

KAMADA LTD  
Form 6-K  
December 23, 2015

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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 December, 2015

Commission File Number 001-35948

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

7 Sapir St.  
Kiryat Weizmann Science Park  
P.O Box 4081  
Ness Ziona 74140  
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 Statement File No. 333-192720.



The following exhibit is attached:

- 99.1 News Release: Kamada's Human Rabies Immune Globulin Successfully Meets Primary Endpoint in U.S. Pivotal Phase 2/3 Clinical Trial as a Post-Exposure Treatment
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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: December 23, 2015

KAMADA LTD.

By: /s/ Gil Efron  
Gil Efron  
Deputy Chief Executive  
Officer and Chief Financial  
Officer

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EXHIBIT INDEX

EXHIBIT DESCRIPTION  
NO.

99.1 News Release: Kamada's Human Rabies Immune Globulin Successfully Meets Primary Endpoint in U.S. Pivotal Phase 2/3 Clinical Trial as a Post-Exposure Treatment

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