UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 8-K

CURRENT REPORT
Pursuant to Section 13 or 15(d)
of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): January 29, 2021

MEDICINOVA, INC.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation) 001-33185 (Commission File Number) 33-0927979 (IRS Employer Identification No.)

4275 EXECUTIVE SQUARE, SUITE 300, LA JOLLA, CA (Address of principal executive offices)

92037 (Zip Code)

Registrant's telephone number, including area code: (858) 373-1500

foll	Check the appropriate box below if the Form 8- owing provisions (<u>see</u> General Instruction A.2. bel	· · ·	filing obligation of the registrant under any of the	
	Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)			
	Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)			
	Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))			
	Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))			
Sec	urities registered pursuant to Section 12(b) of the A	Act:		
	Common Stock, \$0.001 par value	MNOV	The Nasdaq Stock Market LLC	
	(Title of each class)	(Trading symbol(s))	(Name of each exchange on which registered)	
	cate by check mark whether the registrant is an entule 12b-2 of the Securities Exchange Act of 1934			
If aı				

Item 3.02. Unregistered Sale of Equity Securities.

The information contained below in Item 8.01 related to the sale of the Shares (as defined below) is hereby incorporated by reference into this Item 3.02.

On January 29, 2021 MediciNova, Inc. (the "Company") sold and issued 3,656,307 shares of its Common Stock, par value \$0.001 per share (the "Shares") at a price of \$5.47 per share for approximately \$20 million in cash to 3D Opportunity Master Fund (the "Investor") in a private placement (the "Private Placement") pursuant to the terms and conditions of that certain Securities Purchase Agreement (the "Purchase Agreement") dated as of January 11, 2021 by and between the Company and the Investor. The sale of the Shares in the Private placement was exempt from registration under the Securities Act of 1933, as amended (the "Act") pursuant to the provisions of Section 4(a)(2) of the Act. The Purchase Agreement was filed as Exhibit 10.1 to the Company's Current Report on Form 8-K filed on January 12, 2021, and incorporated into this Item 3.02 by reference, and the foregoing summary of the Purchase Agreement is qualified in its entirety by reference to Exhibit 10.1.

Item 8.01. Other Events.

On January 31, 2021, the Company issued a press release announcing the closing of the Private Placement and the sale of the Shares to the Investor. The full text of the press release is attached hereto as Exhibit 99.1 to this Current Report on Form 8-K.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit

No.	<u>Description</u>
No. 10.1	Securities Purchase Agreement, dated January 11, 2021, between the Company and the Investor (incorporated by reference to Exhibit 10.1
	of the Company's Current Report on Form 8-K, filed with the SEC on January 12, 2021).
99.1	Press Release issued January 31, 2021.
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURES

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

MEDICINOVA, INC.

DATE: February 1, 2021

y: /s/ Yuichi Iwaki

Yuichi Iwaki, M.D., Ph.D.

President and Chief Executive Officer



MediciNova Announces Closing of US\$20 Million Private Placement Transaction

LA JOLLA, Calif., January 31, 2021 (GLOBE NEWSWIRE) — MediciNova, Inc., a biopharmaceutical company traded on the NASDAQ Global Market (NASDAQ:MNOV) and the JASDAQ Market of the Tokyo Stock Exchange (Code Number: 4875), today announced the closing of the previously announced private placement transaction under a Securities Purchase Agreement, dated January 11, 2021, pursuant to which MediciNova issued US\$20 million in shares of its common stock to 3D Opportunity Master Fund, a fund managed by 3D Investment Partners Pte. Ltd. ("3D").

MediciNova intends to use the proceeds received from the private placement primarily for the following three programs:

- 1) To initiate a new clinical trial of MN-166 (ibudilast) for glioblastoma, which could be a pivotal trial.
- 2) To develop an intravenous formulation of MN-166 (ibudilast), which is ideal for amyotrophic lateral sclerosis (ALS) patients who have difficulty with swallowing.
- 3) To initiate a Phase 2 clinical trial of MN-001 (tipelukast) in nonalcoholic steatohepatitis (NASH).

