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): **July 28, 2026**

RELMADA THERAPEUTICS, INC.
(Exact name of registrant as specified in its charter)

Nevada	**001-39082**	**45-5401931**
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

2222 Ponce de Leon Blvd., Floor 3 Coral Gables, FL	**33134**
(Address of principal executive offices)	(Zip Code)

Registrant's telephone number, including area code: **(786) 629-1376**

(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 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	Name of each exchange on which registered
Common stock, $0.001 par value per share	**RLMD**	**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 7.01 Regulation FD Disclosure.

On July 28, 2026, Relmada Therapeutics, Inc. (the "Company"), issued a press release announcing the appointment of Bipin Dalmia, PhD, MBA as the Company's Chief Business. Pursuant to Regulation FD, the press release is furnished with this Current Report as Exhibit 99.1.

The information set forth in Item 7.01 of this Current Report on Form 8-K and in the attached Exhibit 99.1 is deemed to be "furnished" and 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. The information set forth in Item 7.01 of this Current Report on Form 8-K, including Exhibit 99.1, shall not be deemed incorporated by reference into any filing under the Exchange Act or the Securities Act of 1933, as amended (the "Securities Act"), regardless of any general incorporation language in such filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No.	Description
99.1*	Press release issued on July 28, 2026
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

* *This Exhibit attached to this Form 8-K shall not be deemed "filed" for purposes of Section 18 of the Exchange Act or otherwise subject to liability under that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act or the Exchange Act, except as expressly set forth by specific reference in such filing.*

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.

Dated: July 28, 2026

RELMADA THERAPEUTICS, INC.

By: */s/ Sergio Traversa*
Name: Sergio Traversa
Title: Chief Executive Officer

Exhibit 99.1

Relmada Therapeutics Appoints Uro-Oncology Leader Bipin Dalmia as Chief Business Officer

- *Brings nearly three decades of leadership spanning NMIBC, corporate strategy and business development to help advance NDV-01 and support the Company's long-term growth and value creation.*

CORAL GABLES, FL – July 28, 2026 (GlobeNewswire) – **Relmada Therapeutics, Inc.** (Nasdaq: RLMD, "Relmada" or the "Company"), a clinical-stage biotechnology company advancing innovative therapies for oncology and central nervous system disorders, today announced the appointment of **Bipin Dalmia, PhD, MBA**, as Chief Business Officer. Bipin will lead Relmada's corporate strategy, commercial planning, and business development activities, reporting directly to Chief Executive Officer Sergio Traversa.

Bipin Dalmia joins Relmada with nearly three decades of biopharmaceutical leadership, combining extensive experience in corporate strategy and business development with deep expertise in non-muscle invasive bladder cancer (NMIBC). As the Company advances NDV-01, its novel sustained-release intravesical therapy for patients with NMIBC, Bipin's appointment reflects Relmada's rapid evolution toward late-stage development and its commitment to building the commercial and strategic capabilities needed to maximize the value of NDV-01.

"Bipin combines a deep understanding of the evolving NMIBC treatment and competitive landscape, with exceptional experience in helping bring the first FDA-approved intravesical gene therapy to bladder cancer patients. His experience and perspective will be invaluable in helping us maximize the clinical, commercial and long-term potential of NDV-01," said **Sergio Traversa, Chief Executive Officer of Relmada Therapeutics.** "In addition to advancing NDV-01, Bipin's experience in corporate strategy, business development and value creation will strengthen Relmada as we evaluate opportunities to maximize long-term shareholder value."

"I am excited to join Relmada at such an important stage in the Company's evolution. What drew me to this opportunity was the chance to help realize the full potential of NDV-01. The NMIBC treatment landscape is entering a period of rapid innovation, yet patients continue to face significant unmet needs. NDV-01's differentiated product profile and encouraging clinical data generated to date position it to become a best-in-class intravesical therapy across the NMIBC disease spectrum," said **Bipin Dalmia, Chief Business Officer.** "I look forward to working alongside the outstanding Relmada team to help advance the next phase of NDV-01's development and, if successful, bring this innovative therapy to patients as quickly as possible."

