

---

---

**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): May 05, 2025**

---

**RELAY THERAPEUTICS, INC.**

(Exact name of Registrant as Specified in Its Charter)

---

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

**001-39385**  
(Commission File Number)

**47-3923475**  
(IRS Employer  
Identification No.)

**399 Binney Street**  
**Cambridge, Massachusetts**  
(Address of Principal Executive Offices)

**02142**  
(Zip Code)

**Registrant's Telephone Number, Including Area Code: (617) 370-8837**

(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, par value \$0.001 per share	RLAY	Nasdaq Global 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 2.02 Results of Operations and Financial Condition.**

On May 5, 2025, Relay Therapeutics, Inc. announced its financial results for the quarter ended March 31, 2025. A copy of the press release is being furnished as Exhibit 99.1 to this Current Report on Form 8-K.

The information in this Current Report on Form 8-K and Exhibit 99.1 attached hereto is intended to be 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 filing.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits

99.1	<a href="#">Press release issued by Relay Therapeutics, Inc. on May 5, 2025, furnished herewith.</a>
104	Cover Page Interactive Data File (embedded within Inline XBRL 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.

**RELAY THERAPEUTICS, INC.**

Date: May 5, 2025

By: /s/ Brian Adams  
Brian Adams  
Chief Legal Officer

---



## Relay Therapeutics Reports First Quarter 2025 Financial Results and Corporate Updates

*Cash runway extended into 2029*

*Initiation of Phase 3 ReDiscover-2 trial on track for mid-2025*

*Initiated Phase 1 RLY-2608 vascular malformations clinical trial in Q1*

*Extended cash runway expected to fully fund the Company meaningfully past topline data for ReDiscover-2 trial and through clinical proof-of-concept data in Vascular Malformations*

*Approximately \$710 million in cash, cash equivalents and investments at end of Q1 2025*

Cambridge, Mass. – May 5, 2025 – Relay Therapeutics, Inc. (Nasdaq: RLAY), a clinical-stage precision medicine company transforming the drug discovery process by combining leading-edge computational and experimental technologies, today reported first quarter 2025 financial results and corporate updates.

“2025 is a year of execution across a range of high value clinical programs,” said Sanjiv Patel, M.D., President and Chief Executive Officer of Relay Therapeutics. “The ongoing changes to our cost base are designed to enable a full funding of key initiatives including generating topline data from the ReDiscover-2 trial and clinical proof-of-concept data in vascular malformations.”

### **Strategic Cost Reductions Implemented to Extend Runway into 2029 and to Fully Fund Key Value Drivers**

- These reductions help extend operating runway into 2029 and are aimed at funding key objectives, including:
    - Completion of ReDiscover-2 Phase 3 trial of RLY-2608 + fulvestrant in metastatic breast cancer well past topline data readout and additional breast cancer clinical trials
    - Execution of RLY-2608 Phase 1 vascular malformations trial through clinical proof-of-concept data
    - Advance Fabry and NRAS program to Investigational New Drug application (IND) readiness
    - Advance one research-stage program
  - **Specific cost reductions include:**
    - Over the past year, focused the research portfolio and platform on the highest value areas resulting in a reduction in the research run rate spend by approximately 80%
      - Reduced research-stage programs from four to one
      - Recent reduction in force by approximately 70 people
    - Executed a global out-license of RLY-4008 with Elevar Therapeutics, Inc. (Elevar) with potential for downstream economics
    - Phased the timing of entry into the clinic for Fabry and NRAS targeted programs
-

## RLY-2608 Highlights

- Breast Cancer
  - Initiation of Phase 3 ReDiscover-2 trial of RLY-2608 + fulvestrant in PI3K $\alpha$ -mutated, CDK4/6 pre-treated, HR+/HER2- advanced breast cancer remains on track for mid-2025
  - Abstract accepted to ASCO for update of Phase 1b ReDiscover trial of RLY-2608 + fulvestrant
    - Focus of the abstract is updated 600mg BID (fasted) doublet data with median follow-up now greater than 12 months
    - Poster Title: Updated efficacy of mutant-selective PI3K $\alpha$  inhibitor RLY-2608 in combination with fulvestrant in patients with PIK3CA-mutant HR+HER2- advanced breast cancer: ReDiscover trial
    - Date/Time: Monday, June 2, 10:00-1:00 p.m. ET (9:00-12:00 p.m. CT)
  - Continued advancement of the ongoing triplet cohorts with RLY-2608 + fulvestrant + atimociclib or ribociclib
  - Planning continues for development of next-generation endocrine therapy combinations with RLY-2608
- Vascular Malformations
  - Initiation of Phase 1 vascular malformations clinical trial in the first quarter of 2025

