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 12, 2026

BIOAGE LABS, INC.

(Exact name of Registrant as Specified in Its Charter)

Delaware	**001-42279**	**47-4721157**
(State or Other Jurisdiction of Incorporation)	**(Commission File Number)**	**(IRS Employer Identification No.)**
5885 Hollis Street Suite 370 Emeryville, California		**94608**
(Address of Principal Executive Offices)		**(Zip Code)**

Registrant's Telephone Number, Including Area Code: 510 806-1445

(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, $0.00001 Par Value Per Share	BIOA	Nasdaq Global Select 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 7.01 Regulation FD Disclosure.

On January 12, 2026, BioAge Labs, Inc. (the "Company") issued a press release announcing additional positive interim Phase 1 data for BGE-102, a novel brain-penetrant NLRP3 inhibitor, demonstrating Potential for Best-in-class hsCRP reduction in participants with elevated cardiovascular risk.

A copy of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K.

The information furnished in this Item 7.01, including Exhibit 99.1, shall not be deemed to be "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 or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended (the "Securities Act"). The information contained in this Item 7.01 and in the accompanying Exhibit 99.1 shall not be incorporated by reference into any other filing under the Exchange Act or under the Securities Act, except as shall be expressly set forth by specific reference in such filing.

Item 8.01 Other Events

Phase 1 Data for BGE-102

On January 12, 2026, the Company announced additional positive interim Phase 1 data for BGE-102, a novel brain-penetrant NLRP3 inhibitor, demonstrating potential for best-in-class hsCRP reduction in participants with elevated cardiovascular risk.

- First BGE-102 multiple ascending dose ("MAD") cohort completed in obese individuals with elevated hsCRP receiving 120 mg once daily ("QD"); demonstrated rapid and profound reduction in inflammatory markers
- BGE-102 achieved 86% reduction in hsCRP at Day 14, with 93% of participants reaching normalized levels (<2 mg/L)
- BGE-102 demonstrated significant reductions in IL-6, a key driver of systemic inflammation and cardiovascular risk, and fibrinogen, an independent predictor of cardiovascular events
- BGE-102 was well tolerated with a favorable safety profile
- Patent issued covering additional composition of matter and novel NLRP3 binding site

Anticipated Milestones

- **1H 2026:** Completion of Phase 1 trial with full data readout, including two additional MAD cohorts in obese participants with elevated hsCRP
- **1H 2026:** Initiation of Phase 2a proof-of-concept study in patients with obesity and cardiovascular risk factors. The trial is planned to enroll approximately 100 patients randomized 1:1 to BGE-102 monotherapy or placebo for 12 weeks. The anticipated primary endpoint is percent change in hsCRP. The trial will also assess inflammatory and metabolic biomarkers, and will include liver MRI
- **2H 2026:** Phase 2a data readout

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
99.1	Press release issued by BioAge Labs, Inc. dated January 12, 2026.
104	Cover Page Interactive Data File (embedded within the Inline XBRL document).

