disputes related to the Company's products or products using the Company's proprietary technologies; the Company's commercial activities may not result in the benefits that the Company anticipates; clinical development activities may not be completed on time or at all; the results of the Company's development activities, including those related to ALKS 2680, may not be positive, or predictive of final results from such activities, results of future development activities or real-world results; potential changes in the cost, scope, design or duration of the Company's development activities, including the ALKS 2680 development program; the U.S. Food and Drug Administration ("FDA") or other regulatory authorities may not agree with the Company's regulatory approval strategies or components of the Company's marketing applications and may make adverse decisions regarding the Company's products; the Company and its licensees may not be able to continue to successfully commercialize their products or support growth of such products; there may be a reduction in payment rate or reimbursement for the Company's products or an increase in the Company's financial obligations to government payers; the Company's products may prove difficult to manufacture, be precluded from commercialization by the proprietary rights of third parties, or have unintended side effects, adverse reactions or incidents of misuse; and those risks, assumptions and uncertainties described under the heading "Risk Factors" in the Company's Annual Report on Form 10-K for the year ended Dec. 31, 2023 and in subsequent filings made by the Company with the U.S. Securities and Exchange Commission ("SEC"), which are available on the SEC's website at www.sec.gov, and on the Company's website at www.alkermes.com in the 'Investors – SEC filings' section. Existing and prospective investors are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date hereof. Except as required by law, the Company disclaims any intention or responsibility for updating or revising any forward-looking statements contained in this presentation.

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Note Regarding Trademarks: The Company and its affiliates are the owners of various U.S. federal trademark registrations (*) and other trademarks (TM), including ARISTADA*, ARISTADA INITIO*, LYBALVI* and VIVITROL*. Any other trademarks referred to in this presentation are the property of their respective owners. Appearances of such other trademarks herein should not be construed as any indicator that their respective owners will not assert their rights thereto.

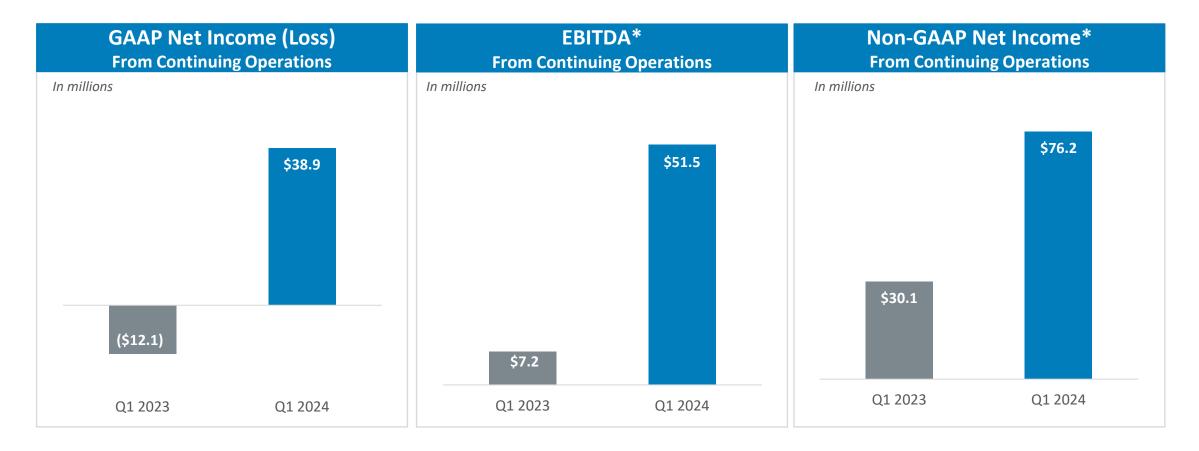
Q1 2024 Financial and Operational Performance

Q1 2024 Financial Results Summary



^{*}Reflects reinstatement of certain U.S. royalties following the successful outcome of the Company's arbitration with Janssen Pharmaceutica N.V. ("Janssen"), a subsidiary of Johnson & Johnson, announced in June 2023.

Q1 2024 Profitability From Continuing Operations



^{*}Reconciliation of this non-GAAP financial measure to the most directly comparable GAAP financial measure can be found in the Appendix of this presentation. EBITDA (earnings before interest, taxes, depreciation and amortization)



Q1 2024 Revenue Summary

In millions, except %	Q1′24	Q1′23	Δ Q1'24 vs. Q1'23
Total Proprietary Net Sales	\$233.5	\$214.7	9%
VIVITROL®	\$97.7	\$96.7	1%
ARISTADA®*	\$78.9	\$80.1	(2%)
LYBALVI®	\$57.0	\$38.0	50%
Manufacturing & Royalty Revenue**	\$116.8	\$72.9	60%
Total Revenue**	\$350.4	\$287.6	22%

Amounts in the table above may not sum due to rounding.

