

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 February 2024

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 □

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, 333-233267 and 333-265866.

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: February 28, 2024 KAMADA LTD.

By: /s/ Nir Livneh

Nir Livneh

Vice President General Counsel and

Corporate Secretary

EXHIBIT INDEX

EXHIBIT NO. 99.1 DESCRIPTION Kamada to Anno Kamada to Announce Fourth Quarter and Fiscal Year 2023 Financial Results, and Provide 2024 Financial Guidance on March 6, 2024

Kamada to Announce Fourth Quarter and Fiscal Year 2023 Financial Results, and Provide 2024 Financial Guidance on March 6, 2024

Company to Host Conference Call at 8:30am ET

Rehovot, Israel, and Hoboken, NJ, February 28, 2024 -- Kamada Ltd. (NASDAQ: KMDA; TASE: KMDA.TA), a commercial stage global biopharmaceutical company with a portfolio of marketed products indicated for rare and serious conditions and a leader in the specialty plasma-derived field, today announced that it will release financial results for the fourth quarter and fiscal year ended December 31, 2023, as well as provide 2024 financial guidance, prior to the open of the U.S. financial markets on Wednesday, March 6, 2024.

Kamada management will host an investment community conference call on Wednesday, March 6, 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 1-877-407-0792 (from within the U.S.), 1-809-406-247 (from Israel) or 1 201-689-8263 (International) using conference ID 13744277. The call will also be webcast live on the Internet at: https://viavid.webcasts.com/starthere.jsp?ei=1655132&tp key=6e560eef4a.

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 commercial stage global biopharmaceutical company with a portfolio of marketed products indicated for rare and serious conditions and a leader in the specialty plasma-derived field, focused on diseases of limited treatment alternatives. The Company is also advancing an innovative development pipeline targeting areas of significant unmet medical need. The Company's strategy is focused on driving profitable growth from its significant commercial catalysts as well as its manufacturing and development expertise in the plasma-derived and biopharmaceutical fields. The Company's commercial products portfolio includes six FDA approved plasma-derived biopharmaceutical products: KEDRAB®, CYTOGAM®, VARIZIG®, WINRHO SDF®, HEPAGAM B® and GLASSIA®, as well as KAMRAB®, KAMRHO (D)® and two types of equine-based anti-snake venom (ASV) products. The Company distributes its commercial products portfolio directly, and through strategic partners or third-party distributors in more than 30 countries, including the U.S., Canada, Israel, Russia, Argentina, Brazil, India, Australia and other countries in Latin America, Europe, the Middle East, and Asia. The Company leverages its expertise and presence in the Israeli market to distribute, for use in Israel, more than 25 pharmaceutical products that are supplied by international manufacturers and in addition have eleven biosimilar products in its Israeli distribution portfolio, which, subject to European Medicines Agency (EMA) and Israeli Ministry of Health approvals, are expected to be launched in Israel through 2028. The Company owns an FDA licensed plasma collection center in Beaumont, Texas, which currently specializes in the collection of hyper-immune plasma used in the manufacture of KAMRHO (D), KAMRAB and KEDRAB. In addition to the Company's commercial operation, it invests in research and development of new product candidates. The Company's leading investigational product is an inhaled AAT for the treatment of AAT deficiency, for which it is continuing to progress the InnovAATe clinical trial, a randomized, double-blind, placebo-controlled, pivotal Phase 3 trial. FIMI Opportunity Funds, the leading private equity firm in Israel, is the Company's controlling shareholder, beneficially owning approximately 38% of the outstanding ordinary shares.

CONTACTS:

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