

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): **May 16, 2022**

Dermata Therapeutics, Inc.

(Exact name of registrant as specified in its charter)

Delaware

(State or Other Jurisdiction
of Incorporation)

001-40739

(Commission
File Number)

86-3218736

(I.R.S. Employer
Identification No.)

3525 Del Mar Heights Rd., #322

San Diego, CA 92130

(Address of principal executive offices, including zip code)

(858) 800-2543

(Registrant's telephone number, including area code)

N/A

(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- 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))

Securities registered pursuant to Section 12(b) of the Act:

Title of Each Class:	Trading Symbol	Name of Each Exchange on which Registered
Common Stock, par value \$0.0001 per share	DRMA	The Nasdaq Capital Market
Warrants, exercisable for one share of Common Stock	DRMAW	The Nasdaq Capital Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (17 CFR §230.405) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR §240.12b-2).

Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 2.02. Results of Operations and Financial Condition.

On May 16, 2022, Dermata Therapeutics, Inc. (the “Company”) issued a press release disclosing certain information regarding its results of operations for the fiscal period ended March 31, 2022. A copy of the press release is furnished under Item 2.02 as Exhibit 99.1.

The information included in this Item 2.02, and Exhibit 99.1 to this Current Report on Form 8-K, shall not be deemed “filed” for the purposes of or otherwise subject to the liabilities under Section 18 of the Securities Exchange Act of 1934 as amended (the “Exchange Act”). Unless expressly incorporated into a filing of the Company under the Securities Act of 1933, as amended, or the Exchange Act made after the date hereof, the information contained in this Item 2.02 and Exhibit 99.1 hereto shall not be incorporated by reference into any filing of the Company, whether made before or after the date hereof, regardless of any general incorporation language in such filing.

Item 7.01. Regulation FD Disclosure.

See “Item 2.02 Results of Operations and Financial Condition” above.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No.	Description
99.1	Press Release, dated May 16, 2022, issued by Dermata Therapeutics, Inc. entitled “Dermata Therapeutics Provides Corporate Update and Reports First Quarter 2022 Financial Results.”
104	Cover Page Interactive Data File (embedded within the Inline XBRL document).

Signature

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.

DERMATA THERAPEUTICS, INC.

Dated: May 16, 2022

By: /s/ Gerald T. Proehl
Gerald T. Proehl
Chief Executive Officer



Dermata Therapeutics Provides Corporate Update and Reports First Quarter 2022 Financial Results

*Dermata completes a \$5.0 million private placement financing
DMT310 Phase 2 rosacea trial topline results expected in H2 2022*

SAN DIEGO, CA, May 16, 2022— Dermata Therapeutics, Inc. (Nasdaq: DRMA; DRMAW) (“Dermata” or the “Company”), a clinical-stage biotechnology company focusing on the treatment of medical and aesthetic skin conditions, today highlighted recent corporate progress and reported financial results for the quarter ended March 31, 2022.

“We are excited to be 90% enrolled in our Phase 2 moderate-to-severe rosacea trial using once weekly treatments of DMT310. Once completed, we expect to receive topline results in the second half of 2022,” commented Gerry Proehl, Dermata’s Chairman, President, and Chief Executive Officer. “We are also happy to report that we recently closed a \$5.0 million private placement financing priced at-the-market under Nasdaq rules. This additional capital will help us prepare for an end of Phase 2 meeting with the FDA for DMT310 for the treatment of moderate-to-severe acne in anticipation of the Phase 3 program,” concluded Mr. Proehl.

Corporate Highlights

- **In April 2022, Dermata successfully closed a \$5.0 million private placement priced at-the-market under Nasdaq rules.** The Company closed a private placement with a single institutional investor of 898,585 shares (“Shares”) of the Company’s common stock, pre-funded warrants to purchase up to 2,875,000 shares of common stock and warrants to purchase up to 3,773,585 shares of common stock, for aggregate gross proceeds of \$5.0 million, before deducting the placement agent’s fees and other offering expenses payable by the Company.
- **Dermata initiated the nonclinical program for DMT310 in preparation for requesting an end of Phase 2 meeting with the Food and Drug Administration (FDA).** In February 2022, the Company received feedback from the FDA on the nonclinical studies and the pharmacokinetic (PK) study that must be conducted at this stage of development for DMT310. The Company has initiated the required nonclinical studies and plans to initiate the PK study in the next months with the plan to complete these required studies and request an end of Phase 2 meeting with the FDA in H1 2023.

Anticipated Upcoming Milestones

- **DMT310 Phase 2 results in moderate-to-severe rosacea.** In November 2021, the Company enrolled its first patient in a Phase 2 rosacea trial. As of May 16, 2022, the trial is 90% enrolled, with the last patient expected to be enrolled in the first half of 2022. The Company expects to receive topline results in the second half of 2022. Upon successful results, the Company will look to request an end of phase 2 meeting with the FDA. The trial is a 12-week, double-blinded, randomized, placebo-controlled study with approximately 180 patients expected to be enrolled at 20 clinical sites in the United States. The co-primary endpoints are (i) absolute reduction in inflammatory lesion count and (ii) Investigator Global Assessment (IGA), which will be graded on a 5-point scale (0-4). To be considered a responder, a patient needs to have at least a 2-grade reduction and an IGA score of 0 or 1.
- **DMT310 Phase 3 trials in moderate-to-severe acne.** After the end of Phase 2 meeting with FDA planned for the first half of 2023, the Company plans to initiate the Phase 3 program. Results from the Phase 3 program are expected in the second half of 2024, with an intended filing of the new drug application approximately 6 months after successful completion of the Phase 3 program.

