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): August 14, 2023

Emmaus Life Sciences, Inc.

(Exact name of registrant as specified in its charter)

Delaware	001-35527	87-0419387
(State or other jurisdiction of incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)
21250 Hawthorne Boulevard, Suite 800, Torran	ce, CA	90503
(Address of principal executive offices)		(Zip Code)
Registran	t's telephone number, including area code (310	214-0065
(Forme	er name or former address, if changed, since las	et report.)
Check the appropriate box below if the Form 8-K filing is intended	led to simultaneously satisfy the filing obligation	on of the registrant under any of the following provisions:
$\hfill \Box$ Written communications pursuant to Rule 425 under the Sec	curities Act (17 CFR 230.425)	
☐ Soliciting material pursuant to Rule 14a-12 under the Excha	ange 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	4(c) under the Exchange Act (17 CFR 240.13a-	-4(c))
Securities registered pursuant to Section 12(b) of the Act:		
Title of each class None	Trading Symbol	Name of each exchange on which registered
Emerging growth company If an emerging growth company, indicate by check mark if the accounting standards provided pursuant to Section 13(a) of the E		ransition period for complying with any new or revised financial
Item 2.02 Results of Operation and Financial Condition.		
On August 14, 2023, Emmaus Life Sciences, Inc. issued a p 30, 2023. A copy of the press release is included as Exhibit 99.1		ns and financial condition as of and for the six months ended June by reference.
	hall it be deemed incorporated by reference in	of Section 18 of the Securities Exchange Act of 1934 ("Exchange any filing under the Securities Act of 1933 or the Exchange Act,
Item 9.01 Financial Statements and Exhibits		
(d) Exhibits		
See the accompanying Index to Exhibits, which information	is incorporated herein by reference.	
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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.

Date: August 14, 2023 Emmaus Life Sciences, Inc.

By: /s/ YASUSHI NAGASAKI

Name: Yasushi Nagasaki Title: Chief Financial Officer

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INDEX TO EXHIBITS

Exhibit	
Number	Description
99.1	August 14, 2023 press release
104	Cover Page Interactive Date File (embedded within Inline XBRL document)
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Emmaus Life Sciences Reports Q2 2023 Financial Results

Record Net Revenues Contributed to Income from Operations Sixth Straight Quarterly Increase in Net Revenues

Torrance CA, August 14, 2023 - Emmaus Life Sciences, Inc. (OTCQX: EMMA), a commercial-stage biopharmaceutical company and leader in the treatment of sickle cell disease, today reported on its financial condition and results of operations as of and for the three and six months ended June 30, 2023.

Recent Highlights

"We are pleased to report record net revenues for the quarter and six months ended June 30 owing to a jump in sales in the Middle East North Africa region and continuing recovery in U.S. sales compared to 2022. As a result, we were able to generate over \$3.3 million in quarterly income from operations. Net revenue growth accelerated in the quarter, and we hope to continue this trend through the end of the year even without regard to the prospects for potential marketing approval of Endari in the Kingdom of Saudi Arabia," remarked Yutaka Niihara, M.D., M.P.H., Chairman and Chief Executive Officer of Emmaus.

Financial and Operating Results

Net Revenues. Net revenues for the three months and six months ended June 30, 2023 were \$10.8 million and \$17.5 million, respectively, compared to \$4.3 million and \$7.5 million, respectively, for same periods in 2022. The increased net revenues were primarily attributable to a \$4.1 million increase in net revenues from sales in the Middle East North Africa (MENA) region in Q2 2023. Net revenues in Q2 and the three months and six months ended June 30, 2023 also were positively affected by increased U.S. sales compared to the same periods in 2022.

Operating Expenses. Total operating expenses for the three months ended June 30, 2023 were \$6.9 million, compared with \$5.3 million for the same periods in 2022. Of the increased operating expenses in Q2 2023, \$0.6 million was attributable to an increase in payroll expenses related to sales personnel and \$1.0 million increase in general and administrative expenses. Total operating expenses for the six months ended June 30, 2023 were \$14.4 million, compared with \$10.6 million for the same period in 2022. The increase was due to a \$1.2 million increase in share-based compensation, a \$0.8 million increase in payroll expenses and a \$0.6 million increase in consulting fees.

