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KWALITY PHARMACEUTICALS LIMITED

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To
The BSE Limited
Department of Corporate Services,
P.J. Towers, Dalal Street,
Mumbai-400001

Date: March 10, 2026

Scrip Code: 539997

Announcement under Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements), Regulations, 2015 - Update on Bioequivalence (BE) Program - Successful Completion of Phase 1 by Kwality Pharmaceuticals Ltd.

Dear Sir/Madam,

Kwality Pharmaceuticals Ltd. (KPL) had initiated an extensive **Bioequivalence (BE)** program covering more than 30 molecules across key therapeutic segments, including Oncology, Cardiology, Anti-diabetics, and Anti-hypertensives. The portfolio comprises a balanced mix of both patented and off-patent complex molecules. To execute this initiative, KPL has partnered with four globally accredited Contract Research Organizations (CROs).

The program has been structured in a phased manner to ensure efficient execution and regulatory alignment. We are pleased to announce the successful completion of Phase 1, with BE studies for **Hydroxycarbamide, Nilotinib, and Liposomal Amphotericin B** completed successfully.

These dossiers are now prepared for submission across regulated, semi-regulated, and Rest of the World (ROW) markets, further strengthening KPL's global regulatory footprint and commitment to expanding access to high-quality pharmaceutical products.

The above is for your information and dissemination to the stakeholders.

Thanking You,

For Kwality Pharmaceuticals Limited

Aditya Arora

Whole Time Director & CFO

DIN: 07320410

