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 13, 2025

Xenetic Biosciences, Inc.

(Exact name of registrant as specified in charter)

Nevada (State or other jurisdiction of incorporation) **001-37937** (Commission File Number)

45-2952962 (IRS Employer Identification No.)

945 Concord Street
Framingham, Massachusetts
(Address of principal executive offices)

01701 (Zip Code)

(781) 778-7720

(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 is intended to sin Instruction A.2. below):	multaneously satisfy the filing obligation o	of the registrant under any of the following provisions (see General
$\hfill \Box$ Written communications pursuant to Rule 425 under the Security	rities Act (17 CFR 230.425)	
\square Soliciting material pursuant to Rule 14a-12 under the Exchange	ge Act (17 CFR 240.14a-12)	
☐ Pre-commencement communications pursuant to Rule 14d-2(b	b) under the Exchange Act (17 CFR 240.14	d-2(b))
☐ Pre-commencement communications pursuant to Rule 13e-4(c	c) under the Exchange Act (17 CFR 240.13	e-4(c))
Securiti	ies registered pursuant to Section 12(b) of t	the Act:
Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, \$0.001 par value per share	XBIO	The Nasdaq Stock Market
Indicate by check mark whether the registrant is an emerging grov Securities Exchange Act of 1934 (17 CFR §240.12b-2).	wth company as defined in Rule 405 of th	e Securities Act of 1933 (17 CFR §230.405) or Rule 12b-2 of the
		Emerging growth company \square
If an emerging growth company, indicate by check mark if the reg accounting standards provided pursuant to Section 13(a) of the Exc		transition period for complying with any new or revised financial

Item 2.02. Results of Operations and Financial Condition.

On August 13, 2025, Xenetic Biosciences, Inc. (the "Company") issued a press release announcing results for the three months ended June 30, 2025.

The full text of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K and hereby incorporated in this Item 2.02 by reference. The information in this Item 2.02, including Exhibit 99.1 attached hereto, shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, except as shall be expressly set forth by specific reference in such filing.

FORWARD-LOOKING STATEMENTS

This Form 8-K, including the press release, contains forward-looking statements that we intend to be subject to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. All statements contained in this Form 8-K, including the press release, other than statements of historical facts may constitute forward-looking statements within the meaning of the federal securities laws. These statements can be identified by words such as "expects," "plans," "projects," "will," "may," "anticipates," "believes," "should," "intends," "estimates," and other words of similar meaning. Any forward-looking statements contained herein are based on current expectations, and are subject to a number of risks and uncertainties. Many factors could cause our actual activities, performance, achievements or results to differ materially from the activities and results anticipated in forward-looking statements. These risks and uncertainties include those described in the "Risk Factors" section as detailed from time to time in the Company's reports filed with the Securities and Exchange Commission ("SEC"), including the Company's annual report on Form 10-K, periodic quarterly reports on Form 10-Q, current reports on Form 8-K and other documents filed with the SEC. In addition, forward-looking statements may also be adversely affected by general market factors, general economic and business conditions, including potential adverse effects of public health issues and geopolitical events, such as the conflicts in the Ukraine and in the Middle East, on economic activity, competitive product development, product availability, federal and state regulations and legislation, the regulatory process for new product candidates and indications, manufacturing issues that may arise, patent positions, litigation, and shareholder activism, among other factors. The forward-looking statements contained in this Form 8-K, including the press release, speak only as of the date the statements were made, and the Company does not undertake an

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
99.1	Press Release dated August 13, 2025 pertaining to the financial results of the Company for the three months ended June 30, 2025.
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.

Date: August 13, 2025

XENETIC BIOSCIENCES, INC.

By: /s/ James Parslow
Name: James Parslow

Title: Interim Chief Executive Officer & Chief Financial Officer



Xenetic Biosciences, Inc. Reports Second Quarter 2025 Financial Results

Expanded strategic partnership with The Scripps Research Institute to advance proof-of-concept studies and further develop its program combining systemic DNase I with CAR T-cell therapies

Ended the quarter with \$4.8 million of cash to fund operations

FRAMINGHAM, MA – (August 13, 2025) – Xenetic Biosciences, Inc. (NASDAQ: XBIO) ("Xenetic" or the "Company"), a biopharmaceutical company focused on advancing innovative immuno-oncology technologies addressing difficult to treat cancers, today reported its financial results for the second quarter 2025.

Recent Highlights

- · Expanded its collaboration with The Scripps Research Institute ("TSRI") to advance the development of the Company's development program evaluating the combination of systemic DNase I and CAR T-cell therapies;
- · Announced advancements from its collaboration partner, PeriNess Ltd. ("PeriNess") including:
 - o Entered into a Clinical Study Agreement to support an exploratory clinical study of DNase I in combination with anti-CD19 CAR T cells in patients with large B cell lymphoma;
 - o Commenced patient dosing in an exploratory clinical study of systemic DNase I in combination with FOLFIRINOX for the first line treatment of unresectable, locally advanced or metastatic pancreatic cancer at Bnei Zion Medical Center; and
- · Continued pursuit of other strategic collaborations to advance the Company's technology.