## First Quarter 2025 Financial Results

**Cash, Cash Equivalents and Investments:** As of March 31, 2025, cash, cash equivalents and investments totaled \$710.3 million, as compared to \$781.3 million as of December 31, 2024. The company expects its current cash, cash equivalents, and investments will be sufficient to fund its operating expenses and capital expenditure requirements into 2029.

**Revenue:** Revenue was \$7.7 million for the first quarter of 2025, as compared to \$10.0 million for the first quarter of 2024. The revenue recognized in the first quarter of 2025 was due to completion of all performance obligations under the company's Exclusive License Agreement with Elevar. The revenue recognized in the first quarter of 2024 was due to a milestone achieved under the Collaboration and License Agreement with Genentech, Inc.

**R&D Expenses:** Research and development expenses were \$73.8 million for the first quarter of 2025, as compared to \$82.4 million for the first quarter of 2024. The decrease was primarily due to the series of strategic choices made to streamline the research organization throughout 2024.

**G&A Expenses:** General and administrative expenses were \$18.7 million for the first quarter of 2025, as compared to \$19.8 million for the first quarter of 2024. The decrease was primarily due to a decrease in stock compensation expense, partially offset by costs to obtain the agreement with Elevar, which were expensed commensurate with the timing of revenue recognized.

**Net Loss:** Net loss was \$77.1 million for the first quarter of 2025, or a net loss per share of \$0.46, as compared to a net loss of \$81.4 million for the first quarter of 2024, or a net loss per share of \$0.62.

---

## **About Relay Therapeutics**

Relay Therapeutics (Nasdaq: RLAY) is a clinical-stage precision medicine company transforming the drug discovery process by combining leading-edge computational and experimental technologies with the goal of bringing life-changing therapies to patients. As the first of a new breed of biotech created at the intersection of complementary techniques and technologies, Relay Therapeutics aims to push the boundaries of what's possible in drug discovery. Its Dynamo® platform integrates an array of leading-edge computational and experimental approaches designed to drug protein targets that have previously been intractable or inadequately addressed. Relay Therapeutics' initial focus is on enhancing small molecule therapeutic discovery in targeted oncology and genetic disease indications. For more information, please visit [www.relaytx.com](http://www.relaytx.com) or follow us on Twitter.

## **Cautionary Note Regarding Forward-Looking Statements**

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, including, without limitation, implied and express statements regarding Relay Therapeutics' strategy, business plans and focus; the progress and timing of the clinical development of the programs across Relay Therapeutics' portfolio; the expected therapeutic benefits and potential efficacy and tolerability of RLY-2608, both as a monotherapy and in combination with other agents, and its other programs, as well as the clinical data for RLY-2608; the interactions with regulatory authorities and any related approvals; the potential market opportunity for RLY-2608; the cash runway projection; the expected benefits resulting from the implementation of the cost saving measures and potential ability to fund key value drivers; and the expectations regarding Relay Therapeutics' use of capital and expenses. The words "may," "might," "will," "could," "would," "should," "plan," "anticipate," "intend," "believe," "expect," "estimate," "seek," "predict," "future," "project," "potential," "continue," "target" and similar words or expressions, or the negative thereof, 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 and are subject to a number of risks, uncertainties and important factors that may cause actual events or results to differ materially from those expressed or implied by any forward-looking statements contained in this press release, including, without limitation, risks associated with: Relay Therapeutics' restructuring activities may be more costly or time-consuming than we expect or may not achieve their intended results; the timing, execution, and expected impact of Relay Therapeutics' restructuring plans (including the scope and timing of workforce reductions); the expected decrease in annual spending; the expected sufficiency of Relay Therapeutics' existing cash resources; the internal and external costs required for Relay Therapeutics' ongoing and planned activities, and the resulting impact on expense and use of cash, may be higher than expected, which may cause the company to use cash more quickly than expected or to change or curtail some of Relay Therapeutics' plans or both; the impact of global economic uncertainty, geopolitical instability and conflicts, or public health epidemics or outbreaks of an infectious disease on countries or regions in which Relay Therapeutics has operations or does business, as well as on the timing and anticipated results of its clinical trials, strategy, future operations and profitability; significant political, trade, or regulatory developments, such as tariffs, beyond Relay Therapeutics' control; the delay or pause of any current or planned clinical trials or the development of Relay Therapeutics' drug candidates; the risk that the preliminary or interim results of its preclinical or clinical trials may not be predictive of future or final results in connection with future clinical trials of its product candidates and that interim and early clinical data may change as more

