# 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 9, 2023

# **BARINTHUS BIOTHERAPEUTICS PLC**

(Exact name of registrant as specified in its charter)

England and Wales (State or other jurisdiction of incorporation) 001-40367 (Commission File Number) Not Applicable (I.R.S. Employer Identification No.)

Barinthus Biotherapeutics plc Unit 6-10, Zeus Building Rutherford Avenue, Harwell, Didcot, OX11 0DF United Kingdom (Address of principal executive offices, including zip code)

+44 (0) 1865 818 808 (Registrant's telephone number, including area code)

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:

□ 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> American Depositary Shares <u>Trade Symbol(s)</u> BRNS <u>Name of each exchange on which</u> <u>registered</u> The Nasdaq Global Market

Ordinary shares, nominal value £0.000025 per share\*

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 Xiii

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.

\*American Depositary Shares may be evidenced by American Depositary Receipts. Each American Depositary Share represents one (1) ordinary share. Not for trading, but only in connection with the listing of the American Depositary Shares on The Nasdaq Global Market. The American Depositary Shares represent the right to receive ordinary shares and are being registered under the Securities Act of 1933, as amended, pursuant to a separate Registration Statement on Form F-6. Accordingly, the American Depositary Shares are exempt from the operation of Section 12(a) of the Securities Exchange Act of 1934, as amended, pursuant to Rule 12a-8.

## Item 2.02. Results of Operations and Financial Condition.

On November 9, 2023, Barinthus Therapeutics plc (the "Company") provided an update on its financial information and recent corporate developments in the quarter ended September 30, 2023. The full text of the press release issued in connection with the update is furnished as Exhibit 99.1 to this Current Report on Form 8-K.

The information in this Form 8-K (including Exhibit 99.1) 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

<u>99.1</u>	Press Release dated November 9, 2023.
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.

Date: November 9, 2023

Barinthus Biotherapeutics plc

By: /s/ William Enright

William Enright Chief Executive Officer



#### Barinthus Bio Reports Third Quarter 2023 Financial Results and Recent Corporate Developments

OXFORD, United Kingdom, Nov. 09, 2023 (GLOBE NEWSWIRE) – Barinthus Biotherapeutics plc (NASDAQ: BRNS), formerly Vaccitech plc, today announced its financial results for the third quarter of 2023 and provided an overview of its progress. The Company is a clinical-stage biopharmaceutical company developing novel T cell immunotherapeutic candidates designed to guide the immune system to overcome chronic infectious disease, autoimmunity and cancer.

"The third quarter of 2023 was a period of solid progress for the Company. We have since revealed the evolution of the company identity from Vaccitech to Barinthus Bio. This change reflects the evolution of our pipeline beyond vaccines, towards T cell immunotherapeutic candidates designed to guide the capabilities of T cells, to stimulate their disease-fighting capacity or to restore immune balance," said Bill Enright, Chief Executive Officer of Barinthus Bio. "We remain focused on progressing our pipeline, particularly in HBV, for which upcoming preliminary data from both of our Phase 2 trials will be presented at The American Association for the Study of Liver Diseases – The Liver Meeting<sup>®</sup> 2023 this month."

#### **Recent Corporate Developments**

- In November 2023, the Company announced it changed its name to Barinthus Biotherapeutics plc to represent the evolution and expansion of its focus beyond vaccines. The Company's new name takes inspiration from "Barinthus," the mythological navigator who guided King Arthur of Britain by ship to the island of Avalon to be healed when he was wounded. The story of the legendary king being guided to a place of healing is reflective of the way the Company's proprietary platforms and technology are designed to guide the immune system to treat infectious diseases, autoimmunity and cancer.
- On November 6, 2023, the Company held a general meeting where its shareholders approved resolutions granting the Company's board of directors or any duly authorized committee of the board of directors an increase in the authority to allot shares in the Company or grant rights to subscribe for or to convert any security into shares in the Company free from pre-emption rights. Pursuant to such approval, our board of directors was authorized to allot shares up to an aggregate nominal amount of £1,928, free from statutory pre-emption rights.

