

NEWS RELEASE

Citius Oncology, Inc. Reports Fiscal Second Quarter 2025 Financial Results and Provides Business Update

2025-05-14

CRANFORD, N.J., May 14, 2025 /PRNewswire/ -- Citius Oncology, Inc. ("Citius Oncology" or the "Company") (Nasdaq: CTOR), a specialty biopharmaceutical company focused on the development and commercialization of novel targeted oncology therapies, today reported business and financial results for the fiscal second quarter ended March 31, 2025.

"In Q2 2025, Citius Oncology advanced its transformation from a development-stage company to a commercial-stage organization. Following FDA approval of LYMPHIR, we intensified our focus on disciplined capital deployment and operational execution to support the success of our planned U.S. launch," said Leonard Mazur, Chairman and CEO of Citius Oncology and Citius Pharmaceuticals.

"This quarter's progress underscores our commitment to creating long-term value by ensuring LYMPHIR reaches patients with cutaneous T-cell lymphoma. Discussions with prospective commercial and strategic partners are underway as we concurrently pursue opportunities to secure additional capital to enhance our financial flexibility. These efforts are critical as we lay the foundation for sustained commercial success. With disciplined execution and a focused strategic vision, we believe Citius Oncology is poised to deliver meaningful near-term impact and durable shareholder value," concluded Mazur.

FISCAL SECOND QUARTER 2025 FINANCIAL RESULTS:

Liquidity

Citius Oncology is a subsidiary of Citius Pharma. Citius Pharma plans to continue to fund Citius Oncology until Citius Oncology raises adequate capital through equity financings from outside investors and/or generates revenue from the future sales of LYMPHIR. Citius Oncology has also retained Jefferies LLC as exclusive financial advisor to evaluate strategic alternatives aimed at maximizing stockholder value.

As of March 31, 2025, the Company had \$112 in cash and cash equivalents and 71,552,402 common shares outstanding. Citius Oncology will need to secure additional capital to support operations beyond May 2025.

Research and Development (R&D) Expenses

R&D expenses were \$3.1 million for the quarter ended March 31, 2025, as compared to \$1.3 million for the quarter ended March 31, 2024. For the six months ended March 31, 2025, R&D expenses were \$4.4 million, as compared to \$2.5 million for the six months ended March 31, 2024. The increase is primarily related to costs associated with the expense of a drug substance batch needed for the pre-license inspection of the manufacturer.

General and Administrative (G&A) Expenses

G&A expenses were \$2.2 million for the quarter ended March 31, 2025, as compared to \$1.4 million for the quarter ended March 31, 2024. For the six months ended March 31, 2025, G&A expenses were \$5.5 million, as compared to \$2.9 million for the six months ended March 31, 2024. The increase was primarily due to costs associated with precommercial and commercial launch activities of LYMPHIR including market research, marketing, distribution and drug product reimbursement from health plans and payers.

Stock-based Compensation Expense

For the quarter ended March 31, 2025, stock-based compensation expense was \$2.1 million, as compared to \$2.0 million for the quarter ended March 31, 2024. For the six months ended March 31, 2025, stock-based compensation expense was \$3.9 million, as compared to \$3.9 million for the six months ended March 31, 2024. The increase was primarily due to new options granted in December 2024.

Net loss

Net loss was \$7.7 million, or (\$0.11) per share, for the quarter ended March 31, 2025, as compared to a net loss of \$4.8 million, or (\$0.07) per share, for the quarter ended March 31, 2024. Net loss for the six months ended March 31, 2025 was \$14.4 million, as compared to a net loss of \$9.6 million for the six months ended March 31, 2024. The increase in net loss was primarily due to the increase in our operating expenses.

About Citius Oncology, Inc.

Citius Oncology specialty is a biopharmaceutical company focused on developing and commercializing novel targeted oncology therapies. In August 2024, its primary asset, LYMPHIR, was approved by the FDA for the treatment of adults with relapsed or refractory CTCL who had had at least one prior systemic therapy. Management estimates the initial market for LYMPHIR currently exceeds \$400 million, is growing, and is underserved by existing therapies. Robust intellectual property protections that span orphan drug designation, complex technology, trade secrets and pending patents for immuno-oncology use as a combination therapy with checkpoint inhibitors would further support Citius Oncology's competitive positioning. Citius Oncology is a publicly traded subsidiary of Citius Pharmaceuticals. For more information, please visit www.citiusonc.com

