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): February 6, 2026

Roivant Sciences Ltd.

(Exact Name of Registrant as Specified in Charter)

Bermuda	**001-40782**	**98-1173944**
(State or Other Jurisdiction of Incorporation)	**(Commission File Number)**	**(I.R.S. Employer Identification No.)**

7th Floor
50 Broadway
London SW1H 0DB
United Kingdom

1 Pennsylvania Plaza
Floor 54
New York, NY
10119[1]

Viaduktstrasse 8
4051 Basel
Switzerland[1]
(Addresses of Principal Executive Offices, and Zip Code)

+44 207 400 3347
Registrant's Telephone Number, Including Area Code

Not Applicable
(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 (*see* General Instruction A.2. below):

☐ Written communication 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 Shares, $0.0000000341740141 per share	**ROIV**	**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 (17 CFR §230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR §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. ☐

[1] Addresses of wholly-owned subsidiaries of the Registrant.

Item 2.02. Results of Operations and Financial Condition.

On February 6, 2026, Roivant Sciences Ltd. (the "Company") issued a press release announcing its financial results for the fiscal quarter ended December 31, 2025. A copy of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K.

The information set forth under this "Item 2.02. Results of Operations and Financial Condition" (including Exhibit 99.1) shall not be deemed to be "filed" for the purposes of Section 18 of the Securities Exchange Act of 1934, as amended, nor shall it be incorporated by reference in any filing made by the Company pursuant to the Securities Act of 1933, as amended, other than to the extent that such filing incorporates by reference any or all of such information by express reference thereto.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No.	Description of Exhibit
99.1	Roivant Science Ltd. Press Release, dated February 6, 2026
104	Cover Page Interactive Data File (embedded with 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.

ROIVANT SCIENCES LTD.

By: /s/ Keyur Parekh

Name: Keyur Parekh

Title: Authorized Signatory

Dated: February 6, 2026

Exhibit 99.1

Roivant Announces Positive Phase 2 Results for Brepocitinib in Cutaneous Sarcoidosis (CS) and Reports Financial Results for the Third Quarter Ended December 31, 2025

- Brepocitinib 45 mg significantly improved cutaneous sarcoidosis disease activity, achieving a 22.3-point improvement in mean CSAMI-A at Week 16 versus a 0.7-point improvement in placebo (Δ21.6 $p<0.0001$). Brepocitinib demonstrated rapid, deep and sustained improvements across all other efficacy endpoints measured with consistent safety profile

- Priovant plans to progress CS to a pivotal program with a Phase 3 study starting in calendar year 2026 following engagement with the FDA, representing the third indication with a pivotal program for brepocitinib

- New Drug Application (NDA) was submitted to the FDA for brepocitinib in dermatomyositis (DM). Topline data from Phase 3 studies in non-infectious uveitis (NIU) are expected in the second half of calendar year 2026. Roivant continues to actively explore other indications for brepocitinib

- IMVT-1402 potentially registrational trial in difficult-to-treat rheumatoid arthritis (D2T RA) fully enrolled, with topline data expected in the second half of calendar year 2026; topline data from the proof-of-concept trial in cutaneous lupus erythematous (CLE) expected in the second half of calendar year 2026

- Mosliciguat Phase 2 trial in pulmonary hypertension associated with interstitial lung disease (PH-ILD) fully enrolled, with topline data expected in the second half of calendar year 2026

- LNP litigation continues to progress, with favorable summary judgment decision in U.S. Moderna case affirming Genevant's view of section 1498: the significant majority of liability belongs in the current case against Moderna. Jury trial in the U.S. Moderna case scheduled for March 2026 and first major hearings in international proceedings expected in the first half of calendar year 2026

- Roivant-led Immunovant financing alongside key institutional investors generated gross proceeds to Immunovant of approximately $550 million, extending Immunovant’s cash runway to the launch of IMVT-1402 in Graves’ disease (GD)

- Roivant reported consolidated cash, cash equivalents, restricted cash and marketable securities of $4.5 billion as of December 31, 2025, supporting cash runway into profitability

- Roivant will host a live conference call and webcast at 8:00 a.m. ET on Friday, February 6, 2026, to discuss positive Phase 2 results for brepocitinib in CS and report its financial results for the third quarter ended December 31, 2025

BASEL, Switzerland and LONDON and NEW YORK, February 6, 2026 – Roivant (Nasdaq: ROIV) today announced positive Phase 2 results for brepocitinib in cutaneous sarcoidosis (CS) and reported its financial results for the third quarter ended December 31, 2025.

