
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 March 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.

The following exhibit is attached:

99.1 [Kamada to Present at the Jefferies Global Plasma Summit](#)

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: March 9, 2021

KAMADA LTD.

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

EXHIBIT INDEX

<u>EXHIBIT NO.</u>	<u>DESCRIPTION</u>
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99.1	Kamada to Present at the Jefferies Global Plasma Summit
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Kamada to Present at the Jefferies Global Plasma Summit

REHOVOT, Israel – March 9, 2021 – Kamada Ltd. (NASDAQ and TASE: KMDA), a plasma-derived biopharmaceutical company, announced today that Amir London, Chief Executive Officer, will present a corporate overview at the inaugural Jefferies Virtual Global Plasma Summit, being held on March 11, 2021.

Mr. London is scheduled to present at 9:35 a.m. Eastern Time. The presentation will be accessible via a live webcast at: <https://www.com/webcast/jeff171/kmda/1823556>. A replay of the presentation will be accessible via the link and will be available for 30 days following the event.

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.

CONTACTS:

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