Forward-Looking Statements

This press release may contain "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934. Such statements are made based on our expectations and beliefs concerning future events impacting Citius Oncology. You can identify these statements by the fact that they use words such as "will," "anticipate," "estimate," "expect," "plan," "should," and "may" and other words and terms of similar meaning or use of future dates. Forward-looking statements are based on management's current expectations and are subject to risks and uncertainties that could negatively affect our business, operating results, financial condition and stock price. Factors that could cause actual results to differ materially from those currently anticipated, and, unless noted otherwise, that apply to Citius Oncology are: our need for substantial additional funds and our ability to raise additional money to fund our operations beyond May 2025 and for at least the next 12 months as a going concern; our ability to commercialize LYMPHIR, including covering the costs of licensing payments, product manufacturing and other third-party goods and services, and any of our other product candidates that may be approved by the FDA; our ability to obtain, perform under and maintain financing and strategic agreements and relationships; the estimated markets for our product candidates and the acceptance thereof by any market; our ability to regain compliance with Nasdag's continued listing standards; the ability of our product candidates to impact the quality of life of our target patient populations; our dependence on third-party suppliers; our ability to procure cGMP commercial-scale supply; risks related to research using our assets but conducted by third parties; uncertainties relating to preclinical and clinical testing; market and other conditions; risks related to our growth strategy; patent and intellectual property matters; our ability to identify, acquire, close and integrate product candidates and companies successfully and on a timely basis; government regulation; competition; as well as other risks described in our Securities and Exchange Commission ("SEC") filings. These risks have been and may be further impacted by any future public health risks. Accordingly, these forward-looking statements do not constitute guarantees of future performance, and you are cautioned not to place undue reliance on these forward-looking statements. Risks regarding our business are described in detail

in our SEC filings which are available on the SEC's website at www.sec.gov, including in Citius Oncology's Annual Report on Form 10-K for the year ended September 30, 2024, filed with the SEC on December 27, 2024, as amended on January 27, 2025, Citius Oncology's Quarterly Report on Form 10-Q for the quarter ended March 31, 2025, filed with the SEC on May 14, 2025, and as updated by our subsequent filings with the SEC. These forward-looking statements speak only as of the date hereof, and we expressly disclaim any obligation or undertaking to release publicly any updates or revisions to any forward-looking statements contained herein to reflect any change in our expectations or any changes in events, conditions or circumstances on which any such statement is based, except as required by law.

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-- Financial Tables Follow -

CITIUS ONCOLOGY, INC. CONDENSED CONSOLIDATED BALANCE SHEETS (Unaudited)

	March 31, 2025	September 30, 2024		
Current Assets: Cash and cash equivalents Inventory Prepaid expenses Total Current Assets	\$ 112 15,339,253 2,700,000 18,039,365	\$ 112 8,268,766 2,700,000 10,968,878		
Other Assets: In-process research and development Total Other Assets	73,400,000 73,400,000	73,400,000 73,400,000		
Total Assets LIABILITIES AND STOCKHOLDERS' EQUITY Current Liabilities:	\$ 91,439,365	\$ 84,368,878		

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4,941,664 49,740,142		588,806 32,700,428
2,256,480 3,800,111 55,796,733		1,728,000 3,800,111 38,228,539
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7,155 89,308,821 (53,673,344) 35,642,632		7,155 85,411,771 (39,278,587) 46,140,339 84,368,878
	4,941,664 49,740,142 2,256,480 3,800,111 55,796,733 7,155 89,308,821 (53,673,344)	49,740,142 2,256,480 3,800,111 55,796,733 7,155 89,308,821 (53,673,344) 35,642,632

CITIUS ONCOLOGY, INC. CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS FOR THE THREE AND SIX MONTHS ENDED MARCH 31, 2025 AND 2024 (Unaudited)

	Three Months Ended		Six Months Ended			nded		
	N	/larch 31, 2025	N	March 31, 2024		March 31, 2025	N	March 31, 2024
Revenues	\$		\$	_	\$		\$	
Operating Expenses Research and development General and administrative Stock-based compensation – general and administrative Total Operating Expenses		3,139,413 2,243,327 2,088,572 7,471,312		1,348,966 1,385,580 1,957,000 4,691,546		4,403,921 5,565,306 3,897,050 13,866,277		2,497,461 2,903,488 3,874,000 9,274,949
Loss before Income Taxes Income tax expense		(7,471,312) 264,240		(4,691,546) 144,000		(13,866,277) 528,480		(9,274,949) 288,000
Net Loss	\$	(7,735,552)	\$	(4,835,546)	\$	(14,394,757)	\$	(9,562,949)
Net Loss Per Share - Basic and Diluted	\$	(0.11)	\$	(0.07)	\$	(0.20)	\$	(0.14)
Weighted Average Common Shares Outstanding Basic and diluted		71,552,402		67,500,000		71,552,402		67,500,000

CITIUS ONCOLOGY, INC.
CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS
FOR THE SIX MONTHS ENDED MARCH 31, 2025 AND 2024
(Unaudited)

2025 2024

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Cash Flows From Operating Activities: Net loss Adjustments to reconcile net loss to net cash provided by operating activities: Stock-based compensation expense Deferred income tax expense Changes in operating assets and liabilities: Inventory Prepaid expenses Accounts payable Accrued expenses Due to related party Net Cash Provided By Operating Activities	\$	(14,394,757) 3,897,050 528,480 (7,070,487) 3,964,688 8,722,168 4,352,858	\$	(9,562,949) 3,874,000 288,000 (1,171,920) (785,132) (259,071) 7,617,072
Net Change in Cash and Cash Equivalents Cash and Cash Equivalents – Beginning of Period Cash and Cash Equivalents – End of Period	<u> </u>	- 112 112	*	<u>-</u>

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