UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, DC 20549

| | | FORM 8-K | |
|------|---|--|--|
| | | CURRENT REPORT SUANT TO SECTION 13 OR 15(d) OF ECURITIES EXCHANGE ACT OF 19 | |
| | Date of Rep | oort (Date of earliest event reported): Aug | ust 10, 2023 |
| | | Onconova Therapeutics, Inc | |
| | Delaware (State or Other Jurisdiction of Incorporation or Organization) | 001-36020 (Commission File Number) | 22-3627252 (I.R.S. Employer Identification No.) |
| | (Address Including Zin Code and Tele | 12 Penns Trail Newtown, PA 18940 (267) 759-3680 phone Number, Including Area Code, of I | Registrant's Principal Executive Offices) |
| | | Not Applicable name or former address, if changed since l | |
| | ck the appropriate box below if the Form 8-K fili | ng is intended to simultaneously satisfy | the filing obligation of the registrant under any of the |
| | Written communications pursuant to Rule 425 un | der the Securities Act (17 CFR 230.425) | |
| | Soliciting material pursuant to Rule 14a-12 under | the Exchange Act (17CFR 240.14a-12) | |
| | Pre-commencement communications pursuant to | Rule 14d-2(b) under the Exchange Act (1 | 7 CFR 240.14d-2(b)) |
| | Pre-commencement communications pursuant to | Rule 13e-4(c) under the Exchange Act (17 | 7 CFR 240.13e-4(c)) |
| Secu | urities registered pursuant to Section 12(b) of the Act | : | |
| | Title of each class | Trading Symbol(s) | Name of each exchange on which registered |
| | Common Stock, par value \$.01 per share | ONTX | The Nasdaq Stock Market LLC |
| | cate by check mark whether the registrant is an enoter) or Rule 12b-2 of the Securities Exchange Act of | | rule 405 of the Securities Act of 1933 (§230.405 of this |
| Eme | erging growth company \Box | | |
| | | | |
| | | | |
| | | | |

Item 2.02 Results of Operations and Financial Condition.

On August 10, 2023, Onconova Therapeutics, Inc. (the "Company") issued a press release announcing its financial results for the quarter and six months ended June 30, 2023, a copy of which is attached hereto as Exhibit 99.1 and incorporated herein by reference. The information contained in this Form 8-K (including the exhibit hereto) shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as shall be expressly set forth by specific reference in such a filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

| Exhibit No. | Exhibit |
|-------------|--|
| <u>99.1</u> | Press release issued by the Company dated August 10, 2023 |
| 104 | The cover page from this Current Report on Form 8-K, formatted in Inline XBRL. |

SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, as amended, the Registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Dated: August 10, 2023 Onconova Therapeutics, Inc.

By: /s/ MARK GUERIN

Name: Mark Guerin

Title: Chief Operating Officer & Chief Financial Officer

Onconova Therapeutics Reports Corporate Update and Announces Second Quarter 2023 Financial Results

Anticipate topline results from the Phase 1 monotherapy and Phase 1/2 combination study with letrozole in Q4 2023

Plans are underway for a registrational trial with rigosertib in patients with RDEB-associated squamous cell carcinoma based on a constructive Type B FDA meeting held in June

Company to host conference call and webcast at 4:30 p.m. ET on Thursday, August 10, 2023

NEWTOWN, PA August 10, 2023 (GLOBE NEWSWIRE) -- <u>Onconova Therapeutics, Inc.</u> (NASDAQ: ONTX), ("Onconova" or "the Company"), a clinical-stage biopharmaceutical company focused on discovering and developing novel products for patients with cancer, today reported second quarter 2023 financial results and provided an update on recent pipeline progress. Management plans to host a conference call and live webcast at 4:30 p.m. ET today to discuss these results.

"We are very encouraged about the recent progress that the Onconova team has made for our two lead programs, narazaciclib, a differentiated multikinase CDK4/6 inhibitor targeting proteins involved in resistance pathways, and rigosertib, a cell signaling inhibitor, over the last few months, while effectively managing our financial resources. In addition, we are pleased that Victor Moyo, M.D., a highly experienced and successful clinical researcher and drug developer, has agreed to join the Company as Consulting Chief Medical Officer. We look forward to sharing several important updates in the coming months," said Steve Fruchtman, M.D., President and Chief Executive Officer.

