UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

Report of Foreign Private Issuer
Pursuant to Rule 13a-16 or 15d-16
of the Securities Exchange Act of 1934

For the Month of May 2021

Commission File Number 001-35948

Kamada Ltd.
(Translation of registrant's name into English)

2 Holzman Street
Science Park, P.O. Box 4081
Rehovot 7670402
Israel
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F.

Form 20-F ☒ Form 40-F ☐

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ____

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7): ____

Indicate by check mark whether the registrant by furnishing the information contained in this Form is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes ☐ No ☒

If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b): 82-____

This Form 6-K is being incorporated by reference into the Registrant's Form S-8 Registration Statements, File Nos. 333-192720, 333-207933, 333-215983, 333-222891 and 333-233267.

On May 5, 2021, Kamada Ltd. issued a press release titled “Kamada to Announce First Quarter Ended March 31, 2021 Financial Results and Host Conference Call on May 12, 2021”. A copy of the press release is attached to this Form 6-K as Exhibit 99.1.

Exhibit No.	Description
99.1	Press Release, dated May 5, 2021

SIGNATURE

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.

Date: May 5, 2021

KAMADA LTD.

By: /s/ Yifat Philip
Yifat Philip
Vice President General Counsel and Corporate Secretary

EXHIBIT INDEX

EXHIBIT NO.	DESCRIPTION
99.1	Press Release dated May 5, 2021: "Kamada to Announce First Quarter Ended March 31, 2021 Financial Results and Host Conference Call on May 12, 2021"

Exhibit 99.1

Kamada to Announce First Quarter Ended March 31, 2021 Financial Results and Host Conference Call on May 12, 2021

Rehovot, Israel, May 5, 2021 -- Kamada Ltd. (NASDAQ & TASE: KMDA), a plasma-derived biopharmaceutical company, today announced that it will release financial results for the first quarter ended March 31, 2021, prior to the open of the U.S. financial markets on Wednesday, May 12, 2021.

Kamada management will host an investment community conference call on Wednesday, May 12, at 8:30am Eastern Time to discuss these results and answer questions. Shareholders and other interested parties may participate in the conference call by dialing 877-407-0792 (from within the U.S.), or 201-689-8263 (International) and entering the conference identification number: 13719388. The call will also be webcast live on the Internet on the Company's website at www.kamada.com.

The call will also be archived for 90 days on the Company's website at www.kamada.com.

About Kamada
Kamada Ltd. (the "Company") is a global specialty plasma-derived biopharmaceutical company with a diverse portfolio of marketed products, a robust development pipeline and industry-leading manufacturing capabilities. The Company's strategy is focused on driving profitable growth from its current commercial products, its plasma-derived development pipeline and its manufacturing expertise, while evolving into a vertically integrated plasma-derived company. The Company's two leading commercial products are GLASSIA® and KEDRRAB®. GLASSIA was the first liquid, ready-to-use, intravenous plasma-derived AAT product approved by the FDA. The Company markets GLASSIA in the U.S. through a strategic partnership with Takeda Pharmaceuticals Company Limited ("Takeda") and in other countries through local distributors. Pursuant to an agreement with Takeda, the Company will continue to produce GLASSIA for Takeda through 2021 and Takeda will initiate its own production of GLASSIA for the U.S. market in 2021, at which point Takeda will commence payment of royalties to the Company until 2040. KEDRAB is an FDA approved anti-rabies immune globulin (Human) for post-exposure prophylaxis treatment. KEDRAB is being marketed in the U.S. through a strategic partnership with Kedrion S.p.A. The Company has additional four plasma-derived products administered by injection or infusion, that are marketed through distributors in more than 15 countries, including Israel, Russia, Brazil, Argentina, India and other countries in Latin America and Asia. The Company has two leading development programs; a plasma-derived hyperimmune immunoglobulin (IgG) product as a potential treatment for coronavirus disease (COVID-19) and an inhaled AAT for the treatment of AAT deficiency for which the Company is currently conducting the InnovAATe clinical trial, a randomized, double-blind, placebo-controlled, pivotal Phase 3 trial. The Company leverages its expertise and presence in the Israeli pharmaceutical market to distribute in Israel more than 20 products that are manufactured by third parties and have recently added nine biosimilar products to its Israeli distribution portfolio, which, subject to EMA and the Israeli MOH approvals, are expected to be launched in Israel between the years 2022 and 2025. FIMI Opportunity Fund, the leading private equity investor in Israel, is the Company's lead shareholder, beneficially owning approximately 21% of the outstanding ordinary shares.

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