Income From Operations. Income from operations for the three months ended June 30, 2023 was \$3.3 million, compared to a loss from operations of \$1.4 million in the same periods in 2022. Income from operations for the six months ended June 30, 2023 increased to \$2.2 million, compared to a loss from operations of \$4.5 million for the same period last year. The increase income from operation resulted from higher new revenues in 2023 compared to 2022. Income from operations in Q2 2023 also increased by \$4.5 million, or 385%, from \$1.2 million loss from operations in Q1 2023 as a result of the increase in net revenues in Q2 2023.

Other Expense. Other expenses decreased to \$4.8 million for the three months ended June 30, 2023, compared to \$7.3 million in the same period in 2022. Other expenses for the six months ended June 30, 2023 increase to \$7.2 million from \$5.8 million in the same period in 2022. Other expenses in Q2 2023 included a decrease of \$2.6 million in change in fair value of embedded conversion option of convertible promissory notes, partially offset by a \$0.5 million increase in interest expense compared to Q2 2022.

Net Loss. For the quarter, the company realized a net loss of \$1.5 million, or \$0.03 per share based on approximately 52.9 million weighted average basic and diluted common shares. This compares to a net loss of \$8.9 million, or \$0.18 per share based on approximately 49.3 million weighted average basic and diluted common shares for the second quarter of 2022. The decrease in net loss was primarily attributable to the increase in income from operations and decrease in other expenses discussed above.

For the six months ended June 30, 2023, the company reported a net loss of \$5.0 million, or \$0.10 per share, based on approximately 51.8 million weighted average basic and diluted common shares. This compares to a net loss of \$10.4 million, or \$0.21 per share, based on approximately 49.3 million weighted average basic and diluted common shares for the six months ended June 30, 2022. The decrease was due to the increase in net revenues, partially offset by the increase in operating expenses.

Liquidity and Capital Resources. On June 30, 2023, the company had cash and cash equivalents of \$1.4 million, compared with \$2.0 million on December 31, 2022.

About Emmaus Life Sciences

Emmaus Life Sciences, Inc. is a commercial-stage biopharmaceutical company and leader in the treatment of sickle cell disease. Endari® (L-glutamine oral powder), indicated to reduce the acute complications of sickle cell disease in adults and children 5 years and older, is approved for marketing in the United States, Israel, Kuwait, Qatar, the United Arab Emirates, Bahrain and Oman and is available on a named patient or early access basis in France, the Netherlands, and the Kingdom of Saudi Arabia, where Emmaus' application for marketing authorization is awaiting final action by the Saudi Food & Drug Authority. For more information, please visit www.emmausmedical.com.

About Endari® (prescription grade L-glutamine oral powder) Endari®, Emmaus' prescription grade L-glutamine oral powder, was approved by the U.S. Food and Drug Administration (FDA) in July 2017 for treating sickle cell disease in adult and pediatric patients five years of age and older.

Indication

Endari® is indicated to reduce the acute complications of sickle cell disease in adult and pediatric patients five years of age and older.

Important Safety Information

The most common adverse reactions (incidence >10 percent) in clinical studies were constipation, nausea, headache, abdominal pain, cough, pain in extremities, back pain, and chest pain.

Adverse reactions leading to treatment discontinuation included one case each of hypersplenism, abdominal pain, dyspepsia, burning sensation, and hot flash.

The safety and efficacy of Endari® in pediatric patients with sickle cell disease younger than five years of age has not been established.

