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Building the Premier Women's Health Company Q2 2022 Earnings August 15, 2022



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Forward-Looking Statements

This presentation by TherapeuticsMD, Inc. may contain forward-looking statements. Forward-looking statements may include, but are not limited to, statements relating to TherapeuticsMD's objectives, plans and strategies as well as statements, other than historical facts, that address activities, events or developments that the company intends, expects, projects, believes or anticipates will or may occur in the future. These statements are often characterized by terminology such as "believes," "hopes," "may," "anticipates," "should," "intends," "plans," "will," "expects," "estimates," "projects," "positioned," "strategy" and similar expressions and are based on assumptions and assessments made in light of management's experience and perception of historical trends, current conditions, expected future developments and other factors believed to be appropriate. Forward-looking statements in this press release are made as of the date of this press release, and the company undertakes no duty to update or revise any such statements, whether as a result of new information, future events or otherwise. Forward-looking statements are not guarantees of future performance and are subject to risks and uncertainties, many of which are outside of the company's control. Important factors that could cause actual results, developments and business decisions to differ materially from forward-looking statements are described in the sections titled "Risk Factors" in the company's filings with the Securities and Exchange Commission, including its most recent Annual Report on Form 10-K and Quarterly Reports on Form 10-Q, as well as reports on Form 8-K, and include the following: the effects of the COVID-19 pandemic; how the proceeds from the divestiture of the company's vitaCare business will be utilized; the company's ability to maintain or increase sales of its products; the company's ability to develop and commercialize IMVEXXY®, ANNOVERA®, and BIJUVA® and obtain additional financing necessary therefor; whether the company will be able to comply with the covenants and conditions under its term loan facility and the company's ability to refinance such facility; the effects of supply chain issues on the supply of the company's products; the potential of adverse side effects or other safety risks that could adversely affect the commercialization of the company's current or future approved products or preclude the approval of the company's future drug candidates; whether the FDA will approve the manufacturing supplement for ANNOVERA; the company's ability to protect its intellectual property, including with respect to the Paragraph IV notice letters the company received regarding IMVEXXY; the length, cost and uncertain results of future clinical trials; the company's reliance on third parties to conduct its manufacturing, research and development and clinical trials; the ability of the company's licensees to commercialize and distribute the company's products; the ability of the company's marketing contractors to market ANNOVERA; the availability of reimbursement from government authorities and health insurance companies for the company's products; the impact of product liability lawsuits; the influence of extensive and costly government regulation; the impact of leadership transitions; the volatility of the trading price of the company's common stock and the concentration of power in its stock ownership.

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Hugh O'Dowd CEO

Q2 Accomplishments

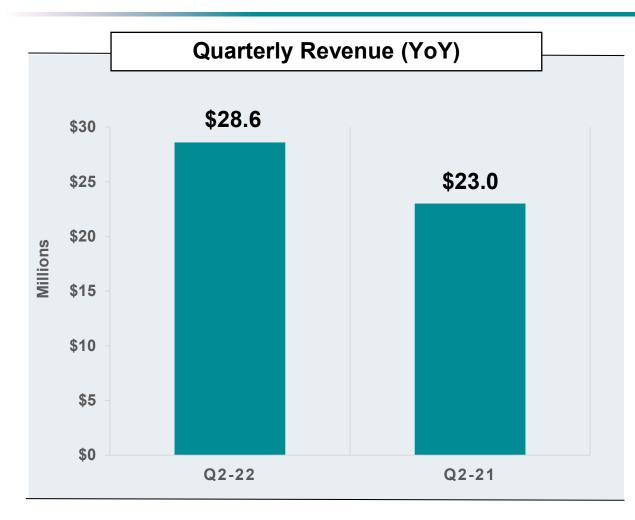
Top-line growth and overall operating performance

Successfully completed the divestiture of vitaCare, enabling the repayment of \$120 million of debt

Gained approval of our supplemental new drug application for ANNOVERA, which increases our current and future supply

Q2 2022 Financial Overview

Quarterly Revenue Trends



Q2-22 Highlights

- Total net revenue increased 24% in Q2-22 vs Q2-21
- Total net product revenue increased 24% in Q2-22 vs Q2-21
- ANNOVERA net revenue increased 91% in Q2-22 vs Q2-21 as supply and demand improved
- BIJUVA net revenue increased 23% in Q2-22 vs Q2-21
- IMVEXXY net revenue decreased 32% Q2-22 vs Q2-21

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Financial Results: Comparison Q2 2022 to Q2 2021 to Q1 2022

