
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): July 27, 2023

VANDA PHARMACEUTICALS INC.

(Exact name of Registrant as specified in its charter)

Delaware
(State or other jurisdiction of incorporation)

001-34186
(Commission File No.)

03-0491827
(IRS Employer Identification No.)

2200 Pennsylvania Avenue NW
Suite 300E
Washington, DC 20037
(Address of principal executive offices and zip code)

Registrant's telephone number, including area code: (202) 734-3400

Not Applicable
(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 (see General Instruction A.2. below):

- 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:

<u>Title of each class</u>	<u>Trading Symbol</u>	<u>Name of each exchange on which registered</u>
Common Stock, par value \$0.001 per share	VNDA	The Nasdaq Global Market

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).

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 July 27, 2023, Vanda Pharmaceuticals Inc. (“Vanda”) issued a press release and is holding a conference call regarding its results of operations and financial condition for the quarter ended June 30, 2023. The full text of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated by reference herein.

Various statements to be made during the conference call are “forward-looking statements” under the securities laws, including, but not limited to, statements regarding Vanda’s commercial products, plans and opportunities, as well as statements about Vanda’s products in development and the related clinical development and regulatory timelines and commercial potential for such products. Words such as, but not limited to, “believe,” “expect,” “anticipate,” “estimate,” “intend,” “plan,” “project,” “target,” “goal,” “likely,” “will,” “would,” and “could,” or the negative of these terms and similar expressions or words, identify forward-looking statements. Forward-looking statements are based upon current expectations and assumptions that involve risks, changes in circumstances and uncertainties.

Important factors that could cause actual results to differ materially from those reflected in Vanda’s forward-looking statements include, among others, Vanda’s assumptions regarding the strength of its business in the U.S. and Vanda’s ability to complete the clinical development of, and obtain regulatory approval for, the products in its pipeline. Therefore, no assurance can be given that the actual results or developments anticipated by Vanda will be realized or, even if substantially realized, that they will have the expected consequences to, or effects on, Vanda. Forward-looking statements made during the call should be evaluated together with the various risks and uncertainties that affect Vanda’s business and market, particularly those identified in the “Cautionary Note Regarding Forward-Looking Statements”, “Risk Factors” and “Management’s Discussion and Analysis of Financial Condition and Results of Operations” sections of Vanda’s most recent Annual Report on Form 10-K, as updated by Vanda’s subsequent Quarterly Reports on Form 10-Q, Current Reports on Form 8-K and other filings with the U.S. Securities and Exchange Commission, which are available at www.sec.gov.

All written and verbal forward-looking statements attributable to Vanda or any person acting on its behalf are expressly qualified in their entirety by the cautionary statements contained or referred to herein. The information contained in this Current Report on Form 8-K is intended to be considered in the context of Vanda’s filings with the SEC and other public announcements that Vanda makes, by press release or otherwise, from time to time. Vanda cautions investors not to rely too heavily on the forward-looking statements Vanda makes or that are made on its behalf. The information conveyed on the conference call will be provided only as of the date of the call, and Vanda undertakes no obligation, and specifically declines any obligation, to update or revise publicly any forward-looking statements made during the call after the date thereof, whether as a result of new information, future events or otherwise, except as required by law.

The information in Item 2.02 of this Current Report on Form 8-K and Exhibit 99.1 attached hereto shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such a filing.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
99.1	Press release of Vanda Pharmaceuticals Inc. dated July 27, 2023.
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.

Dated: July 27, 2023

VANDA PHARMACEUTICALS INC.

By: /s/ Timothy Williams

Name: Timothy Williams

Title: Senior Vice President, General Counsel and Secretary



Vanda Pharmaceuticals Reports Second Quarter 2023 Financial Results

- Q2 2023 total revenues were \$46.1 million
- Total revenues in the first six months of 2023 were \$108.6 million
- Vanda provides update on pipeline advancements and regulatory plans

WASHINGTON – July 27, 2023 – Vanda Pharmaceuticals Inc. (Vanda) (Nasdaq: VNDA) today announced financial and operational results for the second quarter ended June 30, 2023.