"2025 was a transformative year for Roivant, driven by the quality of our clinical execution and its resulting data. We are entering 2026 with early momentum across our late stage pipeline with ambitions of another groundbreaking year. The positive data for brepocitinib in CS represents a breakthrough for the field, generating clinically meaningful outcomes for patients in need in the first ever positive placebo-controlled trial for any therapy in CS," said Matt Gline, CEO of Roivant. "Together with this data, I’m proud of our continued execution elsewhere in our pipeline, including with today's announcement of the NDA filing for brepocitinib in dermatomyositis and the completion of enrollment in multiple large studies setting the stage for the year ahead."

BEACON Study Results

The BEACON study enrolled 31 patients across 15 sites in the United States, randomized 3:2:2 to once daily brepocitinib 45 mg, 15 mg, or placebo with a 16-week treatment period. The brepocitinib 45 mg arm comprised the most treatment-refractory group, with the highest percentage of patients with longstanding disease, damage, and difficult-to-treat plaque-predominant morphology. Despite this, patients in the 45 mg arm achieved meaningful clinical improvement compared to

placebo, including 100% response rates on multiple endpoints. Brepocitinib 15 mg patients also improved considerably, with numerically similar improvement to the 45 mg arm on lower-bar endpoints and evidence of dose-dependent benefit seen on higher bar endpoints and patient reported outcomes. Placebo patients experienced almost no improvement, consistent with natural disease course.

On the Cutaneous Sarcoidosis Activity and Morphology Instrument – Activity score (CSAMI-A), brepocitinib 45 mg achieved a 22.3-point mean improvement at Week 16 versus a 0.7-point improvement in placebo (Δ21.6 P<0.0001). Statistically significant separation was observed as early as Week 4 and maintained at all timepoints thereafter. One hundred percent of brepocitinib 45 mg patients achieved at least a 10-point improvement on CSAMI-A compared to 14% of placebo patients, and 62% of brepocitinib 45 mg patients achieved CSAMI-A <5 (functional remission), compared to 0% of placebo patients.

Treatment Arm	**Mean CSAMI-A Change from Baseline at Week 16**	**Achievement of CSAMI-A ≥ 10-point Reduction at Week 16**	**Achievement of CSAMI-A < 5 (Functional Remission) at Week 16**
45 mg (n = 13)	-22.3	100%	62%
15 mg (n = 11)	-22.2	73%	46%
Placebo (n = 7)	-0.7	14%	0%
45 mg vs. placebo	Δ -21.6; P<0.0001	Δ 86%; P=0.0002	Δ 62%; P=0.0147
15 mg vs. placebo	Δ -21.5; P<0.0001	Δ 58%; P=0.0498	Δ 46%; P=0.1013

On the Investigator's Global Assessment (IGA), 69% of brepocitinib 45 mg patients achieved the gold standard two-point improvement to Clear (0) / Almost Clear (1), compared to 0% of placebo patients (Δ 69% P=.0047). Brepocitinib 45 mg also demonstrated statistically significant improvement over placebo on key patient reported outcomes, including the King's Sarcoidosis Questionnaire (KSQ) Skin Domain, the Skindex-16, and the Patient's Global Impression of Change (PGI-C). On PGI-C, 100% of patients receiving the 45 mg dose reported improvement from baseline, compared to 29% of placebo patients (Δ 71% P=.0014).

Brepocitinib was well tolerated during the study treatment period, with no Serious Adverse Events (SAEs) and all Adverse Events (AEs) graded mild or moderate in severity. Brepocitinib has been evaluated in more than 1,500 patients and subjects, with an observed safety profile consistent with approved JAK1 and TYK2 inhibitors.

Priovant plans to progress CS to a pivotal program with a Phase 3 study starting in calendar year 2026 following engagement with the FDA, representing the third indication with a pivotal program for brepocitinib.

