

**Kamada to Announce Second Quarter and Six Months Ended June 30, 2018 Financial  
Results and Host Conference Call on August 7**

**Rehovot, Israel – August 1, 2018** -- Kamada Ltd. (NASDAQ & TASE: KMDA), a plasma-derived protein therapeutics company, today announced that it will release financial results for the quarter and six months ended June 30, 2018, prior to the open of the U.S. financial markets on Tuesday, August 7, 2018.

Kamada management will host an investment community conference call on Tuesday, August 7 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 800-263-0877 (from within the U.S.), 1809 212 883 (from Israel), or 646-828-8143 (International) and entering the conference identification number: 4166824. The call will also be webcast live on the Internet on the Company's website at [www.kamada.com](http://www.kamada.com).

A replay of the call will be accessible two hours after its completion through August 21 by dialing 844-512-2921 (from within the U.S.) or 412-317-6671 (from outside the U.S.) and entering the conference identification number: 4166824. The call will also be archived for 90 days on the Company's website at [www.kamada.com](http://www.kamada.com).

***About Kamada***

Kamada Ltd. is focused on plasma-derived protein therapeutics for orphan indications, and has a commercial product portfolio and a late-stage product pipeline. The Company uses its proprietary platform technology and know-how for the extraction and purification of proteins from human plasma to produce Alpha-1 Antitrypsin (AAT) in a highly-purified, liquid form, as well as other plasma-derived Immune globulins. AAT is a protein derived from human plasma with known and newly-discovered therapeutic roles given its immunomodulatory, anti-inflammatory, tissue-protective and antimicrobial properties. The Company's flagship product is GLASSIA®, the first liquid, ready-to-use, intravenous plasma-derived AAT product approved by the U.S. Food and Drug Administration. Kamada markets GLASSIA® in the U.S. through a strategic partnership with Baxalta (now part of Shire plc) and in other countries through local distributors. In addition to GLASSIA®, Kamada has a product line of six other plasma-derived pharmaceutical products administered by injection or infusion, that are marketed through distributors in more than 15 countries, including Israel, Russia, Brazil, India and other countries in Latin America and Asia. Kamada has late-stage products in development, including an inhaled formulation of AAT for the treatment of AAT deficiency, and in addition, its intravenous AAT is in development for other indications, such as type-1 diabetes, GvHD and prevention of lung transplant rejection. Kamada's rabies immune globulin (Human) product received FDA approval for Post-Exposure Prophylaxis against rabies infection in August 2017 and was launched in the US during Q1-2018. Kamada also leverages its expertise and presence in the plasma-derived protein therapeutics market by distributing more than 10 complementary products in Israel that are manufactured by third parties.

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***Cautionary Note Regarding Forward-Looking Statements***

This release includes forward-looking statements within the meaning of Section 27A of the U.S. Securities Act of 1933, as amended, Section 21E of the U.S. Securities Exchange Act of 1934, as amended, and the safe harbor provisions of the U.S. Private Securities Litigation Reform Act of 1995. Forward-looking statements are statements that are not historical facts. Forward-looking statements are based on Kamada's current knowledge and its present beliefs and expectations regarding possible future events and are subject to risks, uncertainties and assumptions. Actual results and the timing of events could differ materially from those anticipated in these forward-looking statements as a result of several factors. The forward-looking statements made herein speak only as of the date of this announcement and Kamada undertakes no obligation to update publicly such forward-looking statements to reflect subsequent events or circumstances, except as otherwise required by law.

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