First Quarter 2022 Financial Results

As of March 31, 2022, Dermata had \$8.2 million in cash, compared to \$10.8 million as of December 31, 2021. Dermata received gross proceeds of \$5.0 million in April 2022 from a private placement of its securities, which, together with existing cash resources, are expected to fund operations into the second quarter of 2023.

Research and development expenses were \$1.6 million for the quarter ended March 31, 2022, compared to \$0.7 million for the quarter ended March 31, 2021. The increase in research and development expenses was due to increased clinical trial and manufacturing costs, as well as increases in salaries, offset by decreased stock-based compensation expense. Stock-based compensation expense attributable to research and development totaled \$0.06 million for the quarter ended March 31, 2022 compared to \$0.3 million for the quarter ended March 31, 2021.

General and administrative expenses were \$1.2 million for the quarter ended March 31, 2022, compared to \$1.6 million for the quarter ended March 31, 2021. The decrease in general and administrative expenses was due to decreased legal fees and stock-based compensation expense, offset by increased insurance and public company costs. Stock-based compensation expense attributable to general and administrative totaled \$0.2 million for the quarter ended March 31, 2022 compared to \$0.9 million for the quarter ended March 31, 2021.

About Dermata Therapeutics

Dermata Therapeutics, Inc. is a clinical-stage biotechnology company focusing on the treatment of medical and aesthetic skin conditions. The Company's lead product candidate, DMT310, is the first product candidate being developed from its *Spongilla* technology platform. DMT310 is a once weekly topical product candidate derived from a naturally sourced freshwater sponge with multiple unique mechanisms of action. DMT310 is currently under clinical development for the treatment of acne, psoriasis, and rosacea. The Company's second product candidate, DMT410, uses its *Spongilla* technology as a new method for topical intradermal delivery of botulinum toxin for the treatment of multiple aesthetic skin conditions. Dermata is headquartered in San Diego, California. For more information, please visit <http://www.dermatarx.com/>.

Forward-Looking Statements

Statements in this press release that are not strictly historical in nature are forward-looking statements. These statements are based on the Company's current beliefs and expectations and new risks may emerge from time to time. Forward-looking statements are subject to known and unknown risks, uncertainties, assumptions and other factors including, but are not limited to, statements related to: expectations with regard to the timing of data events; expectations with regard to the timing and/or results from meetings with regulatory bodies; expectations with regard to any potential partnership opportunities for the Company's product candidates; the Company's expectations with regard to current cash and the amount of time it will fund operations; the success, cost, and timing of its product candidates DMT310 and DMT410 development activities and ongoing and planned clinical trials; and whether the results of DMT310 or DMT410 will lead to future product development. These statements are only predictions based on current information and expectations and involve a number of risks and uncertainties. Actual events or results may differ materially from those projected in any of such statements due to various factors, including the risks and uncertainties inherent in drug development, approval and commercialization, and the fact that past results of clinical trials may not be indicative of future trial results. For a discussion of these and other factors, please refer to Dermata's filings with the Securities and Exchange Commission. You are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date hereof. This caution is made under the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. All forward-looking statements are qualified in their entirety by this cautionary statement and Dermata undertakes no obligation to revise or update this press release to reflect events or circumstances after the date hereof, except as required by law.

DERMATA THERAPEUTICS, INC.
(Formerly Dermata Therapeutics, LLC)
Balance Sheets

In thousands, except share and per share data

	March 31, 2022	December 31,
	(unaudited)	2021
Assets		
Cash	\$ 8,191	\$ 10,799
Prepaid expenses and other current assets	853	825
Total assets	9,044	11,624
Liabilities		
Accounts payable	662	515
Accrued liabilities	529	1,002
Total liabilities	1,191	1,517
Equity	7,853	10,107
Total liabilities and equity	\$ 9,044	\$ 11,624

DERMATA THERAPEUTICS, INC.
(Formerly Dermata Therapeutics, LLC)
Statements of Operations

	Three Months Ended March 31,	
	2022	2021
	(unaudited)	(unaudited)
<i>In thousands, except share and per share data</i>		
Operating expenses		
Research and development (1)	\$ 1,596	\$ 681
General and administrative (1)	1,190	1,581
Total operating expenses	2,786	2,262
Loss from operations	(2,786)	(2,262)
Interest expense, net	-	43
Net loss	\$ (2,786)	\$ (2,305)
Net loss per common share, basic and diluted	\$ (0.33)	\$ (1.21)
Weighted average common shares outstanding, basic and diluted	8,352,459	1,911,009
 (1) Includes the following stock-based compensation expense (in thousands)		
Research and development	\$ 55	\$ 250
General and administrative	\$ 158	\$ 910

Investors:
Sean Proehl
Senior Director, Legal and Business Development
info@dermatarx.com