"The BEACON study is a watershed moment for the sarcoidosis field, and most importantly, for our patients," said Dr. Misha Rosenbach, MD, Professor of Dermatology and Rheumatology and Director of the Cutaneous Sarcoidosis Program at the Hospital of the University of Pennsylvania. "This is an incredible milestone for a historically neglected disease – the study drug showed a clear difference in patients who received the medication compared to placebo, both from the patient and the physician perspective, and appeared to be well tolerated. This is the sort of data you dream of seeing when you look at trial results – and I would call this a transformational moment for sarcoidosis."

Recent Developments

- **Priovant**: In February 2026, Priovant announced positive Phase 2 results for brepocitinib in cutaneous sarcoidosis (CS). Brepocitinib 45 mg significantly improved CS disease activity, achieving a 22.3-point improvement in mean CSAMI-A at Week 16 versus a 0.7-point improvement in placebo (Δ21.6 p<0.0001). Brepocitinib demonstrated rapid, deep and sustained improvements across all other efficacy endpoints measured with consistent safety profile. New Drug Application (NDA) was submitted to the FDA for brepocitinib in dermatomyositis (DM). Topline data from Phase 3 studies in non-infectious uveitis (NIU) are expected in the second half of calendar year 2026.

- **Immunovant**: Potentially registrational trial for IMVT-1402 in difficult-to-treat rheumatoid arthritis (D2T RA) fully enrolled, with topline data expected in the second half of calendar year 2026. All other clinical development

timelines remain on track for IMVT-1402 across announced indications, including potentially registrational trials in Graves' disease (GD), myasthenia gravis (MG), chronic inflammatory demyelinating polyneuropathy (CIDP), and Sjögren's disease (SjD), and a proof-of-concept trial in cutaneous lupus erythematosus (CLE). Roivant-led Immunovant financing alongside key institutional investors generated gross proceeds to Immunovant of approximately $550 million, extending Immunovant's cash runway to the launch of IMVT-1402 in GD.

- **Pulmovant**: Phase 2 trial of mosliciguat in pulmonary hypertension associated with interstitial lung disease (PH-ILD) fully enrolled, with topline data expected in the second half of calendar year 2026.

- **Genevant**: Favorable summary judgment decision in U.S. Moderna case affirms Genevant's view of section 1498: the significant majority of liability belongs in the current case against Moderna. Jury trial in U.S. Moderna case scheduled for March 2026 and international proceedings continue with first major hearings expected in the first half of calendar year 2026.

- **Roivant**: Roivant reported consolidated cash, cash equivalents, restricted cash and marketable securities of $4.5 billion as of December 31, 2025, supporting cash runway into profitability.

Major Upcoming Milestones

- **Priovant** plans to progress CS to a pivotal program with a Phase 3 study starting in calendar year 2026 following engagement with the FDA, representing the third indication with a pivotal program for brepocitinib. Topline data from the Phase 3 trial of brepocitinib in NIU are expected in the second half of calendar year 2026. Roivant continues to actively explore other indications for brepocitinib.

- **Immunovant** expects to report topline data from the potentially registrational trial of IMVT-1402 in D2T RA and proof-of-concept trial of IMVT-1402 in CLE in the second half of calendar year 2026. Immunovant anticipates sharing topline data from both of its Phase 3 studies to evaluate batoclimab as a treatment for active moderate to severe thyroid eye disease (TED) in the first half of calendar year 2026. In calendar year 2027, topline data are expected across potentially registrational trials of IMVT-1402 in GD and MG.

- **Pulmovant** expects to report topline data from the ongoing Phase 2 trial of mosliciguat in PH-ILD in the second half of calendar year 2026.

- **Genevant** LNP litigation continues to progress, with summary judgment phase in the U.S. Moderna case ongoing and jury trial in the U.S. Moderna case scheduled for March 2026. First major hearings in international proceedings expected in the first half of calendar year 2026. Discovery phase of Pfizer/BioNTech case is ongoing.

Third Quarter Ended December 31, 2025 Financial Summary

Cash Position and Marketable Securities

As of December 31, 2025, the Company had consolidated cash, cash equivalents, restricted cash and marketable securities of approximately $4.5 billion.

Research and Development Expenses

Research and development (R&D) expenses increased by $23.8 million to $165.4 million for the three months ended December 31, 2025, compared to $141.6 million for the three months ended December 31, 2024. This increase was primarily driven by an increase in program-specific costs of $11.3 million and share-based compensation of $8.4 million.