	Comparison of Key Financial Statement Items			
	[\$1,000's]			
		Q2-22	Q2-21	
	Balance Sheet ⁽¹⁾			
	Cash	\$ 26,303	\$ 111,424	
	Debt	\$ 90,780	\$ 185,300	
	Income Statement			
	Net Revenue	\$ 28,561	\$ 23,001	
	Gross Profit	\$ 23,821	\$ 18,869	
	Gross Margin %	83%	82%	
	Total Operating Expenses	\$ 42,662	\$ 54,048	
	Net Profit / (Loss)	\$ 112,281	\$ (42,652)	
٢	Cash Flow			
	Net Cash Used In Operating Activities	\$ (15,404)	\$ (26,532)	

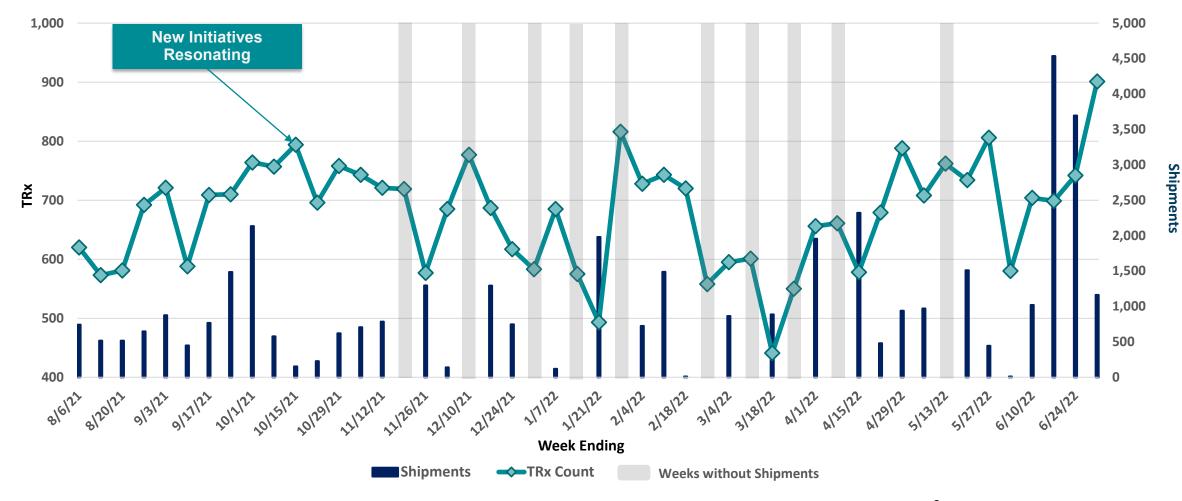
- In April, \$120M in proceeds from the vitaCare divestiture were used to paydown the debt.
- Gross margin of 83% in Q2-22 is relatively consistent to Q2-21.
- Operating expenses in Q2-22 decreased \$11.4M from Q2-21.
- Net Profit in Q2-22 resulted from the sale of vitaCare. Net Operating Loss;
 - \$18.8M Q2-2022
 - \$35.2M Q2-2021
- Net cash used in operating activities in Q2-22 decreased \$11.1M from Q2-21.

(1) Balance Sheet as of quarter end.

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Q2 2022 Commercial Overview

Inventory Impact



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Sources: TRx: Symphony Metys - 7/1/22 Shipments: CORD