Cautionary Note Regarding Forward-Looking Statements

This Current Report on Form 8-K contains "forward-looking statements" within the meaning of, and made pursuant to the safe harbor provisions of, the Private Securities Litigation Reform Act of 1995. All statements contained in this press release that do not relate to matters of historical fact should be considered forward-looking statements, including, but not limited to, statements regarding the Company's plans to develop and commercialize its product candidates, including BGE-102, the potential for BGE-102 as a treatment for cardiovascular risk and the expected timeline for future data readouts from our ongoing Phase 1 clinical trial. These forward-looking statements may be accompanied by such words as "aim," "anticipate," "believe," "could," "estimate," "expect," "forecast," "goal," "intend," "may," "might," "plan," "potential," "possible," "will," "would," and other words and terms of similar meaning. These statements involve risks and uncertainties that could cause actual results to differ materially from those reflected in such statements, including: the Company's ability to develop, obtain regulatory approval for and commercialize its product candidates; the timing and results of preclinical studies and clinical trials; the risk that positive results in a clinical trial may not be replicated in subsequent trials or success in early stage clinical trials may not be predictive of results in later stage clinical trials; risks associated with clinical trials, including its ability to adequately manage clinical activities, unexpected concerns that may arise from additional data or analysis obtained during clinical trials, regulatory authorities may require additional information or further studies, or may fail to approve or may delay approval of its drug candidates; the occurrence of adverse safety events; failure to protect and enforce its intellectual property, and other proprietary rights; failure to successfully execute or realize the anticipated benefits of its strategic and growth initiatives; risks relating to technology failures or breaches; its dependence on collaborators and other third parties for the development of product candidates and other aspects of its business, which are outside of the Company's full control; risks associated with current and potential delays, work stoppages, or supply chain disruptions, including due to the imposition of tariffs and other trade barriers; risks associated with current and potential future healthcare reforms; risks relating to attracting and retaining key personnel; changes in or failure to comply with legal and regulatory requirements, including shifting priorities within the U.S. Food and Drug Administration; risks relating to access to capital and credit markets; and the other risks and uncertainties that are detailed under the heading "Risk Factors" included in the Company's Quarterly Report on Form 10-Q filed with the U.S. Securities and Exchange Commission (SEC) on November 6, 2025, and Company's other filings with the SEC filed from time to time. The Company undertakes no obligation to publicly update any forward-looking statement, whether written or oral, that may be made from time to time, whether as a result of new information, future developments or otherwise.

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.

BIOAGE LABS, INC.

Date: January 12, 2026

By: /s/ Dov Goldstein
Dov Goldstein, M.D.
Chief Financial Officer

Exhibit 99.1

BioAge Announces Additional Positive Interim Phase 1 Data for BGE-102, a Novel Brain-Penetrant NLRP3 Inhibitor, Demonstrating Potential for Best-in-Class hsCRP Reduction in Participants with Elevated Cardiovascular Risk

First BGE-102 MAD cohort completed in obese individuals with elevated hsCRP receiving 120 mg QD; demonstrated rapid and profound reduction in inflammatory markers

BGE-102 achieved 86% reduction in hsCRP at Day 14, with 93% of participants reaching normalized levels (<2 mg/L)

BGE-102 demonstrated significant reductions in IL-6, a key driver of systemic inflammation and cardiovascular risk, and fibrinogen, an independent predictor of cardiovascular events

BGE-102 was well tolerated with a favorable safety profile

Patent issued covering additional composition of matter and novel NLRP3 binding site

Full Phase 1 data, including additional MAD cohorts in obese participants with elevated hsCRP, anticipated 1H 2026; Phase 2a study on track to initiate in 1H 2026

EMERYVILLE, Calif.--(BUSINESS WIRE)--BioAge Labs, Inc. (Nasdaq: BIOA) ("BioAge"), a clinical-stage biopharmaceutical company developing therapeutic product candidates for metabolic diseases by targeting the biology of human aging, today announced additional positive interim data from the ongoing Phase 1 clinical trial evaluating BGE-102, a potent, structurally novel, orally available, brain-penetrant small molecule NLRP3 inhibitor being developed for the treatment of patients with cardiovascular risk factors.

In a multiple ascending dose (MAD) cohort of participants with obesity (BMI 32–42) and elevated baseline inflammation (hsCRP >3 mg/L), BGE-102 120 mg once daily achieved an 86% median reduction in high-sensitivity C-reactive protein (hsCRP) at Day 14. Notably, 93% of BGE-102-dosed participants (13 of 14) achieved hsCRP levels below 2 mg/L—the clinically recognized threshold for reduced cardiovascular risk.

These findings build on positive interim data announced in December 2025 from SAD and initial MAD cohorts, which demonstrated that BGE-102 was well tolerated, achieved dose-proportional pharmacokinetics supporting once-daily dosing, and produced 90-98% suppression of IL-1β at Day 14 trough. Those data also confirmed high brain penetration, with cerebrospinal fluid (CSF) concentrations exceeding the IC90 at doses of 60 mg and above.

