**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 2023

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, 333-233267 and 333-265866.**

The following exhibit is attached:

| | |
|---|---|
| 99.1 | Kamada Files Annual Report for the Year Ended December 31, 2022 |

**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 15, 2023

**KAMADA LTD.**

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

EXHIBIT INDEX

| EXHIBIT NO. | DESCRIPTION |
| --- | --- |
| 99.1 | Kamada Files Annual Report for the Year Ended December 31, 2022 |

**Exhibit 99.1**

**Kamada Files Annual Report for the Year Ended December 31, 2022**

**REHOVOT, Israel, and Hoboken, NJ – March 15, 2023** -- 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 has filed its annual report on Form 20-F for the fiscal year ended December 31, 2022, with the U.S. Securities and Exchange Commission (the "SEC").

The annual report, including the Company's audited consolidated financial statements, can be accessed via the SEC's website at https://www.sec.gov/edgar.shtml, as well as under the SEC Filings section on Kamada's investor relations website at https://www.kamada.com/.

The Company will deliver a hard copy of its annual report containing its audited consolidated financial statements, free of charge, to its shareholders upon request. Requests should be directed to Kamada's Investor Relations Department at IR@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: CYTOGAM®, KEDRAB®, WINRHO SDF®, VARIZIG®, 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, 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 during recent years added eleven biosimilar products to its Israeli distribution portfolio, which, subject to the European Medicines Agency (EMA) and the Israeli Ministry of Health approvals, are expected to be launched in Israel through 2028. The Company owns an FDA registered plasma collection center in Beaumont, Texas, which currently specializes in the collection of hyper-immune plasma used in the manufacture of KAMRHO (D). 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 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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