

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, DC 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): May 20, 2022

KIORA PHARMACEUTICALS, INC.

(Exact name of registrant as specified in its charter)

Delaware

(State or other jurisdiction of incorporation)

001-36672

(Commission File Number)

98-0443284

(IRS Employer Identification No.)

1371 East 2100 South
Suite 200

Salt Lake City, Utah 84105
(Address of principal executive offices)

84105
(Zip Code)

(781) 788-9043

(Registrant's telephone number, including area code)

(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:

Title of each class:	Trading Symbol(s)	Name of each exchange on which registered:
Common Stock, \$0.01 par value	KPRX	The 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.

Item 4.02 Non-Reliance on Previously Issued Financial Statements or a Related Audit Report or Completed Interim Review.

On May 20, 2022, the Audit Committee of the Board of Directors (the "Audit Committee") of Kiora Pharmaceuticals, Inc. (the "Company"), after discussion with management of the Company and the Company's independent registered public accounting firm, EisnerAmper LLP ("EisnerAmper"), concluded that the Company's previously issued audited consolidated financial statements as of and for the fiscal years ended December 31, 2021 and December 31, 2020 included in the Company's Annual Reports on Form 10-K for such periods and unaudited condensed consolidated interim financial statements as of and for the fiscal periods ended March 31, 2021, June 30, 2021 and September 30, 2021 included in the Company's Quarterly Reports on Form 10-Q for such periods should no longer be relied upon. Similarly, earnings releases, and investor communications describing the financial statements for the periods described above should no longer be relied upon.

As part of preparing its condensed consolidated financial statements as of and for the quarter ended March 31, 2022, the Company identified inadvertent errors in the accounting for certain contingent consideration.

In connection with the Company's acquisition of Panoptes Pharma Ges.m.b.H in December 2020, shares of the Company's common stock that were held back at closing and that will be issued on the 18-month anniversary of the acquisition, subject to deduction for any indemnification claims, post-closing adjustments and other specified matters (the "Holdback Shares"), were recorded as a liability of \$1.353 million. Upon further evaluation, the Company determined that the Holdback Shares should have been accounted for as equity.

Additionally, in connection with the Company's acquisition of Jade Therapeutics, Inc. ("Jade") in March 2016, the Company will be required to pay the former shareholders of Jade a cash earnout payment in the event any product developed by Jade prior to its acquisition, or derivative of such product, receives FDA approval. However, due to the fact that the Company's KIO-201 product candidate is now being developed as a drug rather than a device, the expected development time for KIO-201 has been extended. As a result of that extended timeline, the Company anticipates a reduction of the contingent consideration for KIO-201 as of December 31, 2021.