About Bipin Dalmia, PhD, MBA

Bipin Dalmia is a senior biopharmaceutical executive with nearly three decades of leadership roles across corporate strategy, portfolio management, commercial planning, business development and M&A in biotechnology, specialty pharmaceuticals, and global pharmaceutical companies.

Most recently, he served as **Senior Vice President, Global Head of Portfolio Strategy & Business Development** at Ferring Pharmaceuticals, where he established and led this new global function. During his 14-year tenure at Ferring, in addition to global business development leadership, he led the company's Uro-Oncology Franchise, including the U.S. launch and global lifecycle and manufacturing strategy for ADSTILADRIN® – the first FDA-approved intravesical gene therapy for BCG-unresponsive NMIBC patients. Beyond ADSTILADRIN, Bipin led and completed more than 20 strategic licensing, partnership and M&A transactions across multiple therapeutic areas, modalities and development stages, while strengthening the company's portfolio management and capital allocation capabilities.

Prior to Ferring, Bipin held business development leadership positions at Biolex Therapeutics and earlier served in strategy, new product planning and business development roles at Syngenta and Novartis. He began his career as a research scientist. He currently serves on the Board of Directors of Ferring Ventures S.A. and on the Board of Directors of FinVector Oy.

Bipin Dalmia holds a PhD and an MS in Chemical Engineering from Iowa State University and an MBA from Drake University.

About Relmada Therapeutics, Inc.

Relmada Therapeutics is a clinical-stage biotechnology company focused on developing transformative therapies for oncology and central nervous system conditions. Its lead candidates, NDV-01 and sepranolone, are advancing through mid-stage clinical development with the potential to address significant unmet needs.

For more information, visit www.relmada.com

Forward-Looking Statements:

The Private Securities Litigation Reform Act of 1995 provides a safe harbor for forward-looking statements made by us or on our behalf. This press release contains statements which constitute "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934. Any statement that is not historical in nature is a forward-looking statement and may be identified by the use of words and phrases such as "if", "may", "expects", "anticipates", "believes", "will", "will likely result", "will continue", "plans to", "potential", "promising", and similar expressions. These statements are based on management's current expectations and beliefs and are subject to a number of risks, uncertainties and assumptions that could cause actual results to differ materially from those described in the forward-looking statements, including potential for Relmada's product candidates to fail to progress, potential for Phase 2 NDV-01 data to fail to continue to deliver positive results supporting further development, potential for clinical trials to fail to deliver statistically and/or clinically significant evidence of efficacy and/or safety, failure of interim or top-line results to accurately reflect the complete results of the trial, failure of planned or ongoing preclinical and clinical studies to demonstrate expected results, potential failure to continue to secure FDA agreement on the regulatory path for NDV-01 and/or sepranolone, or that future NDV-01 and/or sepranolone clinical results will be acceptable to the FDA, failure to secure adequate NDV-01 and/or sepranolone drug supply, failure of pending patent applications to result in issued patents, or issued patents being challenged and invalidated by third parties or not providing us with any competitive advantages, the Company's cash runway and sufficiency of the Company's cash resources and uncertainties inherent in estimating the Company's cash runway, future expenses and other financial results, including its ability to fund future operations, including clinical trials, and the other risk factors described under the heading "Risk Factors" set forth in the Company's reports filed with the SEC from time to time. No forward-looking statement can be guaranteed, and actual results may differ materially from those projected. Relmada undertakes no obligation to publicly update any forward-looking statement, whether as a result of new information, future events, or otherwise. Readers are cautioned that it is not possible to predict or identify all the risks, uncertainties and other factors that may affect future results and that the risks described herein are not a complete list.

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