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): November 10, 2022

Journey Medical Corporation

(Exact Name of Registrant as Specified in Charter)

Delaware (State or Other Jurisdiction of Incorporation)

001-41063 (Commission File Number)

47-1879539 (IRS Employer Identification No.)

9237 E Via de Ventura Blvd, Suite 105 Scottsdale, AZ 85258 (Address of Principal Executive Offices)

(480) 434-6670

(Registrant's telephone number, including area code)

Common Stock	DERM	Nasdag Capital Market
Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Securities registered pursuant to Section 12(b) of the Act:		
$\hfill\square$ Pre-commencement communications pursuant to Rule 13e-4(c) u	nder the Exchange Act.	
$\hfill\square$ Pre-commencement communications pursuant to Rule 14d-2b un	der the Exchange Act.	
☐ Soliciting material pursuant to Rule 14a-12 under the Exchange	Act.	
\square Written communications pursuant to Rule 425 under the Securities	es Act.	
Check the appropriate box below if the Form 8-K ming is intended to	o simultaneously satisfy the filling obliga	ation of the registrant under any of the following provisions.

	Title of each class	rrading Symbol(s)	Name of each exchange on which registered	1
	Common Stock	DERM	Nasdaq Capital Market	Ī
,			•	•

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

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 November 10, 2022, Journey Medical Corporation (the "Company" or "Journey") issued a press release to provide a corporate update and to announce its financial results for the three months ended September 30, 2022. A copy of such press release is being furnished as Exhibit 99.1 to this report.

The information, including Exhibit 99.1, in this Form 8-K is being furnished and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that Section. The information in this Form 8-K shall not be incorporated by reference into any filing under the Securities Act of 1933, as amended, except as shall otherwise be expressly set forth by specific reference in such filing.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits.

The following exhibits are furnished herewith:

Description

Ex	hib	it
Nu	mb	er

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99.1	Press release issued by Journey Medical Corporation, dated November 10, 2022
JJ.1	1 1035 Telease issued by Journey Medical Corporation, dated Novelliber 10, 2022

Cover Page Interactive Data File, formatted in Inline Extensible Business Reporting Language (iXBRL).

SIGNATURES

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.

Date: November 10, 2022

Journey Medical Corporation (Registrant)

By: /s/ Claude Maraoui
Claude Maraoui
Chief Executive Officer, President and Director



Journey Medical Corporation Reports Third Quarter 2022 Financial Results and Recent Corporate Highlights

Revenue for the first nine months of 2022 increased 26% to \$57.7 million versus the same period in 2021 of \$45.6 million

DFD-29 Phase 3 studies are 75% enrolled to date

Top-line data from the Phase 3 clinical program evaluating DFD-29 for the treatment of papulopustular rosacea anticipated in the first half of 2023

Company to hold conference call on November 10, 2022 at 4:30 p.m. ET

Scottsdale, AZ – November 10, 2022 – Journey Medical Corporation (Nasdaq: DERM) ("Journey Medical" or the "Company"), a commercial-stage pharmaceutical company that focuses on the development and commercialization of pharmaceutical products for the treatment of dermatological conditions, today announced financial results and recent corporate highlights for the third quarter and nine months ended September 30, 2022.

Claude Maraoui, Journey Medical's Co-Founder, President and Chief Executive Officer said, "While our sales revenue for the third quarter of 2022 of \$16.1 continues to be negatively impacted primarily by the effect of Targadox generic competition versus the third quarter of 2021, we still expect to report record revenue for the full year 2022. Also, we are pleased to have enrolled 75% of patients throughout the U.S. and Europe in our two DFD-29 Phase 3 studies for the treatment of papulopustular rosacea. We anticipate announcing top-line data from the trials in the first half of 2023 and expect to file a New Drug Application ("NDA") shortly thereafter in the second half of 2023. After approval, we anticipate that DFD-29 will achieve peak annual net sales exceeding \$100 million. We believe that this program, together with our eight branded and three authorized generic products that help treat common skin conditions, position Journey Medical for sales growth in the coming years. Journey Medical also expects to launch another product in the upcoming months."