"During the second quarter we continued our efforts to protect our commercial interests in the HETLIOZ® franchise in the face of the at-risk generic launch. We recorded a higher number of HETLIOZ® prescription dispenses in the second quarter of 2023 as compared to the first quarter of 2023, a testament to our dedication to our patient services and the loyalty of our patients," said Mihael H. Polymeropoulos, M.D., Vanda's President, CEO and Chairman of the Board. "We also advanced towards our goal of marketing authorizations for Fanapt® in bipolar disorder, HETLIOZ® in insomnia and tridipititon in patients with gastroparesis."

Financial Highlights

Second Quarter of 2023

- Total net product sales from HETLIOZ® and Fanapt® were \$46.1 million in the second quarter of 2023, a 28% decrease compared to \$64.4 million in the second quarter of 2022.
- HETLIOZ® net product sales were \$22.0 million in the second quarter of 2023, a 47% decrease compared to \$41.2 million in the second quarter of 2022. The decrease was the result of the at-risk launch of a generic version of HETLIOZ® in the U.S.
- Fanapt® net product sales were \$24.1 million in the second quarter of 2023, a 4% increase compared to \$23.2 million in the second quarter of 2022.
- Net income was \$1.5 million in the second quarter of 2023 compared to net income of \$2.6 million in the second quarter of 2022.
- Cash, cash equivalents and marketable securities (Cash) was \$489.4 million as of June 30, 2023, representing a decrease to Cash of \$12.1 million, or 2%, compared to March 31, 2023.

First Six Months of 2023

- Total net product sales from HETLIOZ® and Fanapt® were \$108.6 million in the first six months of 2023, a 13% decrease compared to \$124.6 million in the first six months of 2022.
- HETLIOZ® net product sales were \$61.6 million in the first six months of 2023, a 21% decrease compared to \$78.2 million in the first six months of 2022. The decrease was the result of the at-risk launch of a generic version of HETLIOZ® in the U.S.
- Fanapt® net product sales were \$47.0 million in the first six months of 2023, a 1% increase compared to \$46.4 million in the first six months of 2022.
- Net income was \$4.8 million in the first six months of 2023 compared to a net loss of \$3.9 million in the first six months of 2022.

- Cash, cash equivalents and marketable securities (Cash) was \$489.4 million as of June 30, 2023, representing an increase to Cash of \$48.5 million, or 11%, compared to June 30, 2022.

Key Operational Highlights

Fanapt® (iloperidone)

- Vanda previously announced positive results in the Phase III clinical study of Fanapt® in acute manic and mixed episodes associated with bipolar I disorder in adults. Vanda continues to pursue U.S. Food and Drug Administration (FDA) approval of a supplemental New Drug Application (sNDA) for Fanapt® in bipolar I disorder.

HETLIOZ® (tasimelteon)

- Vanda is continuing to pursue FDA approvals for HETLIOZ® in the indications of insomnia and jet lag disorder.
- In May 2023, the U.S. Court of Appeals for the Federal Circuit (the Federal Circuit) affirmed the lower court ruling in favor of Teva Pharmaceuticals USA, Inc. (Teva), Apotex Inc. and Apotex Corp. (Apotex) in the HETLIOZ® Abbreviated New Drug Applications litigation. Vanda disagrees with the Federal Circuit's ruling, and in June 2023, Vanda requested a rehearing from the Federal Circuit. On July 18, 2023, the Federal Circuit requested a response from Teva and Apotex, which is due August 1, 2023.

Tramipitant

- Vanda continues to pursue FDA approval of a New Drug Application (NDA) for tramipitant for patients with gastroparesis.
- Vanda announced positive results in the Phase III study of tramipitant in motion sickness. Vanda plans to continue the motion sickness clinical program and pursue FDA approval upon completion of additional efficacy and safety studies.