#### **Upcoming Milestones**

- · In the fourth quarter of 2023, the Company expects to:
  - o Announce interim efficacy data from HBV003 (NCT05343481), a Phase 2b clinical trial of VTP-300, that further evaluates its potential as a component of a functional cure for chronic Hepatitis B.
  - o Announce interim efficacy data from the Phase 2a clinical trial (ACTRN12622000317796) collaboration with Arbutus of VTP-300 in combination with Arbutus' siRNA therapeutic candidate, AB-729 for chronic hepatitis B.
  - o Submit a regulatory submission application in Australia for VTP-1000, the Company's lead SNAP-TI candidate, for the treatment of celiac disease.



#### Third Quarter 2023 Financial Highlights

- Cash position: As of September 30, 2023, the Company had cash and cash equivalents of \$160.3 million, compared to \$173.0 million as of June 30, 2023. The net cash used in operating activities was \$11.2 million, primarily resulting from its net loss of \$14.1 million adjusted by unrealised foreign exchange gains of \$6.2 million, depreciation and amortization of \$1.5 million, and changes in its operating assets and liabilities. Based on current research and development plans, the Company expects its cash runway to fund its operating expenses and capital expenditure requirements into the second quarter of 2025.
- Research and development expenses: Research and development expenses were \$15.1 million in the third quarter of 2023 compared to \$13.5 million in the second quarter of 2023, showing increased spend due to the phasing of ongoing clinical trials. VTP-300 HBV research and development expenses increased as a result of the increased trial cost and manufacturing development costs, VTP-850 Prostate cancer has also increased as a result of the increased trial progresses. The quarter-on-quarter R&D expense per program is outlined in the following table.

	September 30,	June 30,	
Three months ended	2023	2023	Change
	\$000	\$000	\$000
Direct research and development expenses by program:			
VTP-200 HPV	1,288	1,837	(549)
VTP-300 HBV	4,877	3,757	1,120
VTP-600 NSCLC <sup>1</sup>	155	79	76
VTP-850 Prostate cancer	1,724	242	1,482
VTP-1000/VTP-1100 Celiac/HPV Cancer	2,507	3,018	(511)
Other and earlier stage programs	1,069	701	368
Total direct research and development expenses	11,620	9,634	1,986
Indirect research and development expenses:			
Personnel-related (including share-based compensation)	2,711	3,388	(677)
Facility-related	368	202	166
Other internal costs	445	319	126
Total indirect research and development expenses	3,524	3,909	(385)
Total research and development expense	15,144	13,543	1,601

<sup>1</sup> The VTP-600 NSCLC Phase 1/2a trial is sponsored by Cancer Research UK.

- **General and administrative expenses:** General and administrative expenses were \$1.0 million in the third quarter of 2023, compared to \$13.1 million in the second quarter of 2023. The decrease was mainly attributable to the unrealized foreign exchange gain of \$6.6 million in the third quarter of 2023, compared to an unrealized foreign exchange loss of \$4.2 million in the second quarter of 2023. The remainder of the decrease was primarily attributable to a decrease in legal and professional fees of \$0.5 million and decrease in share-based compensation expense.
- **Net loss:** For the third quarter of 2023, the Company generated a net loss attributable to its shareholders of \$14.1 million, or \$0.37 per share on both basic and fully diluted bases, compared to a net loss attributable to shareholders of \$23.8 million, or \$0.62 per share on both basic and fully diluted bases in the second quarter of 2023.



#### **About Barinthus Bio**

We are a clinical-stage biopharmaceutical company developing novel T cell immunotherapeutic candidates designed to guide the immune system to overcome chronic infectious diseases, autoimmunity and cancer. Helping patients and their families is the guiding principle at the heart of Barinthus Bio. The company stands apart through its broad pipeline, built around four proprietary platform technologies; ChAdOx, MVA, SNAP-TI and SNAP-CI. Barinthus Bio is advancing a pipeline of five product candidates across a diverse range of therapeutic areas, including: VTP-300, an immunotherapeutic candidate designed as a potential component of a functional cure for chronic hepatitis B viral (HBV) infection; VTP-200, a non-surgical product candidate for persistent high-risk human papillomavirus (HPV); VTP-1000, an autoimmune candidate designed to utilize the SNAP-TI platform to treat patients with celiac disease; VTP-850, a second-generation immunotherapeutic candidate designed to treat recurrent prostate cancer; and VTP-1100, a preclinical cancer candidate designed to utilize the SNAP-CI platform to treat patients with HPV-related cancer. Barinthus Bio's proven scientific expertise, high-value portfolio and focus on product development uniquely positions the company to navigate towards delivering treatments for patients with infectious diseases, autoimmunity and cancers that have a significant impact on their lives every day. For more information, visit <u>www.barinthusbio.com</u>.