Financial Results:

- Revenues were \$16.1 million for the third quarter of 2022, compared to revenues of \$19.6 million for the third quarter of 2021, representing a decline of \$3.5 million. The decline in revenue was primarily attributed to a combination of, the continued generic competition of Targadox, that represented a \$4.0 million reduction versus the prior year, and a \$0.6 million increase in the net revenue of Accutane, which was 17% favorable versus the third quarter of 2021.
- · Selling, general and administrative expenses were \$15.6 million for the third quarter of 2022, compared to \$10.8 million for the third quarter of 2021, with the increase primarily resulting from the expansion of the sales force, marketing expenses related to the expanded product portfolio of four products (Accutane, Qbrexza, Amzeeq and Zilxi), and professional fees associated with being a public company.
- · Research and development expenses were \$2.8 million in the third quarter of 2022, compared to \$0.7 million in the third quarter of 2021, reflecting ongoing clinical trial expenses to develop DFD-29. These expenses have increased and are a result of the enrollment of additional patients in the two Phase 3 trials.
- · Net loss was (\$10.1) million, or (\$0.57) per share basic and diluted, for the third quarter of 2022, compared to a net loss of (\$10.6) million, or (\$1.16) basic and diluted per share, for the third quarter of 2021.
- Non-GAAP Adjusted EBITDA (Adjusted Operating Net Income (loss)) was (\$4.0) million, or (\$0.23) per share basic and diluted, for the third quarter of 2022, compared to Non-GAAP Adjusted EBITDA (Adjusted Operating Net Income of \$1.3 million, or \$0.14 per share basic and \$0.12 per share diluted, for the third quarter of 2021. Non-GAAP Adjusted EBITDA (Adjusted Operating Net Income (loss)) and Non-GAAP Adjusted EBITDA (Adjusted Operating Net Income (loss)) per share basic and diluted are non-GAAP financial measures, each of which are reconciled to the most directly comparable financial measures calculated in accordance with GAAP below, under the heading "Reconciliation of GAAP to Non-GAAP Adjusted EBITDA (Adjusted Operating Net Income (loss))."
- At September 30, 2022, cash and cash equivalents totaled \$34.9 million, compared to \$49.1 million on December 31, 2021.

Recent Corporate Highlights:

- To date, Journey Medical has achieved 75% enrollment in its DFD-29 Phase 3 clinical program for the treatment of papulopustular rosacea. The two DFD-29 Phase 3 clinical studies are still on target to complete enrollment in 2022, with top-line data readout expected in the first half of 2023. Journey Medical plans to submit the NDA for DFD-29 in the second half of 2023 and FDA approval is anticipated in the second half of 2024. In the Phase 2 clinical trials, DFD-29 (40mg) demonstrated nearly double the efficacy when compared against Oraycea® (European equivalent of Oracea®) on both co-primary endpoints. For the first co-primary endpoint, Investigator's Global Assessment ("IGA") treatment success, Oraycea only had a 33.33% IGA treatment success rate, while DFD-29 achieved a 66.04% IGA treatment success rate. For the second co-primary endpoint, the change in total inflammatory lesion count, Oraycea only had a 10.5 reduction in inflammatory lesions, while DFD-29 achieved a 19.2 reduction in inflammatory lesions.
- · In August 2022, \$5.0 million was drawn from the Company's term loan facility with East West Bank. The additional \$5.0 million is part of the Company's operating plan and additional working capital.
- Wire Security Fraud update Regarding the cybersecurity breach that resulted in losses of \$9.5 million in September of 2021, the federal government has been able to seize a significant amount of cryptocurrency assets associated with the breach. Once the cryptocurrency has been converted back into U.S. dollars, Journey Medical will receive a notification letter to initiate the return of the cash to the Company. This process could take as long as 6 months or more to complete.