---

patient data become available and are subject to audit and verification procedures; Relay Therapeutics' ability to successfully demonstrate the safety and efficacy of its drug candidates; the timing and outcome of its planned interactions with regulatory authorities; and obtaining, maintaining and protecting its intellectual property. These and other risks and uncertainties are described in greater detail in the section entitled "Risk Factors" in Relay Therapeutics' most recent Annual Report on Form 10-K and Quarterly Report on Form 10-Q, as well as any subsequent filings with the Securities and Exchange Commission. In addition, any forward-looking statements represent Relay Therapeutics' views only as of today and should not be relied upon as representing its views as of any subsequent date. Relay Therapeutics explicitly disclaims any obligation to update any forward-looking statements. No representations or warranties (expressed or implied) are made about the accuracy of any such forward-looking statements.

**Contact:**

Pete Rahmer  
[prahmer@relaytx.com](mailto:prahmer@relaytx.com)

**Media:**

Dan Budwick  
1AB  
973-271-6085  
[dan@1abmedia.com](mailto:dan@1abmedia.com)

---

Relay Therapeutics, Inc.  
Condensed Consolidated Statements of Operations and Comprehensive Loss  
(In thousands, except share and per share data)  
(Unaudited)

	<b>Three Months Ended March 31,</b>	
	<b>2025</b>	<b>2024</b>
<b>Revenue:</b>		
License and other revenue	\$ 7,679	\$ 10,007
<b>Total revenue</b>	<b>7,679</b>	<b>10,007</b>
<b>Operating expenses:</b>		
Research and development expenses	\$ 73,809	\$ 82,403
Change in fair value of contingent consideration liability	—	(1,832)
General and administrative expenses	18,739	19,799
<b>Total operating expenses</b>	<b>92,548</b>	<b>100,370</b>
<b>Loss from operations</b>	<b>(84,869)</b>	<b>(90,363)</b>
<b>Other income:</b>		
Interest income	7,813	8,951
Other (expense) income	(9)	25
<b>Total other income, net</b>	<b>7,804</b>	<b>8,976</b>
<b>Net loss</b>	<b>\$ (77,065)</b>	<b>\$ (81,387)</b>
<b>Net loss per share, basic and diluted</b>	<b>\$ (0.46)</b>	<b>\$ (0.62)</b>
<b>Weighted average shares of common stock, basic and diluted</b>	<b>169,233,155</b>	<b>130,843,013</b>
<b>Other comprehensive income (loss):</b>		
Unrealized holding gain (loss)	1,029	(962)
<b>Total other comprehensive income</b>	<b>1,029</b>	<b>(962)</b>
<b>Total comprehensive loss</b>	<b>\$ (76,036)</b>	<b>\$ (82,349)</b>

Relay Therapeutics, Inc.  
Selected Condensed Consolidated Balance Sheet Data  
(In thousands)  
(Unaudited)

	<b>March 31, 2025</b>		<b>December 31, 2024</b>
Cash, cash equivalents and investments	\$ 710,355	\$	781,323
Working capital (1)	702,607		758,475
<b>Total assets</b>	<b>799,362</b>		<b>871,296</b>
Total liabilities	78,281		93,504
<b>Total stockholders' equity</b>	<b>721,081</b>		<b>777,792</b>
Restricted cash	2,119		2,119

(1) Working capital is defined as current assets less current liabilities.

---

