Hikma receives FDA approval for Rifampin Injection

London, 9 March 2016 – Hikma Pharmaceuticals PLC ("Hikma") (LSE: HIK) (NASDAQ Dubai: HIK) (OTC: HKMPY), (rated Ba1 Moody's / BB+ S&P, both stable), the fast growing multinational pharmaceutical group, announces that its abbreviated new drug application ("ANDA") for Rifampin Injection USP, 600mg/vial has been approved by the U.S. Food and Drug Administration ("FDA").

Rifampin Injection is an anti-infective indicated to treat certain infections such as tuberculosis and leprosy. According to IMS Health, sales of Rifampin Injection in the US market were approximately $14.3 million for the 12 months ending January 2016.

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About Hikma

Hikma Pharmaceuticals PLC is a fast growing multinational group focused on developing, manufacturing and marketing a broad range of both branded and non-branded generic and in-licensed products. Hikma operates through three businesses: "Injectables", "Branded" and "Generics", based principally in the United States, the Middle East and North Africa ("MENA") and Europe. In 2014, Hikma achieved revenues of $1,489 million and profit attributable to shareholders of $278 million.