About Sickle Cell Disease

There are approximately 100,000 people living with sickle cell disease (SCD) in the United States and millions more globally. The sickle gene is found in every ethnic group, not just among those of African descent; and in the United States an estimated 1-in-365 African Americans and 1-in-16,300 Hispanic Americans are born with SCD. ¹ The genetic mutation responsible for SCD causes an individual's red blood cells to distort into a "C" or a sickle shape, reducing their ability to transport oxygen throughout the body. These sickled red blood cells break down rapidly, become very sticky, and develop a propensity to clump together, which causes them to become stuck and cause damage within blood vessels. The result is reduced blood flow to distal organs, which leads to physical symptoms of incapacitating pain, tissue and organ damage, and early death.²

¹Source: Data & Statistics on Sickle Cell Disease – National Center on Birth Defects and Developmental Disabilities, Centers for Disease Control and Prevention, December 2020.

Forward-looking Statements

This press release contains forward-looking statements made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995, as amended, including statements regarding the trend in sales in the MENA region and in the U.S. and need for related-party loans or other financing needed to meet our current liabilities and fund our business and operations. These forward-looking statements are subject to numerous assumptions, risks and uncertainties which change over time, including factors disclosed in the company's Annual Report on Form 10-K for the year ended December 31, 2022 and Quarterly Reports on Form 10-Q for the quarters ended March 31, 2023 and June 30, 2023 filed with the Securities and Exchange Commission on March 31, 2023, May 15, 2023 and August 14, 2023, respectively, and actual results may differ materially. Such forward-looking statements speak only as of the date they are made, and Emmaus assumes no duty to update them, except as may be required by law.

Company Contact:

Emmaus Life Sciences, Inc. Willis Lee Chief Operating Officer (310) 214-0065, Ext. 1130 wlee@emmauslifesciences.com

(Financial Tables Follow)

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Emmaus Life Sciences, Inc. Condensed Consolidated Statement of Operations and Comprehensive Income Loss

(In thousands, except share and per share amounts) (Unaudited)

	Three Months Ended June 30			Six Months Ended June 30				
		2023		2022		2023		2022
Revenues, Net	\$	10,759	\$	4,287	\$	17,512	\$	7,521
Cost of Goods Sold		508		396		937		1,403
Gross Profit		10,251		3,891		16,575		6,118
Operating Expenses		6,925		5,331		14,414		10,626
Income (Loss) from Operations		3,326		(1,440)		2,161		(4,508)
Total Other Expenses		(4,842)		(7,270)		(7,155)		(5,847)
Net Loss		(1,482)		(8,892)		(5,009)		(10,434)
Comprehensive Income (Loss)		1,381		(12,664)		(2,504)		(13,518)
Net Loss Per Share	(\$	0.03)	(\$	0.18)	(\$	0.10)	(\$	0.21)
Weighted Average Common Shares Outstanding		52,865,353		49,319,995		51,793,445		49,315,952

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Emmaus Life Sciences, Inc. Condensed Consolidated Balance Sheets (In thousands)

		As of		
	2	ne 30, 2023 audited)	December 31, 2022	
Assets				
Current Assets:				
Cash and cash equivalents	\$	1,361	\$ 2,021	
Accounts receivable, net		5,573	375	
Inventories, net		1,814	2,379	
Prepaid expenses and other current assets		1,099	1,514	

²Source: Committee on Addressing Sickle Cell Disease – A Strategic Plan and Blueprint for Action -- National Academy of Sciences Press, 2020.

Total Current Assets		9,847		6,289
Property and equipment, net		68		75
Equity method investment		18,302		18,828
Right of use assets		2,585		2,799
Investment in convertible bond		19,210		19,971
Other Assets		276		263
Total Assets	\$	50,288	\$	48,225
Liabilities and Stockholders' Deficit				
Current Liabilities:				
Accounts payable and accrued expenses	\$	15,200	\$	13,549
Conversion feature derivative, notes payable		4,217		3,248
Notes payable, current portion		8,462		6,814
Convertible notes payable, net of discount		14,306		14,655
Other current liabilities		19,362		16,057
Total Current Liabilities		61,547		54,323
Notes payable, less current portion		0		380
Other long-term liabilities		23,773		27,613
Total Liabilities		85,320		82,316
Stockholders' Deficit		(35,032)		(34,091)
Total Liabilities & Stockholders' Deficit	•	50,288	e	48,225
Total Liabilities & Stockholders Deficit	Φ	30,288	Φ	40,223