Conference Call and Webcast Information

To listen to the conference call, interested parties within the U.S. should dial 1-866-777-2509 (domestic) or 1-412-317-5413 (international). All callers should dial in approximately 10 minutes prior to the scheduled start time and ask to be joined into the Journey Medical conference call. Participants can register for the conference here: https://dpregister.com/sreg/10172735/f4f2b41486. Please note that registered participants will receive their dial-in number upon registration.

A live audio webcast can be accessed on the News and Events page of the Investors section of Journey Medical's website, www.journeymedicalcorp.com, and will remain available for replay for approximately 30 days after the meeting.

About Journey Medical Corporation

Journey Medical Corporation (Nasdaq: DERM) ("Journey Medical") is focused on identifying, acquiring, developing and strategically commercializing innovative, differentiated dermatology products through its efficient sales and marketing model. The company currently markets eight products that help treat and heal common skin conditions. The Journey Medical team is comprised of industry experts with extensive experience commercializing some of the most successful prescription dermatology brands. Journey Medical is located in Scottsdale, Arizona and was founded by Fortress Biotech, Inc. (Nasdaq: FBIO). Journey Medical's common stock is registered under the Securities Exchange Act of 1934, as amended, and it files periodic reports with the U.S. Securities and Exchange Commission ("SEC"). For additional information about Journey Medical, visit www.journeymedicalcorp.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, as amended. As used below and throughout this press release, the words "we", "us" and "our" may refer to Journey Medical. Such statements include, but are not limited to, any statements relating to our growth strategy and product development programs and any other statements that are not historical facts. The words "anticipate," "believe," "estimate," "may," "expect," "will," "could," "project," "intend" and similar expressions are generally intended to identify forward-looking statements. Forwardlooking 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 include: risks relating to our growth strategy; our ability to obtain, perform under and maintain financing and strategic agreements and relationships; risks relating to the results of research and development activities; uncertainties relating to preclinical and clinical testing; risks relating to the timing of starting and completing clinical trials, including disruptions that may result from hostilities in Europe; our dependence on third-party suppliers; risks relating to the COVID-19 outbreak and its potential impact on our employees' and consultants' ability to complete work in a timely manner and on our ability to obtain additional financing on favorable terms or at all; our ability to attract, integrate and retain key personnel; the early stage of products under development; our need for substantial additional funds; government regulation; patent and intellectual property matters; competition; potential recovery of funds lost from previously disclosed cyber security breaches; as well as other risks described in Part I, Item 1A, "Risk Factors," in our Annual Report on Form 10-K filed on March 28, 2022, subsequent Reports on Form 10-Q, and our other filings we make with the SEC. 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 may be required by law, and we claim the protection of the safe harbor for forward-looking statements contained in the Private Securities Litigation Reform Act of 1995.

Company Contact:

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JOURNEY MEDICAL CORPORATION

Unaudited Condensed Consolidated Balance Sheets

(Dollars in thousands except for share and per share amounts)

	September 30, 2022	Ε	December 31, 2021	
ASSETS				
Current assets				
Cash and cash equivalents	\$ 34,8	91 \$	49,081	
Accounts receivable, net of reserves	28,5	33	23,112	
Inventory	15,2	30	9,862	
Prepaid expenses and other current assets	9	42	2,438	
Total current assets	79,5	96	84,493	
Intangible assets, net	28,4	24	12,552	
Operating lease right-of-use asset, net	,	22	89	
Other assets	1	03	150	
Total assets	\$ 108,1	45 \$	97,284	
LIABILITIES AND STOCKHOLDERS' EQUITY				
Current liabilities				
Accounts payable	\$ 33,6	26 \$	22,812	
Due to related party		74	641	
Accrued expenses	17,7	33	22,733	
Accrued interest	1	25	-	
Income taxes payable		22	8	