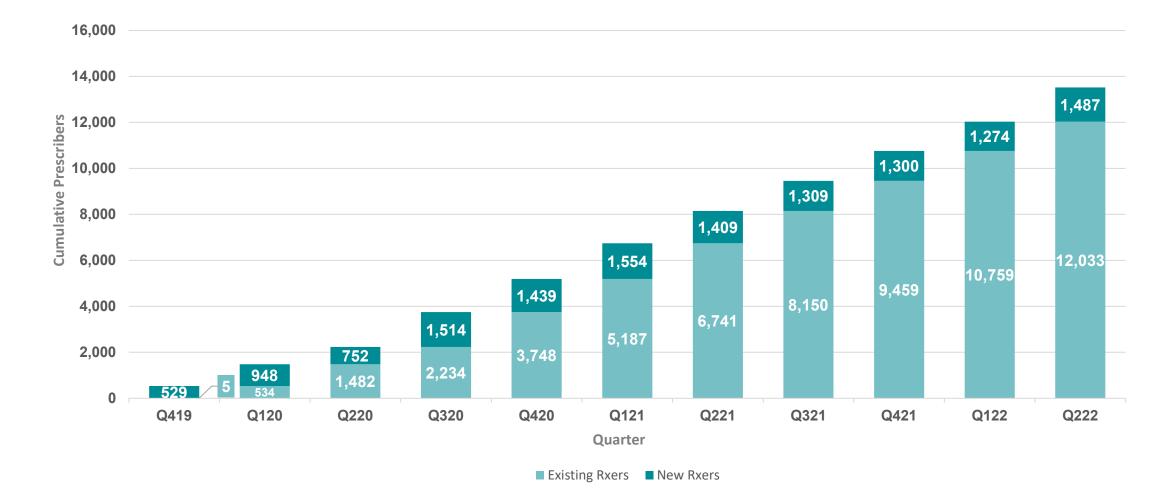
ANNOVERA Monthly TRx



Month

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ANNOVERA Prescribers: Launch to Date



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Source: Symphony Metys – June 2022; 5 prescribers in Q319 New prescribers defined as not having written Annovera in prior periods

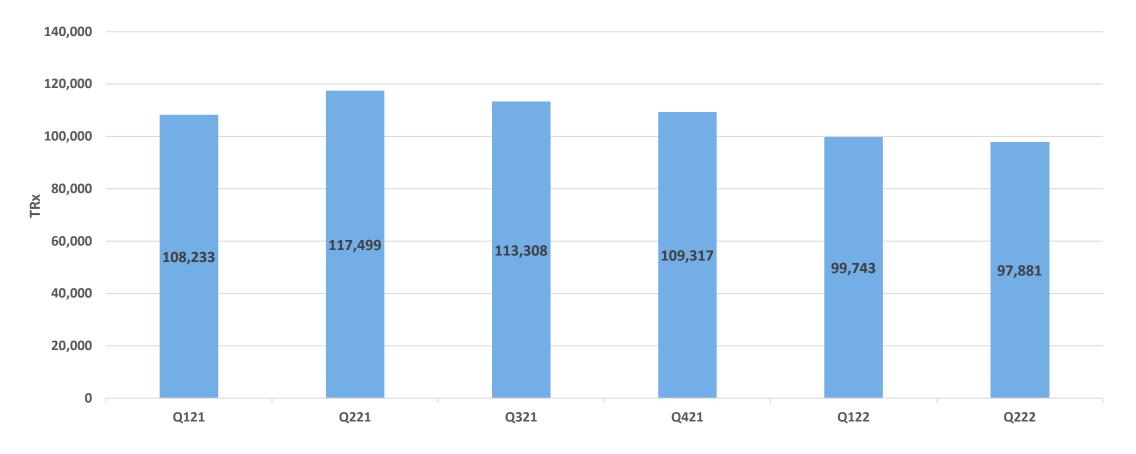
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Requests for Administration/Tri-Agencies & Impact for TXMD

REQUEST	POTENTIAL IMPACT	
1 Standardize the forms for the exceptions process	Vast majority of exceptions (LMNs) approved instantaneously LMN denials are the greatest burden to Rx fulfillment for ANNOVERA A standard form and process would almost eliminate denials 	
2 Greater enforcement is needed Enforce existing requirements and penalties	Enforcement of current legislative language would ensure more Rx fulfillment. – Payors react to penalties and risk; if they fall in-line, coverage and barriers to fulfillment would immediately decrease	
3 Remove reference to FDA-approved methods in HRSA guidelines (after the exceptions process and enforcement are addressed)	If all brands without generic substitution were required to be covered at a \$0 cost share. ANNOVERA would be covered 100% of the time. – With exception to the religious beliefs clause	

IMVEXXY Quarterly TRx

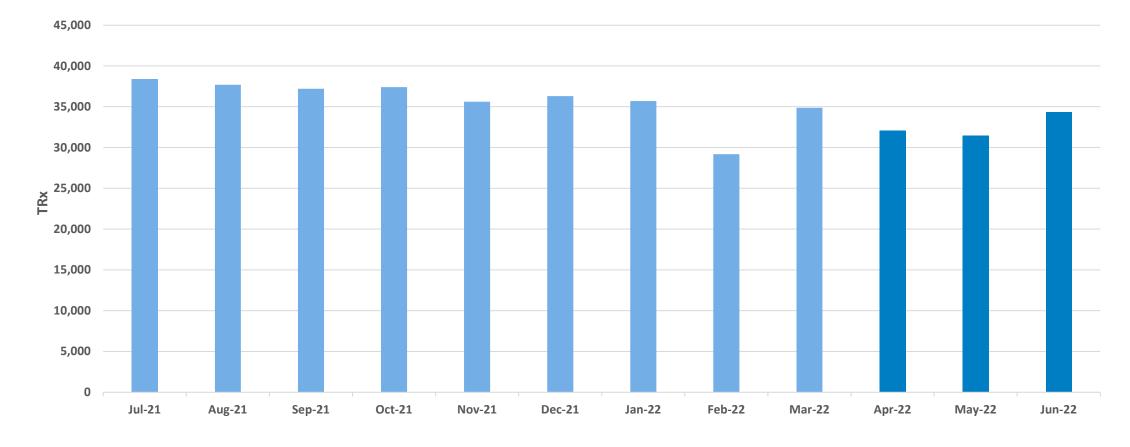
Held up well considering prioritization on Annovera beginning Q321