Early-Stage Programs

- Vanda announced that the FDA had granted Orphan Drug Designation for VCA-894A for the treatment of Charcot-Marie-Tooth disease, axonal, type 2S (CMT2S), caused by cryptic splice site variants within IGHMBP2.

GAAP Financial Results

Net income was \$1.5 million in the second quarter of 2023 compared to net income of \$2.6 million in the second quarter of 2022. Diluted net income per share was \$0.03 in the second quarter of 2023 compared to diluted net income per share of \$0.05 in the second quarter of 2022.

Net income was \$4.8 million in the first six months of 2023 compared to a net loss of \$3.9 million in the first six months of 2022. Diluted net income per share was \$0.08 in the first six months of 2023 compared to diluted net loss per share of \$0.07 in the first six months of 2022.

2023 Financial Guidance

Given uncertainties surrounding the U.S. market for HETLIOZ® for the treatment of Non-24 as a result of the ongoing HETLIOZ® patent litigation and the at-risk launch of a generic version of HETLIOZ®, Vanda is unable to provide 2023 financial guidance at this time. Vanda will continue to evaluate its ability to provide financial guidance as the year progresses.

HETLIOZ® net product sales will likely decline in future periods, potentially significantly, related to the at-risk launch of a generic version of HETLIOZ® in the U.S.

Conference Call

Vanda has scheduled a conference call for today, Thursday, July 27, 2023, at 4:30 PM ET. During the call, Vanda's management will discuss the second quarter 2023 financial results and other corporate activities. Investors can call 1-800-715-9871 (domestic) or 1-646-307-1963 (international) and use passcode number 4063687. A replay of the call will be available on Thursday, July 27, 2023, beginning at 8:30 PM ET and will be accessible until Thursday, August 3, 2023 at 8:30

PM ET. The replay call-in number is 1-800-770-2030 for domestic callers and 1-609-800-9909 for international callers. The passcode number is 4063687.

The conference call will be broadcast simultaneously on Vanda's website, www.vandapharma.com. Investors should click on the Investors tab and are advised to go to the website at least 15 minutes early to register, download, and install any necessary software or presentations. The call will also be archived on Vanda's website for a period of 30 days.

About Vanda Pharmaceuticals Inc.

Vanda is a leading global biopharmaceutical company focused on the development and commercialization of innovative therapies to address high unmet medical needs and improve the lives of patients. For more on Vanda Pharmaceuticals Inc., please visit www.vandapharma.com and follow us on Twitter @vandapharma.

CAUTIONARY NOTE REGARDING FORWARD LOOKING STATEMENTS

Various statements in this press release, including, but not limited to, statements regarding Vanda's plans for pursuit of FDA approval of Fanapt® in the treatment of bipolar I in adults, HETLIOZ® in the treatments of insomnia and jet lag disorder and tridipitant in the treatment of patients with gastroparesis, and the treatment of motion sickness and Vanda's expectations regarding the impact of generic competition on the HETLIOZ® business are "forward-looking statements" under the securities laws. All statements other than statements of historical fact are statements that could be deemed forward-looking statements. Forward-looking statements are based upon current expectations and assumptions that involve risks, changes in circumstances and uncertainties. Important factors that could cause actual results to differ materially from those reflected in Vanda's forward-looking statements include, among others, Vanda's ability to pursue and obtain FDA approval of the sNDAs for Fanapt® and HETLIOZ® and the NDA for tridipitant, the FDA's assessment of the sufficiency of the data packages to be included in Vanda's regulatory submissions for Fanapt®, HETLIOZ® and tridipitant, Vanda's ability to complete the clinical program for tridipitant in the treatment of motion sickness, the outcome of Vanda's request for a rehearing by the Federal Circuit and Vanda's ability to enforce its legal rights to exclusivity for HETLIOZ®. Therefore, no assurance can be given that the results or developments anticipated by Vanda will be realized or, even if substantially realized, that they will have the expected consequences to, or effects on, Vanda. Forward-looking statements in this press release should be evaluated together with the various risks and uncertainties that affect Vanda's business and market, particularly those identified in the "Cautionary Note Regarding Forward-Looking Statements", "Risk Factors" and "Management's Discussion and Analysis of Financial Condition and Results of Operations" sections of Vanda's most recent Annual Report on Form 10-K, as updated by Vanda's subsequent Quarterly Reports on Form 10-Q, Current Reports on Form 8-K and other filings with the U.S. Securities and Exchange Commission, which are available at www.sec.gov.

