
**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): September 24, 2026

CULLINAN THERAPEUTICS, INC.

(Exact name of Registrant as Specified in Its Charter)

Delaware
(State or Other Jurisdiction
of Incorporation)

001-39856
(Commission File Number)

81-3879991
(IRS Employer
Identification No.)

**One Main Street
Suite 1350
Cambridge, Massachusetts**
(Address of Principal Executive Offices)

02142
(Zip Code)

Registrant's Telephone Number, Including Area Code: 617 410-4650

(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, \$0.0001 par value per share	CGEM	The 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 September 24, 2026, Cullinan Therapeutics, Inc. (the "Company") issued a press release outlining anticipated milestones for the fourth quarter of 2026. A copy of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated herein by reference.

The information in this Current Report on Form 8-K furnished pursuant to Item 7.01, including Exhibit 99.1 attached hereto, shall not be deemed to be "filed" for the purposes of Section 18 of the Exchange Act, or otherwise subject to the liabilities of that section and will not be incorporated by reference into any filing by the Company under the Securities Act or the Exchange Act, unless specifically identified as being incorporated therein by reference.

Item 8.01 Other Events.

On September 24, 2026, the Company outlined its plans to share the following immunology and oncology pipeline updates in the fourth quarter of 2026.

Immunology

CLN-978

- The Company plans to share both multi-dose and single target dose regimen data from each of its ongoing Phase 1 clinical trials of CLN-978 in patients with (i) treatment-refractory moderate to severe systemic lupus erythematosus ("SLE"), (ii) difficult-to-treat rheumatoid arthritis ("RA") and (iii) treatment-refractory moderate to severe Sjögren's disease in December.

Velinotamig

- The Company plans to share multi-dose regimen data from the ongoing Phase 1 dose escalation portion of the clinical trial of velinotamig in patients with SLE being conducted by Chongqing Genrix Biopharmaceutical Co., Ltd. in China in a poster session at American College of Rheumatology Convergence 2026 in November.

Oncology

CLN-049

- The Company plans to share updated data from the dose escalation portion of its ongoing Phase 1 trial of CLN-049 in patients with relapsed/refractory acute myeloid leukemia in December.

Item 9.01 Financial Statements and Exhibits.

(d)

<u>Exhibit No.</u>	<u>Description</u>
99.1	Press release issued by Cullinan Therapeutics, Inc. on September 24, 2026, furnished herewith
104	Cover page from this Current Report on Form 8-K, formatted in Inline XBRL

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.

CULLINAN THERAPEUTICS, INC.

Date: September 24, 2026

By: /s/ Mary Kay Fenton
Mary Kay Fenton
Chief Financial Officer

Cullinan Therapeutics Highlights Fourth Quarter 2026 Milestones Across Immunology and Oncology T Cell Engager Portfolio

Velinotamig (BCMA TCE) multi-dose regimen data in SLE at ACR Convergence 2026 in November

CLN-978 (CD19 TCE) multi-dose regimen data in SLE, RA and Sjögren's disease in December

CLN-049 (FLT3 TCE) updated data from the Phase 1 dose escalation study in December

CAMBRIDGE, Mass. September 24, 2026 (GLOBE NEWSWIRE) -- Cullinan Therapeutics, Inc. (Nasdaq: CGEM; "Cullinan"), a clinical-stage biopharmaceutical company accelerating potential first- or best-in-class, disease-modifying T cell engagers in autoimmune diseases and cancer, today outlined fourth quarter 2026 milestones across its immunology and oncology pipeline.

"We look forward to providing several updates across our T cell engager programs in the fourth quarter of 2026. Starting with autoimmune diseases, for CLN-978 we look forward to sharing the most comprehensive clinical data set to date for a CD19 T cell engager across all indications, with multi-dose regimen data reported concurrently for SLE, RA, and now Sjögren's disease also. For velinotamig, we will provide multi-dose regimen data from the ongoing Phase 1 dose escalation study as we advance the program in plasma cell driven diseases. Together, our CD19- and BCMA-targeted programs reflect a differentiated approach to treating autoimmune diseases, aiming to address distinct disease drivers across a broad range of conditions. For CLN-049, we plan to provide an update with longer follow up from the dose escalation portion of our ongoing Phase 1 study in a broad, all-comer population of relapsed/refractory AML patients. We look forward to rapidly progressing this program and initiating our potentially registrational Phase 2 study, following our recent successful meeting with the FDA," said Nadim Ahmed, President and CEO of Cullinan Therapeutics.