^{*}Inclusive of ARISTADA INITIO®

^{**}Reflects reinstatement of certain U.S. royalties following the successful outcome of the Company's arbitration with Janssen announced in June 2023.

Alkermes: 2024 Financial Expectations*

(in millions)	Financial Expectations for Year Ending Dec. 31, 2024
Total Revenues	\$1,500 – \$1,600
COGS	\$230 – \$250
R&D Expense	\$225 – \$255
SG&A Expense	\$625 – \$655
GAAP Net Income	\$350 – \$390
EBITDA [‡]	\$445 – \$485
Non-GAAP Net Income [‡]	\$465 – \$505
Effective Tax Rate	~17%

Expected net sales of proprietary products:

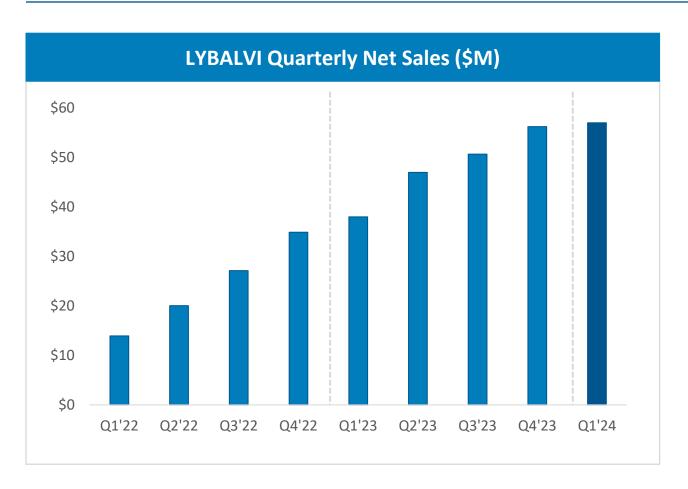
- VIVITROL® net sales of \$410M \$430M
- ARISTADA® net sales of \$340M \$360M
- LYBALVI® net sales of \$275M \$295M

^{*}These expectations were initially provided by the Company on Feb. 15, 2024, are reiterated by the Company on May 1, 2024 and are effective only as of such date. The Company expressly disclaims any obligation to update or reaffirm these expectations.

[‡]Reconciliation of these non-GAAP financial measures to the most directly comparable GAAP financial measures can be found in the Appendix of this presentation.

Q1 2024 Commercial Review

LYBALVI® Performance and Expectations



Q1'24 net sales of \$57.0M reflects 1% sequential growth compared to Q4'23

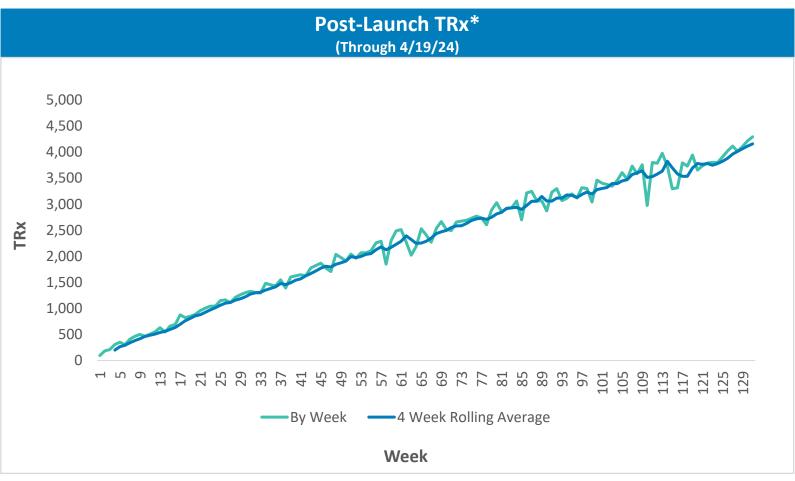
- Q1'24 gross-to-net deductions: ~29%
- Inventory in the channel decreased by ~\$2.3M during Q1'24

Outlook:

FY'24 net sales expected to range from \$275M - \$295M*

^{*}These expectations were initially provided by the Company on Feb. 15, 2024, are reiterated by the Company on May 1, 2024 and are effective only as of such date. The Company expressly disclaims any obligation to update or reaffirm these expectations.