The $11.3 million increase in program-specific costs was primarily driven by increases of $5.7 million related to mosliciguat and $5.3 million related to brepocitinib, reflecting the progression of our programs.

The $8.4 million increase in share-based compensation expense, which is an unallocated internal cost, was primarily driven by incremental share-based compensation expense as a result of the exchange of certain vested equity granted under the Priovant Holdings, Inc. ("Priovant") 2021 Equity Incentive Plan for RSL common shares (the "Exchange Offer").

Non-GAAP R&D expenses were $146.7 million for the three months ended December 31, 2025, compared to $131.2 million for the three months ended December 31, 2024.

General and Administrative Expenses

General and administrative (G&A) expenses increased by $33.5 million to $175.1 million for the three months ended December 31, 2025, compared to $141.5 million for the three months ended December 31, 2024. This increase was primarily due to an impairment loss of $17.1 million related to the relocation of the U.S. corporate headquarters of Roivant Sciences, Inc. and an increase of $17.5 million in share-based compensation expense, primarily driven by the Priovant Exchange Offer. These increases were partially offset by a decrease of $5.3 million in personnel-related expense, primarily due to the conclusion of one-time cash retention awards under the Cash Bonus Program.

Non-GAAP G&A expenses were $71.0 million for the three months ended December 31, 2025, compared to $71.1 million for the three months ended December 31, 2024.

Income from discontinued operations, net of tax

Income from discontinued operations, net of tax was $327.0 million for the three months ended December 31, 2024 and reflects the gain on sale of subsidiary interests resulting from the sale of our entire equity interest in our majority-owned subsidiary, Dermavant, to Organon in October 2024, partially offset by Dermavant's net losses.

Loss from continuing operations, net of tax

Loss from continuing operations, net of tax was $313.7 million for the three months ended December 31, 2025, compared to $208.9 million for the three months ended December 31, 2024. On a basic and diluted per common share basis, loss from continuing operations, net of tax was $0.38 and $0.22, respectively, for the three months ended December 31, 2025 and 2024.

Non-GAAP loss from continuing operations, net of tax was $167.0 million for the three months ended December 31, 2025, compared to $143.7 million for the three months ended December 31, 2024.

ROIVANT SCIENCES LTD.
Selected Balance Sheet Data
(unaudited, in thousands)

	December 31, 2025	March 31, 2025
Cash, cash equivalents and restricted cash	$ 1,488,770	$ 2,725,661
Marketable securities	3,051,708	2,171,480
Total assets	5,225,368	5,436,940
Total liabilities	251,144	249,742
Total shareholders' equity	4,974,224	5,187,198
Total liabilities and shareholders' equity	5,225,368	5,436,940

ROIVANT SCIENCES LTD.
Condensed Consolidated Statements of Operations
(unaudited, in thousands, except share and per share amounts)

	Three Months Ended December 31,		Nine Months Ended December 31,	
	2025	2024	2025	2024
Revenue	$ 1,999	$ 9,018	$ 5,740	$ 21,483
Operating expenses:				
Cost of revenues	700	259	965	706
Research and development (includes $18,123 and $9,685 of share-based compensation expense for the three months ended December 31, 2025 and 2024 and $40,218 and $30,128 for the nine months ended December 31, 2025 and 2024, respectively)	165,380	141,595	482,867	405,175
General and administrative (includes $86,874 and $69,386 of share-based compensation expense for the three months ended December 31, 2025 and 2024 and $228,778 and $165,670 for the nine months ended December 31, 2025 and 2024, respectively)	175,072	141,545	452,216	444,318
Total operating expenses	341,152	283,399	936,048	850,199
Gain on sale of Telavant net assets	—	—	—	110,387
Loss from operations	(339,153)	(274,381)	(930,308)	(718,329)
Change in fair value of investments	(21,592)	21,314	(130,968)	(42,287)
Change in fair value of liability instruments	24,416	(2,147)	47,704	(1,632)
Interest income	(43,266)	(61,851)	(136,929)	(203,751)
Other (income) expense, net	(8,342)	2,816	8,560	7,877
Loss from continuing operations before income taxes	(290,369)	(234,513)	(718,675)	(478,536)
Income tax expense (benefit)	23,332	(25,568)	34,976	(1,147)
Loss from continuing operations, net of tax	(313,701)	(208,945)	(753,651)	(477,389)
Income from discontinued operations, net of tax	—	327,020	—	373,030
Net (loss) income	(313,701)	118,075	(753,651)	(104,359)
Net loss attributable to noncontrolling interests	(47,810)	(51,306)	(150,886)	(138,853)
Net (loss) income attributable to Roivant Sciences Ltd.	$ (265,891)	$ 169,381	$ (602,765)	$ 34,494
Amounts attributable to Roivant Sciences Ltd.:				
Loss from continuing operations, net of tax	$ (265,891)	$ (157,639)	$ (602,765)	$ (338,691)
Income from discontinued operations, net of tax	—	327,020	—	373,185
Net (loss) income attributable to Roivant Sciences Ltd.	$ (265,891)	$ 169,381	$ (602,765)	$ 34,494
Basic and diluted net (loss) income per common share:				
Basic and diluted loss from continuing operations	$ (0.38)	$ (0.22)	$ (0.88)	$ (0.46)
Basic and diluted income from discontinued operations	$ —	$ 0.45	$ —	$ 0.51
Basic and diluted net (loss) income per common share	$ (0.38)	$ 0.23	$ (0.88)	$ 0.05
Weighted average shares outstanding:				
Basic	696,859,682	722,716,168	686,052,379	731,318,202
Diluted	696,859,682	722,716,168	686,052,379	731,318,202