Line of credit	-	812
Deferred cash payment (net of discount of \$76)	4,924	-
Installment payments – licenses, short-term	4,198	4,510
Operating lease liability	25	98
Total current liabilities	60,777	 51,614
Term loan (net of debt discount of \$190)	19,810	-
Installment payments – licenses, long-term	1,374	3,627
Total liabilities	81,961	 55,241
Stockholders' equity		
Common stock, \$.0001 par value, 50,000,000 shares authorized, 11,642,659 and 11,316,344 shares issued and outstanding as		
of September 30, 2022 and December 31, 2021, respectively	1	1
Common stock - Class A, \$.0001 par value, 50,000,000 shares authorized, 6,000,000 shares issued and outstanding as of		
September 30, 2022 and December 31, 2021	1	1
Additional paid-in capital	84,042	80,915
Accumulated deficit	(57,860)	(38,874)
Total stockholders' equity	26,184	42,043
Total liabilities and stockholders' equity	\$ 108,145	\$ 97,284

JOURNEY MEDICAL CORPORATION

Unaudited Condensed Consolidated Statements of Operations

(Dollars in thousands except for share and per share amounts)

		Three-Month Periods Ended Nine- September 30,				ine-Month Periods Ended September 30,		
		2022		2021		2022		2021
Revenue:								
Product revenue, net	\$	16,043	\$	19,610	\$	55,074	\$	45,617
Other revenue		73		-		2,629		-
Total Revenue		16,116		19,610		57,703		45,617
Operating expenses								
Cost of goods sold – product revenue		7,221		11,167		23,057		22,559
Research and development		2,812		718		6,687		747
Research and development - licenses acquired		-		76		· -		13,819
Selling, general and administrative		15,575		10,755		45,481		24,776
Wire transfer fraud loss		-		9,540		-		9,540
Total operating expenses		25,608		32,256		75,225		71,441
Loss from operations	_	(9,492)		(12,646)		(17,522)		(25,824)
Other expense								
Interest income		(3)		-		(10)		-
Interest expense		559		1,373		1,402		2,936
Foreign exchange transaction losses		22		-		22		-
Change in fair value of derivative liability		-		2		-		184
Total other expense		578		1,375		1,414		3,120
Loss before income taxes		(10,070)		(14,021)		(18,936)		(28,944)
Income tax expense (benefit)		10		(3,375)		50		(6,701)
Net Loss	\$	(10,080)	\$	(10,646)	\$	(18,986)	\$	(22,243)
	_							
Net loss per common share:								
Basic and diluted	\$	(0.57)	\$	(1.16)	\$	(1.09)	\$	(2.43)
Weighted average number of common shares:		4 - 640 064		0.464.000				0.450.044
Basic and diluted		17,618,064		9,161,333		17,464,561		9,160,344

Use of Non-GAAP Measures:

In addition to the GAAP financial measures as presented in our Form 10-Q that will be filed with the Securities and Exchange Commission ("SEC"), the Company has, in this press release, included certain non-GAAP measurements, including Adjusted EBITDA (Adjusted Operating Net Income (loss)), Adjusted Operating Net Income (loss) per share basic and Adjusted Net Income (loss) per share diluted. We define Adjusted EBITDA (Adjusted Operating Net Income (loss)) as net income (loss) excluding interest, taxes and depreciation, less certain other non-cash items, namely, share-based compensation expense, amortization of acquired intangible assets, inventory step-ups from the purchases of intangibles assets and products, as more fully described as follows:

- Share-Based Compensation Expense: We exclude share-based compensation from our adjusted financial results because share-based compensation expense, which is non-cash, fluctuates from period to period based on factors that are not within our control, such as our stock price on the dates share-based grants are issued.
- · Non-core and Short-term Research and Development Expense: We exclude costs associated with non-core and short-term related research and development because we do not consider such costs to be normal, recurring operating expenses that are core to our long-term strategy.