LYBALVI® Prescription Growth Trends

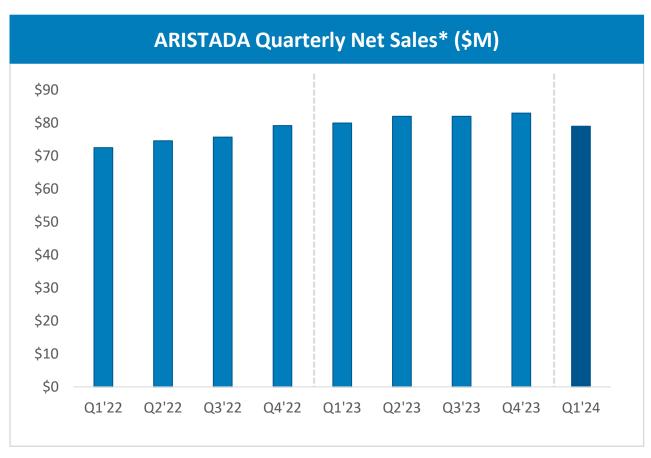


Q1'24 total TRx:

 ~49,600 reflecting 6% sequential growth compared to Q4'23

^{*}Source: IQVIA NPA Weekly

ARISTADA® Performance and Expectations



Q1'24 ARISTADA net sales were \$78.9M

 Inventory in the channel decreased by ~\$3.6M during Q1'24

Outlook:

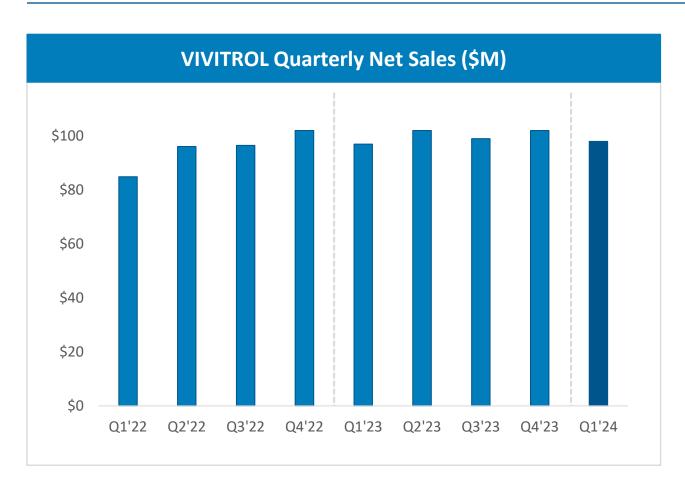
 FY'24 net sales expected to range from \$340M - \$360M^{+*}

[†] These expectations were initially provided by the Company on Feb. 15, 2024, are reiterated by the Company on May 1, 2024 and are effective only as of such date. The Company expressly disclaims any obligation to update or reaffirm these expectations.



^{*}Inclusive of ARISTADA INITIO®

VIVITROL® Performance and Expectations



Q1'24 VIVITROL net sales were \$97.7M

 Inventory in the channel decreased by ~\$4.3M during Q1'24

Outlook:

FY'24 net sales expected to range from \$410M - \$430M*

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Business Update

ALKS 2680: Investigational Oral Orexin 2 Receptor Agonist for the Once-Daily Treatment of Narcolepsy

Recent Progress

Narcolepsy Type 1 (NT1)

- ✓ Initiated Vibrance-1 phase 2 study
- ✓ Submitted data from the phase 1b NT1 cohort for presentation at upcoming medical meeting

Narcolepsy Type 2 (NT2) and Idiopathic Hypersomnia (IH)

- ✓ Reported positive topline results from phase 1b proof-of-concept NT2 and IH cohorts
- ✓ Selected phase 2 NT2 doses

Upcoming Program Priorities

NT1

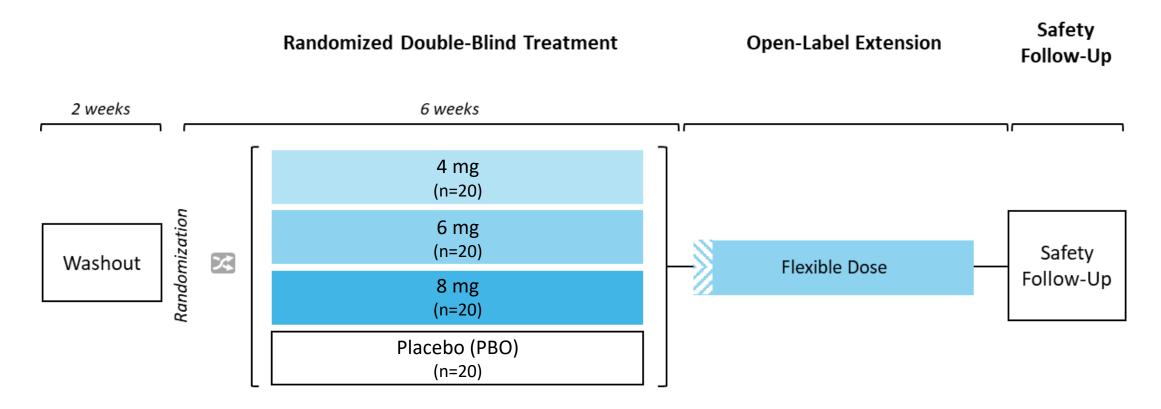
- ☐ Present additional phase 1b data at SLEEP 2024 June
- ☐ Enroll Vibrance-1 phase 2 study