ROIVANT SCIENCES LTD.
Reconciliation of GAAP to Non-GAAP Financial Measures
(unaudited, in thousands)

	Note	Three Months Ended December 31, 2025	Three Months Ended December 31, 2024	Nine Months Ended December 31, 2025	Nine Months Ended December 31, 2024
Loss from continuing operations, net of tax		$ (313,701)	$ (208,945)	(753,651)	(477,389)
Adjustments:					
Research and development:					
Share-based compensation	(1)	18,123	9,685	40,218	30,128
Depreciation and amortization	(2)	548	728	2,010	2,147
General and administrative:					
Share-based compensation	(1)	86,874	69,386	228,778	165,670
Depreciation and amortization	(2)	142	1,083	700	3,267
Impairment loss from relocation of Roivant Sciences, Inc.'s headquarters	(3)	17,098	—	17,098	—
Gain on sale of Telavant net assets	(4)	—	—	—	(110,387)
Other:					
Change in fair value of investments	(5)	(21,592)	21,314	(130,968)	(42,287)
Change in fair value of liability instruments	(6)	24,416	(2,147)	47,704	(1,632)
Estimated income tax impact from adjustments	(7)	21,058	(34,786)	23,174	(38,976)
Adjusted loss from continuing operations, net of tax (Non-GAAP)		**$ (167,034)**	**$ (143,682)**	**$ (524,937)**	**$ (469,459)**

	Note	Three Months Ended December 31, 2025	Three Months Ended December 31, 2024	Nine Months Ended December 31, 2025	Nine Months Ended December 31, 2024
Research and development expenses		**$ 165,380**	**$ 141,595**	**$ 482,867**	**$ 405,175**
Adjustments:					
Share-based compensation	(1)	18,123	9,685	40,218	30,128
Depreciation and amortization	(2)	548	728	2,010	2,147
Adjusted research and development expenses (Non-GAAP)		**$ 146,709**	**$ 131,182**	**$ 440,639**	**$ 372,900**

	Note	Three Months Ended December 31, 2025	Three Months Ended December 31, 2024	Nine Months Ended December 31, 2025	Nine Months Ended December 31, 2024
General and administrative expenses		**$ 175,072**	**$ 141,545**	**$ 452,216**	**$ 444,318**
Adjustments:					
Share-based compensation	(1)	86,874	69,386	228,778	165,670
Depreciation and amortization	(2)	142	1,083	700	3,267
Impairment loss from relocation of Roivant Sciences, Inc.'s headquarters	(3)	17,098	—	17,098	—
Adjusted general and administrative expenses (Non-GAAP)		**$ 70,958**	**$ 71,076**	**$ 205,640**	**$ 275,381**

Notes to non-GAAP financial measures:

(1) Represents non-cash share-based compensation expense.

(2) Represents non-cash depreciation and amortization expense.