Amortization of Acquired Intangible assets: We exclude the impact of certain amounts recorded in connection with the acquisitions of intangible assets that are either non-cash or not normal, recurring operating expenses due to their nature, variability of amounts, and lack of predictability as to occurrence and/or timing. These amounts may include non-cash items such as the amortization of acquired intangible assets and amortization of step-ups of acquisition accounting adjustments to inventories.

Adjusted Operating Net Income (loss) per share basic and Adjusted Net Income (loss) per share diluted are determined by dividing the resulting Adjusted EBITDA (Adjusted Operating Net Income (loss)) by the number of shares outstanding on an actual and fully diluted basis.

Management believes use of these non-GAAP measures provide meaningful supplemental information regarding the Company's performance because (i) it allows for greater transparency with respect to key measures used by management in its financial and operational decision-making, (ii) it excludes the impact of non-cash or, when specified, non-recurring items that are not directly attributable to the Company's core operating performance and that may obscure trends in the Company's core operating performance and (iii) it is used by institutional investors and the analyst community to help analyze the Company's results. However, Adjusted EBTIDA (Adjusted Operating Net Income (loss)), Adjusted Operating Net Income (loss) per share basic, Adjusted Net Income (loss) per share diluted and any other non-GAAP financial measures should be considered as a supplement to, and not as a substitute for, or superior to, the corresponding measures calculated in accordance with GAAP. Further, non-GAAP financial measures used by the Company and the manner in which they are calculated may differ from the non-GAAP financial measures or the calculations of the same non-GAAP financial measures used by other companies, including the Company's competitors.

The table below provides a reconciliation from GAAP to non-GAAP measures:

JOURNEY MEDICAL CORPORATION Reconciliation of GAAP to Non-GAAP Adjusted EBITDA (Adjusted Operating Net Income (loss))

(Dollars in thousands except for share and per share amounts)