“We are very encouraged by these results, which support the potential for BGE-102 to deliver injectable-like inflammation reduction in an oral therapy designed for primary care, the clinical setting where most cardiovascular risk is managed and where oral medicines are preferred by patients and physicians,” said Kristen Fortney, PhD, CEO and co-founder of BioAge. “Chronic inflammation is now recognized as a major driver of cardiovascular disease—on par with cholesterol—yet it remains far less commonly treated. An 86% reduction in hsCRP, with 93% of participants reaching levels associated with reduced cardiovascular risk, positions BGE-102 as a potential best-in-class oral therapy to directly address inflammation. These findings support our plans to advance BGE-102 into a Phase 2a study in the first half of this year.”

Key findings from the MAD cohort in patients with obesity and elevated hsCRP

Rapid and profound hsCRP reduction

- BGE-102 achieved 83% median reduction in hsCRP (from a median baseline of 4.85 mg/L) at Day 7 and 86% at Day 14
- 93% of participants (13/14) on BGE-102 achieved hsCRP <2 mg/L at Day 14; 71% (10/14) reached ≤1 mg/L
- Rapid onset of effect: 86% of BGE-102-dosed participants (12/14) achieved hsCRP levels <2 mg/L at Day 7; 71% (10/14) reached ≤1 mg/L
- hsCRP is the most widely used marker of inflammatory cardiovascular risk; levels below 2 mg/L are associated with reduced risk of cardiovascular events

Significant IL-6 reduction

- BGE-102 achieved a 44% median reduction in serum IL-6 at Day 14
- CSF IL-6 decreased in the two participants with elevated baseline levels, consistent with BGE-102's high brain penetration
- IL-6 is a key upstream driver of hsCRP production and a validated marker of cardiovascular risk

Significant fibrinogen reduction

- BGE-102 achieved a 30% reduction in fibrinogen at Day 14

- Elevated fibrinogen has been shown to be an independent predictor of cardiovascular events and thrombotic risk

Potent IL-1β suppression, consistent with strong target engagement

- In the ex vivo whole blood stimulation assay, BGE-102 achieved 93% suppression of IL-1β at trough (Day 14, pre-dose)
- IL-1β is directly downstream of NLRP3 and drives production of IL-6 and CRP, key markers of cardiovascular risk

Safety and tolerability

- BGE-102 continued to be well tolerated
- Adverse events were infrequent, mild to moderate in severity, and self-limited, with no dose-dependent pattern observed
- No dose-limiting toxicities observed

Additional figures and data from the ongoing Phase 1 study are available in the Company's corporate presentation at **https://ir.bioagelabs.com/**.

"The substantial reductions in hsCRP, IL-6, and fibrinogen we observed in participants with obesity and elevated inflammation demonstrate that BGE-102 potently suppresses the NLRP3-driven inflammatory cascade in a clinically relevant population," said Paul Rubin, MD, Chief Medical Officer of BioAge. "These data provide strong rationale for advancing into our planned Phase 2a study, where we will evaluate BGE-102's effects on a range of key inflammatory biomarkers over a longer duration in patients with elevated cardiovascular risk."

Phase 1 study design

The ongoing Phase 1 study is a randomized, double-blind, placebo-controlled trial in healthy volunteers and participants with obesity. Part 1 evaluated single ascending doses at four dose levels (10, 30, 60, and 120 mg); Part 2 to date has evaluated multiple ascending doses administered once daily for 14 days in healthy volunteers (60 and 120 mg) and in participants with obesity and elevated hsCRP (120 mg QD cohort complete; two lower-dose QD cohorts ongoing). Pharmacodynamic effects were evaluated by assessment of serum biomarkers including hsCRP, IL-6, and fibrinogen, as well as an ex vivo whole blood stimulation assay measuring IL-1β suppression.