(3) Represents a loss on impairment of the operating lease right-of-use asset and leasehold improvements for the former U.S. corporate headquarters of Roivant Sciences, Inc., following relocation to a new office space and execution of an agreement to sublease the former office space during the three months ended December 31, 2025.

(4) Represents a gain on the sale of Telavant net assets to Roche due to achievement of a one-time milestone in June 2024.

(5) Represents the unrealized (gain) loss on equity investments in unconsolidated entities that are accounted for at fair value with changes in value reported in earnings.

(6) Represents the change in fair value of liability instruments, which is non-cash and primarily includes the loss (gain) relating to the measurement and recognition of fair value on a recurring basis of certain liabilities.

(7) Represents the estimated tax effect of the adjustments.

Investor Conference Call Information

Roivant will host a live conference call and webcast at 8:00 a.m. ET on Friday, February 6, 2026, to report its financial results for the third quarter ended December 31, 2025, and provide a corporate update.

To access the conference call by phone, please register online using this registration link. The presentation and webcast details will also be available under "Events & Presentations" in the Investors section of the Roivant website at https://investor.roivant.com/news-events/events. The archived webcast will be available on Roivant's website after the conference call.

About Roivant

Roivant (Nasdaq: ROIV) is a biopharmaceutical company that aims to improve the lives of patients by accelerating the development and commercialization of medicines that matter. Roivant's pipeline includes brepocitinib, a potent small molecule inhibitor of TYK2 and JAK1 in development for the treatment of dermatomyositis, non-infectious uveitis and cutaneous sarcoidosis; IMVT-1402 and batoclimab, fully human monoclonal antibodies targeting FcRn in development across several IgG-mediated autoimmune indications; and mosliciguat, an inhaled sGC activator in development for pulmonary hypertension associated with interstitial lung disease. We advance our pipeline by creating nimble subsidiaries or "Vants" to develop and commercialize our medicines and technologies. Beyond therapeutics, Roivant also incubates discovery-stage companies and health technology startups complementary to its biopharmaceutical business. For more information, visit https://roivant.com.

Roivant Forward-Looking Statements

This press release contains forward-looking statements. Statements in this press release may include statements that are not historical facts and are considered forward-looking within the meaning of Section 27A of the Securities Act of 1933, as amended (the "Securities Act"), and Section 21E of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), which are usually identified by the use of words such as "anticipate," "believe," "continue," "could," "estimate," "expect," "intends," "may," "might," "plan," "possible," "potential," "predict," "project," "should," "would" and variations of such words or similar expressions. The words may identify forward-looking statements, but the absence of these words does not mean that a statement is not forward-looking. We intend these forward-looking statements to be covered by the safe harbor provisions for forward-looking statements contained in Section 27A of the Securities Act and Section 21E of the Exchange Act.

Our forward-looking statements include, but are not limited to, statements regarding our or our management team's expectations, hopes, beliefs, intentions or strategies regarding the future, and statements that are not historical facts, including statements about the clinical and therapeutic potential of our product candidates, the availability and success of topline results from our ongoing clinical trials and any commercial potential of our product candidates following applicable regulatory approvals. In addition, any statements that refer to projections, forecasts or other characterizations of future events, results or circumstances, including any underlying assumptions, are forward-looking statements. Actual results may differ materially from those contemplated in these statements due to a variety of risks, uncertainties and other factors.

Although we believe that our plans, intentions, expectations and strategies as reflected in or suggested by those forward-looking statements are reasonable, we can give no assurance that the plans, intentions, expectations or strategies will be attained or achieved. Furthermore, actual results may differ materially from those described in the forward-looking statements and will be affected by a number of risks, uncertainties and assumptions, including, but not limited to, those risks set forth in the Risk Factors section of our filings with the U.S. Securities and Exchange Commission. Moreover, we operate in a very competitive and rapidly changing environment in which new risks emerge from time to time. These forward-looking statements are based upon the current expectations and beliefs of our management as of the date of this press release, and are subject to certain risks and uncertainties that could cause actual results to differ materially from those described in the forward-looking statements. Except as required by applicable law, we assume no obligation to update publicly any forward-looking statements, whether as a result of new information, future events or otherwise.

Contacts:

Investors
Keyur Parekh
keyur.parekh@roivant.com

Media

Stephanie Lee
stephanie.lee@roivant.com