All written and verbal forward-looking statements attributable to Vanda or any person acting on its behalf are expressly qualified in their entirety by the cautionary statements contained or referred to herein. Vanda cautions investors not to rely too heavily on the forward-looking statements Vanda makes or that are made on its behalf. The information in this press release is provided only as of the date of this press release, and Vanda undertakes no obligation, and specifically declines any obligation, to update or revise publicly any forward-looking statements, whether as a result of new information, future events or otherwise, except as required by law.

VANDA PHARMACEUTICALS INC.
CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS
(in thousands, except for share and per share amounts)
(unaudited)

	Three Months Ended		Six Months Ended	
	June 30 2023	June 30 2022	June 30 2023	June 30 2022
Revenues:				
HETLIOZ® net product sales	\$ 21,979	\$ 41,188	\$ 61,595	\$ 78,219
Fanapt® net product sales	24,077	23,202	46,959	46,363
Total revenues	46,056	64,390	108,554	124,582
Operating expenses:				
Cost of goods sold excluding amortization	3,499	6,059	8,273	11,724
Research and development	16,647	21,490	35,884	42,459
Selling, general and administrative	28,399	33,001	64,503	73,849
Intangible asset amortization	378	379	757	758
Total operating expenses	48,923	60,929	109,417	128,790
Income (loss) from operations	(2,867)	3,461	(863)	(4,208)
Other income	5,459	329	8,983	434
Income (loss) before income taxes	2,592	3,790	8,120	(3,774)
Provision for income taxes	1,072	1,216	3,348	82
Net income (loss)	<u>\$ 1,520</u>	<u>\$ 2,574</u>	<u>\$ 4,772</u>	<u>\$ (3,856)</u>
Net income (loss) per share, basic	\$ 0.03	\$ 0.05	\$ 0.08	\$ (0.07)
Net income (loss) per share, diluted	\$ 0.03	\$ 0.05	\$ 0.08	\$ (0.07)
Weighted average shares outstanding, basic	57,453,916	56,508,533	57,233,878	56,307,999
Weighted average shares outstanding, diluted	57,535,615	56,821,024	57,469,105	56,307,999

VANDA PHARMACEUTICALS INC.
CONDENSED CONSOLIDATED BALANCE SHEETS
(in thousands)
(unaudited)

	June 30 2023	December 31 2022
ASSETS		
Current assets:		
Cash and cash equivalents	\$ 150,031	\$ 135,029
Marketable securities	339,320	331,830
Accounts receivable, net	33,621	33,512
Inventory	1,080	1,194
Prepaid expenses and other current assets	9,452	17,727
Total current assets	533,504	519,292
Property and equipment, net	2,164	2,573
Operating lease right-of-use assets	7,782	8,400
Intangible assets, net	17,808	18,565
Deferred tax assets	70,476	74,039
Non-current inventory and other	9,948	11,378
Total assets	\$ 641,682	\$ 634,247
LIABILITIES AND STOCKHOLDERS' EQUITY		
Current liabilities:		
Accounts payable and accrued liabilities	\$ 31,929	\$ 45,551
Product revenue allowances	55,468	45,885
Total current liabilities	87,397	91,436
Operating lease non-current liabilities	7,942	8,813
Other non-current liabilities	6,445	6,800
Total liabilities	101,784	107,049
Stockholders' equity:		
Common stock	57	57
Additional paid-in capital	693,835	686,235
Accumulated other comprehensive loss	(865)	(1,193)
Accumulated deficit	(153,129)	(157,901)
Total stockholders' equity	539,898	527,198
Total liabilities and stockholders' equity	\$ 641,682	\$ 634,247

Corporate Contact:

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