Anticipated milestones for BGE-102 in cardiovascular disease

- **1H 2026:** Completion of Phase 1 trial with full data readout, including two additional MAD cohorts in obese participants with elevated hsCRP
- **1H 2026:** Initiation of Phase 2a proof-of-concept study in patients with obesity and cardiovascular risk factors. The trial is planned to enroll approximately 100 patients randomized 1:1 to BGE-102 monotherapy or placebo for 12 weeks. The anticipated primary endpoint is percent change in hsCRP. The trial will also assess inflammatory and metabolic biomarkers, and will include liver MRI.
- **2H 2026:** Phase 2a data readout

Background on BGE-102 and NLRP3

BGE-102 is a potent, orally available, brain-penetrant small molecule NLRP3 inhibitor being developed for diseases of inflammation including elevated cardiovascular risk. BGE-102 represents a structurally novel class of NLRP3 inhibitors developed by BioAge with a unique mechanism and binding site. NLRP3 is a key driver of age-related inflammation that has been implicated in a broad range of diseases, including cardiovascular disease, neurodegeneration, and metabolic disorders.

About BioAge Labs, Inc.

BioAge is a clinical-stage biopharmaceutical company developing therapeutic product candidates for metabolic diseases by targeting the biology of human aging. The Company's lead product candidate, BGE-102, is a potent, orally available, brain-penetrant small-molecule NLRP3 inhibitor being developed for cardiovascular risk factors. A Phase 1 SAD/MAD trial of BGE-102 is underway, with topline data including additional MAD cohorts anticipated in 1H26. The Company is also developing long-acting injectable and oral small molecule APJ agonists for obesity. BioAge’s additional preclinical programs, which leverage insights from the Company’s proprietary discovery platform built on human longevity data, address key pathways involved in metabolic aging.

Forward-looking statements

This press release contains "forward-looking statements" within the meaning of, and made pursuant to the safe harbor provisions of, the Private Securities Litigation Reform Act of 1995. All statements contained in this press release that do not relate to matters of historical fact should be considered forward-looking statements, including, but not limited to, statements regarding our plans to develop and commercialize our product candidates,

including BGE-102 and our APJ program, the potential for BGE-102 as a treatment for cardiovascular risk and the expected timeline for future data readouts from our ongoing Phase 1 clinical trial, the timing and results of our clinical trials, risks associated with clinical trials, including our ability to adequately manage clinical activities, the timing of and our ability to obtain and maintain regulatory approvals and the clinical utility of our product candidates. These forward-looking statements may be accompanied by such words as "aim," "anticipate," "believe," "could," "estimate," "expect," "forecast," "goal," "intend," "may," "might," "plan," "potential," "possible," "will," "would," and other words and terms of similar meaning. These statements involve risks and uncertainties that could cause actual results to differ materially from those reflected in such statements, including: our ability to develop, obtain regulatory approval for and commercialize our product candidates; the timing and results of preclinical studies and clinical trials; the risk that positive interim results in a preclinical study or clinical trial may not be replicated in subsequent trials or success in early stage clinical trials may not be predictive of results in later stage clinical trials; risks associated with clinical trials, including our ability to adequately manage clinical activities, unexpected concerns that may arise from additional data or analysis obtained during clinical trials, regulatory authorities may require additional information or further studies, or may fail to approve or may delay approval of our drug candidates; the occurrence of adverse safety events; failure to protect and enforce our intellectual property, and other proprietary rights; failure to successfully execute or realize the anticipated benefits of our strategic and growth initiatives; risks relating to technology failures or breaches; our dependence on collaborators and other third parties for the development of product candidates and other aspects of our business, which are outside of our full control; risks associated with current and potential delays, work stoppages, or supply chain disruptions, including due to the imposition of tariffs and other trade barriers; risks associated with current and potential future healthcare reforms; risks relating to attracting and retaining key personnel; changes in or failure to comply with legal and regulatory requirements, including shifting priorities within the U.S. Food and Drug Administration; risks relating to access to capital and credit markets; and the other risks and uncertainties that are detailed under the heading "Risk Factors" included in BioAge's Quarterly Report on Form 10-Q filed with the U.S. Securities and Exchange Commission (SEC) on November 6, 2025, and BioAge's other filings with the SEC filed from time to time. BioAge undertakes no obligation to publicly update any forward-looking statement, whether written or oral, that may be made from time to time, whether as a result of new information, future developments or otherwise.

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