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): January 10, 2025

XERIS BIOPHARMA HOLDINGS, INC.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation)

001-40880 (Commission File Number) **87-1082097** (I.R.S. Employer Identification No.)

1375 West Fulton Street, Suite 1300 Chicago, Illinois 60607

(Address of principal executive offices, including zip code)

(844) 445-5704

(Registrant's telephone number, including area code)

	(Former name o	Not Applicable or former address, if changed since last	report)	
Check	the appropriate box below if the Form 8-K filing is intended to s	simultaneously satisfy the filing obliga	tion 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))			
Securit	ies 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, par value \$0.0001 per share	XERS	The Nasdaq Global Select Market	
Indicat 12b-2 o	e by check mark whether the registrant is an emerging growth of the Securities Exchange Act of 1934 (§ 240.12b-2 of this chap	company as defined in Rule 405 of the pter).	he Securities Act of 1933 (§ 230.405 of this chapter) or Rule	
Emergir	ng growth company □			
	nerging growth company, indicate by check mark if the regist l accounting standards provided pursuant to Section 13(a) of the		ded transition period for complying with any new or revised	

Item 2.02 Results of Operations and Financial Condition.

On January 10, 2025, Xeris Biopharma Holdings, Inc. (the "Company") issued a press release announcing certain of its financial preliminary results for the fourth quarter and full year ended December 31, 2024. A copy of this press release is attached as Exhibit 99.1 to this Current Report on Form 8-K.

The information disclosed pursuant to Item 2.02 in this Current Report on Form 8-K (including Exhibit 99.1) is being furnished and 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 Number	Description
99.1	Press release issued by the Company dated January 10, 2025
104	Cover Page Interactive Data File (embedded within the Inline XBRI, 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: January 10, 2025 Xeris Biopharma Holdings, Inc.

By: /s/ Steven M. Pieper

Name: Steven M. Pieper Title: Chief Financial Officer



XERIS EXPECTS TO EXCEED FULL-YEAR 2024 FINANCIAL GUIDANCE

Full-year 2024 total revenue projected to be \$203 million, exceeding previous guidance of \$198-\$202 million

Year-end 2024 cash position expected to be over \$71 million, generating positive cash flow in the fourth quarter

Recorlev® net revenue Q4 2024 anticipated to increase by approximately \$5 million or 28% sequentially

2024 financial results and 2025 outlook expected on March 6, 2025

CHICAGO, IL; **January 10, 2025** – Xeris Biopharma Holdings, Inc. (Nasdaq: XERS) today announced it expects to generate total 2024 revenue of \$203 million, which will exceed previously announced 2024 total revenue guidance of \$198 million to \$202 million. The Company also anticipates ending 2024 with over \$71 million in cash, cash equivalents, and short-term investments, generating positive cash flow in the fourth quarter.

"We achieved another record quarter of exceptional total revenue growth with revenues expected to be \$60 million for the fourth quarter and \$203 million for the full year representing growth of 35% and 24%, respectively. These impressive results are driven by accelerating demand for Recorlev and continued strong Gvoke demand," said John Shannon, CEO of Xeris. "Our focus remains on continuing to drive exceptional product revenue growth and advancing our robust pipeline - namely our Phase 3 ready, XP-8121 program. With our commercial engine leading the way and our strong balance sheet, we are well positioned for a transformational 2025."

Other Fourth Quarter 2024 Updates

- Recorlev® achieved record number of new starts and referrals.
- **Gyoke**® ended the year with approximately 35% market share.
- **Kevevis®** maintained a similar number of patients on therapy as Q3 2024.
- Beta Bionics, Inc.: Xeris successfully formulated a unique XeriSol® formulation of glucagon for bi-hormonal pumps and pump systems. This triggered a milestone payment of \$3 million in the fourth quarter.
- Amgen: Amgen chose to terminate its exclusive worldwide license agreement with Xeris following a broader company portfolio assessment. Their decision was not related to the performance of the XeriJect® formulation technology. The termination will not have a material impact on Xeris' outlook.