"We continue to set a strong foundation that we believe positions us for success as we advance our systemic DNase I in combination with immunotherapy, chemotherapy, and radiotherapy across various oncology indications where there remains significant unmet need. We continue to work with our partners and believe the data and information will be invaluable as we look to realize the full potential of our DNase platform technology. Looking ahead, we remain focused on building momentum across all fronts and driving development toward an IND and Phase 1 clinical trial," commented James Parslow, Interim Chief Executive Officer and Chief Financial Officer of Xenetic.

Xenetic continues to advance its DNase-based technology towards Phase 1 clinical development for the treatment of pancreatic carcinoma and other locally advanced or metastatic solid tumors. Preclinical proof-of-concept studies combining DNase I with chemotherapy, immunotherapies, and CAR-T therapy in hematological and solid tumor and metastatic cancer models have been completed. Building on proof-of-concept success, the program has now advanced to mechanism-of-action and translational studies in preparation for a Phase 1 clinical trial.

Additionally, as previously announced in December 2024, Xenetic entered into a Clinical Trial Services Agreement with PeriNess, under which PeriNess will lead in the regulatory approval, operational execution and management of potential exploratory, investigator initiated studies of recombinant DNase as an adjunctive treatment in patients with pancreatic carcinoma and other locally advanced or metastatic solid tumors receiving chemotherapy and immunotherapy in Israeli medical centers.

Summary of Financial Results for Second Quarter 2025

Net loss for the quarter ended June 30, 2025 was approximately \$0.7 million. Research & development expenses for the three months ended June 30, 2025 decreased by approximately \$277,000, or 29.7%, to approximately \$0.7 million from \$0.9 million in the comparable quarter in 2024. General and administrative expenses for the three months ended June 30, 2025 decreased by approximately \$472,000, or 41.8%, to approximately \$0.7 million from approximately \$1.1 million in the comparable quarter in 2024. These decreases were primarily due to certain severance and benefits expensed in connection with separation agreements entered into during the second quarter of 2024 with the Company's former Chief Executive Officer and Chief Scientific Officer.

The Company ended the quarter with approximately \$4.8 million in cash.

About Xenetic Biosciences

Xenetic Biosciences, Inc. is a biopharmaceutical company focused on advancing innovative immune-oncology technologies addressing hard to treat cancers. The Company's DNase platform is designed to improve outcomes of existing treatments, including immunotherapies, by targeting neutrophil extracellular traps (NETs), which are involved in cancer progression. Xenetic is currently focused on advancing its systemic DNase program into the clinic as an adjunctive therapy for pancreatic carcinoma and locally advanced or metastatic solid tumors.

For more information, please visit the Company's website at www.xeneticbio.com and connect on X, LinkedIn, and Facebook.

Forward-Looking Statements

This press release contains forward-looking statements that we intend to be subject to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. All statements contained in this press release other than statements of historical facts may constitute forward-looking statements within the meaning of the federal securities laws. These statements can be identified by words such as "expects," "plans," "projects," "will," "may," "anticipates," "believes," "should," "intends," "estimates," "remain," "focus", "confidence in", "potential", and other words of similar meaning, including, but not limited to, all statements regarding expectations for our DNase-base oncology platform, including statements regarding: (a) advancing our DNase-based oncology program towards Phase 1 clinical development for the treatment of pancreatic carcinoma and other locally advanced or metastatic solid tumors and our focus on building momentum and driving development toward an IND and Phase I clinical trial, (b) setting a strong foundation that we believe positions us for success as we advance DNase I in combination with various types of therapy and (c) working with our partners and our belief regarding the data and information as we look to realize the full potential of DNase. Any forward-looking statements contained herein are based on current expectations and are subject to a number of risks and uncertainties. Many factors could cause our actual activities, performance, achievements, or results to differ materially from the activities and results anticipated in forward-looking statements. Important factors that could cause actual activities, performance, achievements, or results to differ materially from such plans, estimates or expectations include, among others, (1) unexpected costs, charges or expenses resulting from our manufacturing and collaboration agreements; (2) unexpected costs, charges or expenses resulting from the licensing of the DNase platform; (3) uncertainty of the expected financial performance of the Company following the licensing of the DNase platform; (4) failure to realize the anticipated potential of the DNase or PolyXen technologies; (5) the ability of the Company to obtain funding and implement its business strategy; and (6) other risk factors as detailed from time to time in the Company's reports filed with the SEC, including its annual report on Form 10-K, periodic quarterly reports on Form 10-Q, current reports on Form 8-K and other documents filed with the SEC. The foregoing list of important factors is not exclusive. In addition, forward-looking statements may also be adversely affected by general market factors, general economic and business conditions, including potential adverse effects of public health issues, and geopolitical events, such as the conflicts in Ukraine and in the Middle East, on economic activity, competitive product development, product availability, federal and state regulations and legislation, the regulatory process for new product candidates and indications, manufacturing issues that may arise, patent positions, litigation, and shareholder activism, among other factors. The forward-looking statements contained in this press release speak only as of the date the statements were made, and the Company does not undertake any obligation to update forward-looking statements, except as required by law.

Contact:

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