Month

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IMVEXXY Monthly TRx



Month

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IMVEXXY Prescribers: Launch to Date

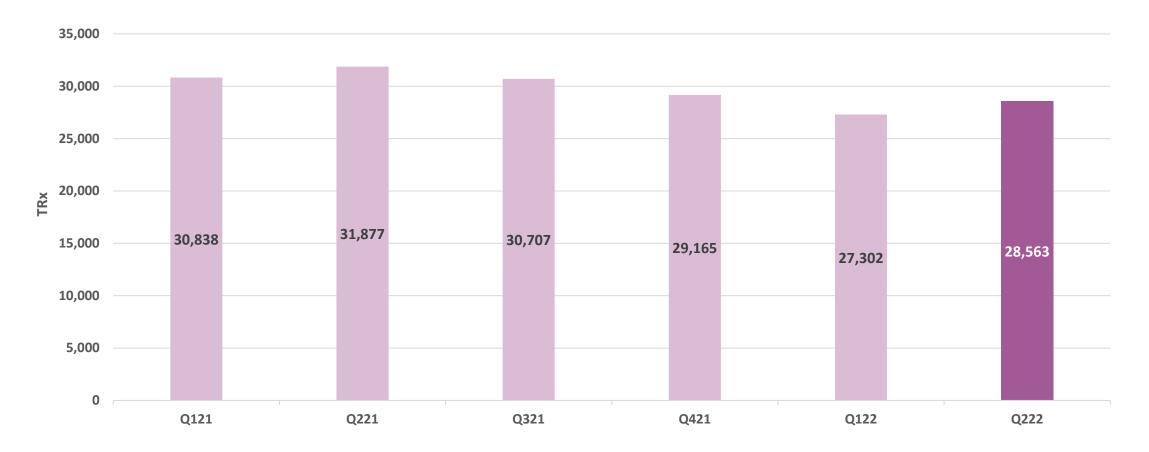


Existing Rxers New Rxers

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Source: Symphony Metys – June 2022; 21,677 prescribers prior to Q121 New prescribers defined as not having written Imvexxy in prior periods

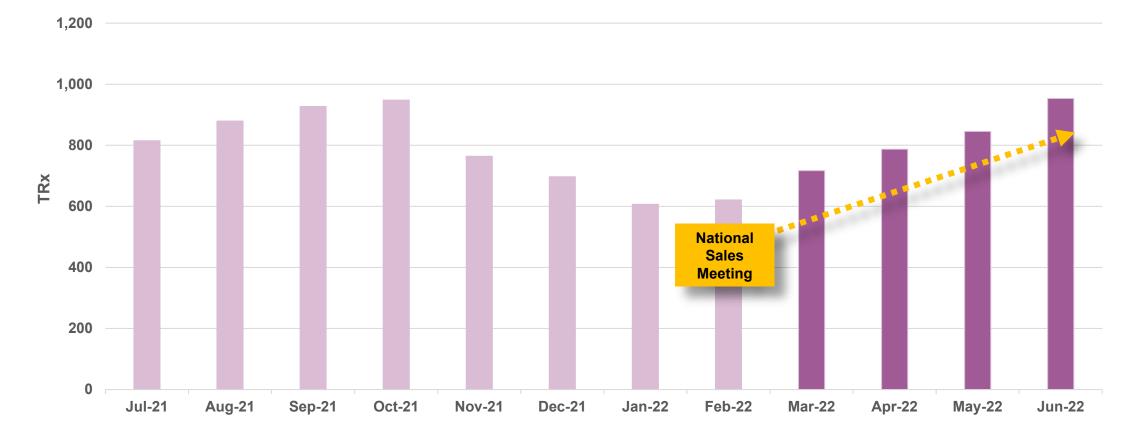
BIJUVA Quarterly TRx



Month

BIJUVA New-to-Brand TRx*

Growth coincides with targeting strategy change initiated post-NSM



Month

* New to brand includes patient switches/adds and new to market patients

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