The Company plans to share the following immunology and oncology pipeline updates in Q4 2026:

- **CLN-978 (CD19xCD3 T cell engager):** treatment-refractory moderate to severe systemic lupus erythematosus (SLE), difficult-to-treat rheumatoid arthritis (RA), and treatment-refractory moderate to severe Sjögren's disease (SjD)
 - Multi-dose and single target dose regimen data in SLE, RA, and SjD in December
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- **Velinotamig (BCMAxCD3 T cell engager):** treatment-refractory autoimmune diseases driven by long-lived plasma cells
 - Multi-dose regimen data from the ongoing Genrix Bio Phase 1 dose escalation study in SLE to be shared in poster session at ACR Convergence 2026 on November 8, 2026, 10:30 a.m. to 12:30 p.m. ET
- **CLN-049 (FLT3xCD3 T cell engager):** relapsed/refractory acute myeloid leukemia (AML)
 - Updated data from the dose escalation portion of the Phase 1 study in December

About Cullinan Therapeutics

Cullinan Therapeutics, Inc. (Nasdaq: CGEM) is a biopharmaceutical company developing potential first- or best-in-class, disease-modifying T cell engagers for autoimmune diseases and cancer. Cullinan pursues promising therapeutic targets while leveraging core expertise in T cell engagers, which are established in oncology and are now advancing into autoimmune diseases. With a clinical-stage pipeline built on a rigorous scientific approach and purposeful innovation, Cullinan is advancing its mission to deliver new standards of care for patients. Learn more about Cullinan at <https://cullinantherapeutics.com/>, and follow Cullinan on LinkedIn and X.

Forward-Looking Statements

This press release contains forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995. These forward-looking statements include, but are not limited to, express or implied statements regarding the company's beliefs and expectations regarding: our clinical development plans and anticipated timelines for our product candidates, the clinical and therapeutic potential of our product candidates, our plans regarding future data presentations and other statements that are not historical facts. The words "believe," "continue," "could," "estimate," "expect," "intends," "may," "plan," "potential," "project," "pursue," "will," and similar expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these identifying words.

Any forward-looking statements in this press release are based on management's current expectations and beliefs of future events and are subject to known and unknown risks and uncertainties that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. These risks include, but are not limited to, the following: uncertainty regarding the timing and results of clinical trial data and regulatory submissions; the risk that any NDAs, INDs, or other global regulatory submissions we may file with the United States Food and Drug Administration or other global regulatory agencies are not accepted or cleared on our expected timelines, or at all; the success of our clinical trials and preclinical studies; the risks related to our ability to protect and

maintain our intellectual property position; the risks related to manufacturing, supply, and distribution of our product candidates; the risk that any one or more of our product candidates, including those that are co-developed, will not be successfully developed and commercialized; the risk that the results of preclinical studies or clinical trials will not be predictive of future results in connection with future studies or clinical trials; the effect of changes in global economic conditions, including uncertainties related to international trade policies, tariffs and supply chain dynamics on our business and operations; and the success of any collaboration, partnership, license or similar agreements. These and other important risks and uncertainties discussed in our filings with the Securities and Exchange Commission, including under the caption “Risk Factors” in our most recent Annual Report on Form 10-K and subsequent filings with the SEC, could cause actual results to differ materially from those indicated by the forward-looking statements made in this press release. While we may elect to update such forward-looking statements at some point in the future, we disclaim any obligation to do so, even if subsequent events cause our views to change, except to the extent required by law. These forward-looking statements should not be relied upon as representing our views as of any date subsequent to the date of this press release. Moreover, except as required by law, neither the company nor any other person assumes responsibility for the accuracy and completeness of the forward-looking statements included in this press release. Any forward-looking statement included in this press release speaks only as of the date on which it was made.

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