About 3D Investment Partners

3D Investment Partners Pte. Ltd. is a value-oriented investment manager founded in 2015 and based in Singapore. 3D seeks value investing opportunities through a process of bottom-up fundamental research and analysis. By unlocking value with an emphasis on alignment of interest with the management teams of their portfolio companies, 3D delivers its clients superior long-term compounding returns with the spirit of "Sampo Yoshi" – a Japanese business core value that one should do business in a way that is good for all three parties: the seller, the buyer, and society at large. 3D, together with its portfolio company's management, pursues the same goal of increasing value for all stakeholders.

About MediciNova

MediciNova, Inc. is a publicly-traded biopharmaceutical company founded upon developing novel, small-molecule therapeutics for the treatment of diseases with unmet medical needs with a primary commercial focus on the U.S. market. MediciNova's current strategy is to focus on BC-PIV SARS-COV-2 vaccine for COVID-19, MN-166 (ibudilast) for neurological disorders such as progressive multiple sclerosis (MS),

amyotrophic lateral sclerosis (ALS), degenerative cervical myelopathy (DCM), substance dependence (e.g., alcohol use disorder, methamphetamine dependence, opioid dependence) and glioblastoma (GBM), as well as prevention of acute respiratory distress syndrome (ARDS) caused by COVID-19, and MN-001 (tipelukast) for fibrotic diseases such as nonalcoholic steatohepatitis (NASH) and idiopathic pulmonary fibrosis (IPF). MediciNova's pipeline also includes MN-221 (bedoradrine) and MN-029 (denibulin). For more information on MediciNova, Inc., please visit www.medicinova.com.

Statements in this press release that are not historical in nature constitute forward-looking statements within the meaning of the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. These forward-looking statements include, without limitation, statements regarding the future development and efficacy of BC-PIV SARS-COV-2 vaccine, MN-166, MN-001, MN-221, and MN-029. These forward-looking statements may be preceded by, followed by, or otherwise include the words "believes," "expects," "anticipates," "intends," "estimates," "projects," "can," "could," "may," "will," "would," "considering," "planning" or similar expressions. These forwardlooking statements involve a number of risks and uncertainties that may cause actual results or events to differ materially from those expressed or implied by such forward-looking statements. Factors that may cause actual results or events to differ materially from those expressed or implied by these forward-looking statements include, but are not limited to, risks of obtaining future partner or grant funding for development of BC-PIV SARS-COV-2 vaccine, MN-166, MN-001, MN-221, and MN-029 and risks of raising sufficient capital when needed to fund MediciNova's operations and contribution to clinical development, risks and uncertainties inherent to the development of formulations as well as the initiation and conduct of clinical trials, including the potential cost, expected timing and risks associated with clinical trials designed to meet FDA guidance and the viability of further development considering these factors, product development and commercialization risks, the uncertainty of whether the results of clinical trials will be predictive of results in later stages of product development, the risk of delays or failure to obtain or maintain regulatory approval, risks associated with the reliance on third parties to sponsor and fund clinical trials, risks regarding intellectual property rights in product candidates and the ability to defend and enforce such intellectual property rights, the risk of failure of the third parties upon whom MediciNova relies to conduct its clinical trials and manufacture its product candidates to perform as expected, the risk of increased cost and delays due to delays in the commencement, enrollment, completion or analysis of clinical trials or significant issues regarding the adequacy of clinical trial designs or the execution of clinical trials, and the timing of expected filings with the regulatory authorities, MediciNova's collaborations with third parties, the availability of funds to complete product development plans and MediciNova's ability to obtain third party funding for programs and raise sufficient capital when needed, and the other risks and uncertainties described in MediciNova's filings with the Securities and Exchange Commission, including its annual report on Form 10-K for the year ended December 31, 2019 and its subsequent periodic reports on Form 10-Q and current reports on Form 8-K. Undue reliance should not be placed on these forward-looking statements, which speak only as of the date hereof. MediciNova disclaims any intent or obligation to revise or update these forward-looking statements.

INVESTOR CONTACT:

Geoff O'Brien Vice President MediciNova, Inc. info@medicinova.com