Kamada Files Annual Report for the Year Ended December 31, 2018

Rehovot, Israel, February 27, 2019 – Kamada Ltd. (NASDAQ & TASE: KMDA), a plasma-derived protein therapeutics company, today announced that it has filed its annual report on Form 20-F for the fiscal year ended December 31, 2018 with the U.S. Securities and Exchange Commission (the "SEC").

The annual report, which contains the Company's audited consolidated financial statements, can be accessed via the SEC's website at http://www.sec.gov/ as well as under the SEC Filings section on the Company's investor relations website at http://www.kamada.com/reports.php.

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 the Company's Investor Relations Department at IR@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 Takeda Pharmaceuticals Company Limited and in other counties through local distributors. Kamada's second leading product is KamRAB, a rabies immune globulin (Human) for Post-Exposure Prophylaxis against rabies infection. KamRAB is FDA approved and is being marketed in the U.S. under the brand name KEDRAB and through a strategic partnership with Kedrion S.p.A. In addition to GLASSIA and KEDRAB, Kamada has a product line of four 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 GvHD, prevention of lung transplant rejection and type-1 diabetes. Kamada also leverages its expertise and presence in the plasma-derived protein therapeutics market by distributing more than 20 complementary products in Israel that are manufactured by third parties.

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