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**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION**

WASHINGTON, D.C. 20549

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**FORM 8-K**

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**CURRENT REPORT  
Pursuant to Section 13 or 15(d) of the  
Securities Exchange Act of 1934**

Date of report (Date of earliest event reported): **January 6, 2025**

**Fortress Biotech, Inc.**

(Exact Name of Registrant as Specified in Charter)

**Delaware**  
(State or Other Jurisdiction  
of Incorporation)

**001-35366**  
(Commission File Number)

**20-5157386**  
(IRS Employer  
Identification No.)

**1111 Kane Concourse, Suite 301  
Bay Harbor Islands, FL 33154**  
(Address of Principal Executive Offices)

**(781) 652-4500**  
(Registrant's telephone number, including area code)

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.
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act.
- Pre-commencement communications pursuant to Rule 14d-2b under the Exchange Act.
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act.

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock	FBIO	Nasdaq Capital Market
9.375% Series A Cumulative Redeemable Perpetual Preferred Stock	FBIO P	Nasdaq Capital 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.

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**Item 8.01. Other Events.**

On January 6, 2025, Cyprium Therapeutics, Inc. (“**Cyprium**”), a majority-owned subsidiary of Fortress Biotech, Inc. (the “**Company**” or “**Fortress**”), announced that the U.S. Food and Drug Administration (the “**FDA**”) accepted the New Drug Application (“**NDA**”) for CUTX-101 (Copper Histidinate) for priority review with a target action date of June 30, 2025.

In December 2023, Sentyln Therapeutics, Inc. (“**Sentyln**”), a U.S.-based biopharmaceutical company wholly-owned by Zydus Lifesciences, Ltd. (“**Zydus Group**”), assumed full responsibility for the development and commercialization of CUTX-101 from Cyprium. The NDA submission was completed by Sentyln who will be responsible for commercialization upon approval.

Cyprium will retain ownership over any Priority Review Voucher that may be issued at NDA approval and is eligible to receive royalties and up to \$129 million in aggregate development and sales milestones.

Fortress founded Cyprium in 2017 and currently owns approximately 76% of Cyprium.

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**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.

**Fortress Biotech, Inc.**  
(Registrant)

Date: January 6, 2025

By: /s/ David Jin  
David Jin  
Chief Financial Officer

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