NT2

- ☐ Present phase 1b data at upcoming medical meeting
- ☐ Initiate phase 2 study (Vibrance-2) expected in H2 2024

Vibrance-1 ALKS 2680 Phase 2 Study in Patients with NT1: Recently Initiated

Vibrance-1 Narcolepsy Type 1 Phase 2 Design



2024 Strategic Priorities



Deliver strong commercial growth and profitability

Driven by core products and streamlined operating structure



Advance orexin 2 receptor agonist program

Initiate phase 2 programs



Expand neuroscience pipeline

Advance internal development candidates and explore external pipeline opportunities



Plan for significant cash generation

Continue focus on capital allocation, including potential opportunities to return capital to shareholders

Appendix

Appendix: Amounts Included in Discontinued Operations

(In thousands)	Three Months	Three Months Ended March 31, 2024		
Cost of goods manufactured and sold	\$			
Research and development		2,516		
Selling, general and administrative				
Income tax benefit	\$	(396)		
Loss from discontinued operations, net of tax	\$	2,120		

Three		Ended March
(In thousands)		31, 2023
Cost of goods manufactured and sold	\$	11
Research and development		29,867
Selling, general and administrative		6,644
Income tax benefit	\$	(6,727)
Loss from discontinued operations, net of tax	\$	29,795

Appendix: Financial Results GAAP to Non-GAAP Adjustments

(In millions)			Three Months Ended March 31, 2023	
Net Income (Loss) from Continuing Operations — GAAP	\$ 38	3.9	\$	(12.1)
Adjustments:				
Share-based compensation expense	32	2.8		21.0
Depreciation expense	7	7.0		9.4
Amortization expense	<u>-</u>	1.1		8.8
Non-cash net interest expense	(0.1		0.1
Separation expense	(0.4		3.8
Income tax effect related to reconciling items	(4	.1)		(1.0)
Non-GAAP Net Income from Continuing Operations	\$ 76	5.2	\$	30.1
Non-GAAP Net Loss from Discontinued Operations	\$ (2	.1)	\$	(27.6)
Non-GAAP Net Income	\$ 74	4.1	\$	2.4

Amounts in the table above may not sum due to rounding.

Appendix: Financial Results GAAP to EBITDA Adjustments

(In millions)	Three Months Ended 1 March 31, 2024		Three Months Ended March 31, 2023	
Net Income (Loss) from Continuing Operations — GAAP	\$ 38.	9 \$	(12.1)	
Adjustments:				
Depreciation expense	7.	0	9.4	
Amortization expense	1.	1	8.8	
Interest income	(9.4	!)	(5.0)	
Interest expense	6.	0	5.3	
Income tax provision	7.	6	0.7	
EBITDA from Continuing Operations	\$ 51.	5 \$	7.2	
EBITDA from Discontinued Operations	\$ (2.5	5) \$	(36.0)	
EBITDA	\$ 49.	0 \$	(28.8)	

Appendix: 2024 Guidance GAAP to Non-GAAP Adjustments

(In millions, except per share data)	Year Ending December 31, 2024	Shares+	Earnings Per Share
Projected Net Income — GAAP	\$ 370.0	173.0	\$ 2.14
Adjustments:			
Share-based compensation expense	86.0		
Depreciation expense	35.0		
Amortization expense	1.0		
Non-cash net interest expense	0.5		
Income tax effect related to reconciling items	<u>(7.5)</u>		
Projected Net Income — Non-GAAP	\$ 485.0	173.0	\$ 2.80

Projected GAAP and non-GAAP measures reflect the mid-points within the Company's financial expectations ranges.



⁺²⁰²⁴ per share expectations are calculated based on a weighted average diluted share count of approximately 173.0 million shares outstanding.

Appendix: 2024 Guidance GAAP to EBITDA Adjustments

(In millions)	Year Ending December 31, 2024
Projected Net Income — GAAP	\$ 370.0
Adjustments:	
Net interest income	(16.0)
Depreciation expense	35.0
Amortization expense	1.0
Provision for income taxes	75.0
Projected EBITDA	\$ 465.0

Projected GAAP and non-GAAP measures reflect the mid-points within the Company's financial expectations ranges.

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