#### **Forward Looking Statements**

This press release contains forward-looking statements, within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, which can generally be identified as such by use of the words "would," "aim," "forward," "expect," "plan," "intend," "believe," "potential," "continue," and similar expressions, although not all forward-looking statements contain these identifying words. These forward looking statements include express or implied statements regarding the Company's future expectations, plans and prospects, and include, without limitation, statements regarding the timing and advancement of the Company's programs, including the clinical trials of VTP-200, VTP-300, and VTP-850, statements regarding the timing for the potential IND application for VTP-1000, statements regarding the presentation of interim data, including with respect to VTP-300, and statements regarding the Company's capital, including its cash runway. Any forward-looking statements in this press release are based on management's current expectations and beliefs and are subject to numerous risks, uncertainties and important factors that may cause actual events or results to differ materially from those expressed or implied by any forward-looking statements contained in this press release, including, without limitation, risks and uncertainties related to: the success, cost and timing of the Company's product development activities and planned and ongoing clinical trials, the Company's ability to execute on its strategy, regulatory developments, approval of the Company's product candidates, the Company's ability to fund its operations and access capital, and global economic uncertainty, including disruptions in the banking industry, the conflict in Ukraine, and the conflict in Israel and Gaza, and other risks identified in the Company's filings with the Securities and Exchange Commission (the SEC), including its Annual Report on Form 10-K for the year ended December 31, 2022, its Quarterly Reports on Form 10-Q and Current Reports on Form 8-K. The Company cautions you not to place undue reliance on any forward-looking statements, which speak only as of the date they are made. The Company expressly disclaims any obligation to publicly update or revise any such statements to reflect any change in expectations or in events, conditions or circumstances on which any such statements may be based, or that may affect the likelihood that actual results will differ from those set forth in the forward-looking statements.



#### BARINTHUS BIOTHERAPEUTICS PLC CONDENSED CONSOLIDATED BALANCE SHEETS (IN THOUSANDS, EXCEPT NUMBER OF SHARES AND PER SHARE AMOUNTS) (UNAUDITED)

		September 30, 2023		December 31, 2022	
ASSETS					
Current assets:					
Cash and cash equivalents	\$	160,309	\$	194,385	
Accounts receivable				323	
Accounts receivable - related parties		—		5,524	
Research and development incentives receivable		4,172		4,541	
Prepaid expenses and other current assets		6,584		8,268	
Total current assets		171,065		213,041	
Goodwill		12,209		12,209	
Property and equipment, net		12,269		7,957	
Intangible assets, net		25,898		28,269	
Right of use assets, net		7,474		7,753	
Other assets		1,055		976	
Total assets	\$	229,970	\$	270,205	
LIABILITIES AND SHAREHOLDERS' EQUITY					
Current liabilities:					
Accounts payable	\$	5,145	\$	3,748	
Accrued expenses and other current liabilities		11,923		8,061	
Operating lease liability - current		1,501		433	
Total current liabilities		18,569		12,242	
Non-Current liabilities:		-,		,	
Operating lease liability		11,202		8,340	
Contingent consideration		1,797		1,711	
Deferred tax liability, net		1,521		3,746	
Other non-current liabilities		1,278		965	
Total liabilities	\$	34,367	\$	27,004	
Commitments and contingencies (Note 14)	<u> </u>	<u> </u>	<u>.</u>		
Shareholders' equity:					
Ordinary shares, £0.000025 nominal value; 38,546,594 shares authorized, issued and outstanding (December 31, 2022:					
authorized, issued and outstanding: 37,683,531)		1		1	
Deferred A shares, £1 nominal value; 63,443 shares authorized, issued and outstanding (December 31, 2022: authorized,					
issued and outstanding: 63,443)		86		86	
Deferred B shares, £0.01 nominal value; nil shares authorized, issued and outstanding (December 31, 2022:authorized,					
issued and outstanding: 570,987)				8	
Deferred C shares, £0.000007 nominal value, nil shares authorized, issued and outstanding (December 31, 2022: authorized, issued and outstanding: 27,828,231)		_		0	
Additional paid-in capital		385,707		379,504	
Accumulated deficit		(159,297)		(103,243)	
Accumulated other comprehensive loss – foreign currency translation adjustments		(31,099)		(33,460)	
Total shareholders' equity attributable to Barinthus Biotherapeutics plc shareholders		195,398		242,896	
Noncontrolling interest		205		305	
Total shareholders' equity	\$	195,603	\$	243,201	
Total liabilities and shareholders' equity	\$	229,970	\$	270,205	