Dr. Fruchtman continued, "For narazaciclib, our efforts have been dedicated to completing a Phase 1 program and defining a recommended Phase 2 dose to support evaluation of narazaciclib in a randomized trial. Onconova believes this CDK4/6 compound has the potential to provide differentiated efficacy based on targeting proteins that have been implicated in resistance mechanisms and the potential for an improved safety profile. We are pleased to see target engagement based on an assay measuring proliferation. We expect to report the results from our Phase 1 monotherapy and Phase 1/2 combination study with letrozole in Q4 2023. The readout will include safety, pharmacokinetics and the definition of a recommended Phase 2 dose."

Dr. Fruchtman concluded, "For rigosertib, we continue to believe this rigosertib's unique action on cell signaling pathways, including K-RAS and PLK-1, combined with an acceptable safety profile, could position it as an attractive anti-cancer agent. In June, we had a constructive Type B meeting with the FDA for the use of rigosertib monotherapy in the lead, ultra-rare indication of RDEB-associated squamous cell carcinoma. Based on that meeting and the impressive clinical responses in previously refractory patients we have seen and presented at major medical meetings, we plan to design a registrational trial and will look to provide an update on next steps in H1 2024. In the meantime, we continue to support two investigator sponsored studies for rigosertib, underway in melanoma and KRAS mutated non-small cell lung cancer which includes any KRAS mutation that may be present."

Second Quarter Financial Results

Cash and cash equivalents as of June 30, 2023, were \$29.7 million, compared to \$38.8 million as of December 31, 2022. The Company believes that its cash and cash equivalents will be sufficient to fund ongoing clinical trials and business into the second quarter of 2024.

Research and development expenses were \$2.5 million for the second guarter of 2023, compared with \$2.0 million for the second guarter of 2022.

General and administrative expenses were \$2.2 million for the second quarter of 2023, compared with \$2.1 million for the second quarter of 2022.

Net loss for the second quarter of 2023 was \$4.3 million, or \$0.20 per share on 21.0 million weighted shares outstanding, compared with a net loss of \$4.0 million, or \$0.19 per share for the second quarter of 2022 on 20.9 million weighted shares outstanding.

Conference Call and Webcast Information

Interested parties who wish to participate in the conference call may do so by dialing:

- · (800) 715-9871 for domestic and
- · (646) 307-1963 for international callers and
- · Using conference ID 9506701

Those interested in listening to the conference call via the internet may do so by visiting the investors and media page on the Company's website at www.onconova.com and clicking on the webcast link. In addition to the live webcast, a replay will be available on the Onconova website for 90 days following the call.

About Onconova Therapeutics, Inc.

Onconova Therapeutics is a clinical-stage biopharmaceutical company focused on discovering and developing novel products for patients with cancer. The Company's product candidates include proprietary targeted anti-cancer agents designed to disrupt specific cellular pathways that are important for cancer cell proliferation.

Onconova's novel, proprietary multi-kinase inhibitor narazaciclib (formerly ON 123300) is being evaluated in a Phase 1/2 combination trial with the estrogen blocker, letrozole, in advanced low grade endometrial cancer (NCT05705505). Based on preclinical and clinical studies of CDK 4/6 inhibitors, Onconova is also evaluating opportunities for combination studies with narazaciclib and letrozole in additional indications.

Onconova's product candidate rigosertib is being studied in multiple investigator-sponsored studies. These studies include a dose-escalation and expansion Phase 1/2a study of oral rigosertib in combination with nivolumab in patients with KRAS+ non-small cell lung cancer (NCT04263090), a Phase 2 program evaluating oral or IV rigosertib monotherapy in advanced squamous cell carcinoma complicating recessive dystrophic epidermolysis bullosa (RDEB-associated SCC (NCT03786237, NCT04177498), and a Phase 2 trial evaluating rigosertib in combination with pembrolizumab in patients with metastatic melanoma(NCT05764395).

For more information, please visit $\underline{www.onconova.com}$.