CAAP Net Loss CAAP Net Los		Three-month periods ended September 30,			Nine-month periods ended September 30,			
Taxes 10 (3,375) 50 (6,701)		2022		2021		2022		2021
Interest 556 1,373 1,392 2,936 Taxes 10 (3,375) 50 (6,701) Depreciation - - - Amortization of acquired intangible assets 1,016 658 3,050 1,983 EBITDA (8,498) (11,990) (14,494) (24,025) Non-GAAP Adjusted EBITDA (Adjusted Operating Net (loss) gain): Share-based compensation 1,438 8 2,985 41 Change in fair value of derivative liabilities - 2 - 184 Inventory step-up expense 214 3,001 525 4,239 Wire transfer fraud loss - 9,540 - 9,540	GAAP Net Loss	\$ (10,080)	\$	(10,646)	\$	(18,986)	\$	(22,243)
Interest 556 1,373 1,392 2,936 Taxes 10 (3,375) 50 (6,701) Depreciation - - - Amortization of acquired intangible assets 1,016 658 3,050 1,983 EBITDA (8,498) (11,990) (14,494) (24,025) Non-GAAP Adjusted EBITDA (Adjusted Operating Net (loss) gain): Share-based compensation 1,438 8 2,985 41 Change in fair value of derivative liabilities - 2 - 184 Inventory step-up expense 214 3,001 525 4,239 Wire transfer fraud loss - 9,540 - 9,540								
Taxes 10 (3,375) 50 (6,701) Depreciation - <td< td=""><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></td<>								
Depreciation				,				,
Amortization of acquired intangible assets 1,016 658 3,050 1,983		10		(3,375)		50		(6,701)
EBITDA (8,498) (11,990) (14,494) (24,025) Non-GAAP Adjusted EBITDA (Adjusted Operating Net (loss) gain): Share-based compensation 1,438 8 2,985 41 Change in fair value of derivative liabilities - 2 - 184 Inventory step-up expense 214 3,001 525 4,239 Wire transfer fraud loss - 9,540 - 9,540		-		-		-		-
Non-GAAP Adjusted EBITDA (Adjusted Operating Net (loss) gain): Share-based compensation 1,438 8 2,985 41 Change in fair value of derivative liabilities - 2 - 184 Inventory step-up expense 214 3,001 525 4,239 Wire transfer fraud loss - 9,540 - 9,540	1 6	 				3,050		1,983
Share-based compensation 1,438 8 2,985 41 Change in fair value of derivative liabilities - 2 - 184 Inventory step-up expense 214 3,001 525 4,239 Wire transfer fraud loss - 9,540 - 9,540	EBITDA	(8,498)		(11,990)		(14,494)		(24,025)
Share-based compensation 1,438 8 2,985 41 Change in fair value of derivative liabilities - 2 - 184 Inventory step-up expense 214 3,001 525 4,239 Wire transfer fraud loss - 9,540 - 9,540								
Change in fair value of derivative liabilities - 2 - 184 Inventory step-up expense 214 3,001 525 4,239 Wire transfer fraud loss - 9,540 - 9,540								
Inventory step-up expense 214 3,001 525 4,239 Wire transfer fraud loss - 9,540 - 9,540	Share-based compensation	1,438				2,985		
Wire transfer fraud loss - 9,540 - 9,540	Change in fair value of derivative liabilities	-		2		-		184
· · · · · · · · · · · · · · · · · · ·	Inventory step-up expense	214		3,001		525		4,239
	Wire transfer fraud loss	-				-		
R&D 2,778 718 6,653 747				718				747
Foreign exchange transaction losses 22 - 22 -	Foreign exchange transaction losses	22		-		22		-
Severance <u>27</u> - <u>27</u> <u>260</u>	Severance	27		<u>-</u>		27		260
Non-GAAP Adjusted EBITDA (Adjusted Operating Net (loss) gain) \$ (4,019) \$ 1,279 \$ (4,282) \$ (9,014)	Non-GAAP Adjusted EBITDA (Adjusted Operating Net (loss) gain)	\$ (4,019)	\$	1,279	\$	(4,282)	\$	(9,014)
							_	
Net loss per common share Basic:	Net loss per common share Basic:							
GAAP Net loss \$ (0.57) \$ (1.16) \$ (1.09) \$ (2.43)		\$ (0.57)	\$	(1.16)	\$	(1.09)	\$	(2.43)
Non-GAAP Net (loss) gain \$ (0.23) \$ 0.14 \$ (0.25) \$ (0.98)	Non-GAAP Net (loss) gain	\$ (0.23)				` /		
· / ·	` , '	` ′				` ′		, ,
Net loss per common share Diluted:	Net loss per common share Diluted:							
GAAP Net loss \$ (0.57) \$ (0.98) \$ (1.09) \$ (2.43)		\$ (0.57)	\$	(0.98)	\$	(1.09)	\$	(2.43)
Non-GAAP Net (loss) gain \$ (0.23) \$ 0.12 \$ (0.25) \$ (0.98)	Non-GAAP Net (loss) gain	\$ (0.23)		0.12	\$	(0.25)		(0.98)
	` , '	` ′				` ′		, í
Weighted average number of common shares Basic: 17,618,064 9,161,333 17,464,561 9,160,344	Weighted average number of common shares Basic:	17,618,064		9,161,333		17,464,561		9,160,344
Weighted average number of common shares Diluted: 17,618,064 10,892,050 17,464,561 9,160,344	Weighted average number of common shares Diluted:	17,618,064		10,892,050		17,464,561		9,160,344

The weighted average number of shares of common stock outstanding used to calculate both basic and diluted income loss per share is the same for the three-month period ended September 30, 2022 and the nine-month periods ended September 30, 2022 and 2021 since the Company reported a net loss for these periods.