About Xeris

Xeris (Nasdaq: XERS) is a growth-oriented biopharmaceutical company committed to improving patient lives by developing and commercializing innovative products across a range of therapies. Xeris has three commercially available products: Recorlev®, for the treatment of endogenous Cushing's syndrome; Gvoke®, a ready-to-use liquid glucagon for the treatment of severe hypoglycemia; and Keveyis®, a proven therapy for primary periodic paralysis. Xeris also has a pipeline of development programs led by XP-8121, a Phase 3-ready, once-weekly subcutaneous injection for hypothyroidism, as well as multiple early-stage programs leveraging Xeris' technology platforms, XeriSol® and XeriJect®, for its partners.

Xeris Biopharma Holdings is headquartered in Chicago, IL. For more information, visit <u>www.xerispharma.com</u>, or follow us on \underline{X} , <u>LinkedIn</u>, or <u>Instagram</u>.

Forward-Looking Statements

Any statements in this press release other than statements of historical fact are forward-looking statements. Forward-looking statements include, but are not limited to, statements about future expectations, plans and prospects for Xeris Biopharma Holdings, Inc. including statements regarding the financial outlook for 2024, including quarterly product revenue, projections regarding year-end 2024 cash estimates and total revenue, company performance in 2025, the potential for growth of revenue, the market, demand and therapeutic potential of its products and product candidates, the potential utility of its formulation platforms, the advancement of its pipeline (including XP-8121), the impact of the termination of the Amgen license agreement on our outlook, the timing of the release of our financial results and outlook, and other statements containing the words "will," "would," "continue," "expect," "should," "anticipate" and similar expressions, constitute forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995. These forward-looking statements are based on numerous assumptions and assessments made in light of Xeris' experience and perception of historical trends, current conditions, business strategies, operating environment, future developments, geopolitical factors and other factors it believes appropriate. By their nature, forward-looking statements involve known and unknown risks and uncertainties because they relate to events and depend on circumstances that will occur in the future. The various factors that could cause Xeris' actual results, performance or achievements, industry results and developments to differ materially from those expressed in or implied by such forward-looking statements, include, but are not limited to, its financial position and need for financing, including to fund its product development programs or commercialization efforts, whether its products will achieve and maintain market acceptance in a competitive business environment, its reliance on third-party suppliers, including single-source suppliers, its reliance on third parties to conduct clinical trials, the ability of its product candidates to compete successfully with existing and new drugs, and its and collaborators' ability to protect its intellectual property and proprietary technology. No assurance can be given that such expectations will be realized and persons reading this communication are, therefore, cautioned not to place undue reliance on these forward-looking statements. Additional risks and information about potential impacts of financial, operational, economic, competitive, regulatory, governmental, technological, and other factors that may affect Xeris can be found in Xeris' filings, including those other risk factors identified in the "Risk Factors" and "Management's Discussion and Analysis of Financial Condition and Results of Operations" sections of Xeris' Annual Report on Form 10-K for the year ended December 31, 2023, and subsequent Quarterly Reports on Form 10-Q, as filed with the Securities and Exchange Commission (SEC), the contents of which are not incorporated by reference into, nor do they form part of, this communication. You are encouraged to read our filings with the SEC, available at www.sec.gov and www.xerispharma.com, for a discussion of these and other risks and uncertainties. Forward-looking statements in this communication are based on information available to us, as of the date of this communication and, while we believe our assumptions are reasonable, actual results may differ materially. Subject to any obligations under applicable law, we do not undertake any obligation to update any forward-looking statement whether as a result of new information, future developments or otherwise, or to conform any forward-looking statement to actual results, future events, or to changes in expectations. Our business is subject to substantial risks and uncertainties, including those referenced above. Investors, potential investors, and others should give careful consideration to these risks and uncertainties.

Investor Contact

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