Forward Looking Statements

Some of the statements in this release are forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended, Section 21E of the Securities Exchange Act of 1934, as amended, and the Private Securities Litigation Reform Act of 1995, and involve risks and uncertainties. These statements relate to Onconova's expectations regarding its clinical development and trials, its product candidates and its business and financial position. Onconova has attempted to identify forward-looking statements by terminology including "believes," "estimates," "anticipates," "expects," "plans," "intends," "may," "could," "might," "will," "should," "preliminary," "encouraging," "approximately" or other words that convey uncertainty of future events or outcomes. Although Onconova believes that the expectations reflected in such forward-looking statements are reasonable as of the date made, expectations may prove to have been materially different from the results expressed or implied by such forward-looking statements. These statements are only predictions and involve known and unknown risks, uncertainties, and other factors, including the success and timing of Onconova's clinical trials, investigator-sponsored trials, regulatory agency and institutional review board approvals of protocols, Onconova's collaborations, market conditions and those discussed under the heading "Risk Factors" in Onconova's most recent Annual Report on Form 10-K and quarterly reports on Form 10-Q. Any forward-looking statements contained in this release speak only as of its date. Onconova undertakes no obligation to update any forward-looking statements contained in this release to reflect events or circumstances occurring after its date or to reflect the occurrence of unanticipated events.

Company Contact:

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Investor Contact:

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ONCONOVA THERAPEUTICS, INC. Condensed Consolidated Balance Sheets (in thousands)

| | June | , | December 31, 2022 | |
|--|--|-----------|----------------------|-----------|
| Assets | , and the second | | | |
| Current assets: | | | | |
| Cash and cash equivalents | \$ | 29,729 | \$ | 38,757 |
| Receivables | | 17 | | 29 |
| Prepaid expenses and other current assets | | 704 | | 561 |
| Total current assets | | 30,450 | | 39,347 |
| Property and equipment, net | | 17 | | 24 |
| Other non-current assets | | 1 | | 1 |
| Total assets | \$ | 30,468 | \$ | 39,372 |
| Liabilities and stockholders' equity | | | | |
| Current liabilities: | | | | |
| Accounts payable | \$ | 5,071 | \$ | 3,860 |
| Accrued expenses and other current liabilities | | 3,369 | | 3,960 |
| Deferred revenue | | 226 | | 226 |
| Total current liabilities | | 8,666 | | 8,046 |
| Deferred revenue, non-current | | 2,904 | | 3,017 |
| Total liabilities | | 11,570 | | 11,063 |
| Stockholders' equity: | | | | |
| Preferred stock | | - | | - |
| Common stock | | 210 | | 209 |
| Additional paid in capital | | 492,424 | | 491,816 |
| Accumulated other comprehensive loss | | (28) | | (33) |
| Accumulated deficit | | (473,708) | | (463,683) |
| Total stockholders' equity | | 18,898 | | 28,309 |
| Total liabilities and stockholders' equity | \$ | | \$ | 39,372 |

ONCONOVA THERAPEUTICS, INC. Condensed Consolidated Statements of Operations (unaudited) (in thousands, except share and per share amounts)

| | Three Months Ended June 30, | | | Six months months ended June 30, | | | | |
|---|-----------------------------|------------|------|----------------------------------|------|------------|----|------------|
| | 2023 | | 2022 | | 2023 | | | 2022 |
| Revenue | \$ | 57 | \$ | 57 | \$ | 113 | \$ | 113 |
| Operating expenses: | | | | | | | | |
| General and administrative | | 2,211 | | 2,139 | | 4,324 | | 4,325 |
| Research and development | | 2,456 | | 2,038 | | 6,536 | | 4,040 |
| Total operating expenses | | 4,667 | | 4,177 | | 10,860 | | 8,365 |
| Loss from operations | | (4,610) | | (4,120) | | (10,747) | | (8,252) |
| Other income, net | | 360 | | 96 | | 722 | | 106 |
| Net loss | | (4,250) | | (4,024) | | (10,025) | | (8,146) |
| Net loss per share of common stock, basic and diluted | \$ | (0.20) | \$ | (0.19) | \$ | (0.48) | \$ | (0.39) |
| Basic and diluted weighted average shares outstanding | | 20,979,766 | | 20,904,085 | | 20,970,022 | | 20,904,085 |