<sup>1</sup> indicates amount less than thousand.



#### BARINTHUS BIOTHERAPEUTICS PLC CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS AND COMPREHENSIVE LOSS (IN THOUSANDS, EXCEPT NUMBER OF SHARES AND PER SHARE AMOUNTS) (UNAUDITED)

	Three months ended		Nine months ended		
	September 30, 2023	September 30, 2022	September 30, 2023	September 30, 2022	
License revenue <sup>(1)</sup>	\$	\$ 6,165	\$ 802	\$ 38,237	
Research grants and contracts			_	9	
Total revenue		6,165	802	38,246	
Operating expenses	·			· · · · · · · · · · · · · · · · · · ·	
Research and development	15,144	9,744	38,501	30,165	
General and administrative	961	(10,815)	26,227	(12,971)	
Total operating expenses	16,105	(1,071)	64,728	17,194	
(Loss)/income from operations	(16,105)	7,236	(63,926)	21,052	
Other income (expense):					
Interest income	196	1,024	2,306	1,776	
Interest expense	(7)	11	(21)	3	
Research and development incentives	1,205	(724)	2,921	1,150	
Other income	(2)	—	308	51	
Total other income (expense)	1,392	311	5,514	2,980	
(Loss)/profit before income tax	(14,713)	7,547	(58,412)	24,032	
Tax benefit	603	674	2,255	2,452	
Net (loss)/income	(14110)	8,221	(56,157)	26,484	
Net loss attributable to noncontrolling interest	38	21	103	47	
Net (loss)/income attributable to Barinthus Biotherapeutics					
plc shareholders	(14,072)	8,242	(56,054)	26,531	
Weighted-average ordinary shares outstanding, basic	38,533,833	37,247,123	38,320,208	37,213,787	
Weighted-average ordinary shares outstanding, diluted	38,533,833	38,156,564	38,320,208	38,226,092	
Net (loss)/income per share attributable to ordinary		<u></u> _	· · · · · · · · · · · · · · · · · · ·	· <u>····</u> ·	
shareholders, basic	\$ (0.37)	\$ 0.22	\$ (1.46)	\$ 0.71	
Net (loss)/income per share attributable to ordinary	<u>`</u>		· · · · · · · · · · · · · · · · · · ·		
shareholders, diluted	\$ (0.37)	\$ 0.22	\$ (1.46)	\$ 0.69	
		- <u></u>			
Net (loss)/income	\$ (14,110)	\$ 8,221	\$ (56,157)	\$ 26,484	
Other comprehensive gain/(loss) – foreign currency	4 (- ·,)	• •,	÷ (co,_co)	<b>4</b> _ 0,10 1	
translation adjustments	(7,820)	(19,940)	2,364	(42,730)	
Comprehensive loss	(21,930)	(11,719)	(53,793)	(16,246)	
Comprehensive loss attributable to noncontrolling interest	48	51	100	122	
Comprehensive loss attributable to Barinthus					
Biotherapeutics plc shareholders	\$ (21,882)	\$ (11,668)	\$ (53,693)	\$ (16,124)	
	+ (11,002)	- (11,000)	+ (88,000)	- (10,121)	

<sup>1</sup> Includes license revenue from related parties for the three and nine month periods ended September 30, 2023 of \$Nil million and \$0.8 million, respectively and for the three and nine month periods ended September 30, 2022 of \$6.2 million and \$38.2 million, respectively.



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