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When repying please quote

PPB/GMP/F/2016/124

LENANA ROAD P.O. BOX 27663-00506 NAIROBI

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

On basis of the inspection carried out in 21st - 22nd March 2016 we certify that, as at the time of inspection, Synokem Pharmaceuticals Ltd at Plot 35-36 Sector-6A, Integrated Industrial Estate (SIDCUL), Ranipur, (BHEL), Haridwar-249 403 India Complies with current PPB requirements and WHO current Good Manufacturing Practice standards for Human dosage forms, categories and activities listed below.

Dosage Form	Category and posson board pharmacy and poison board p	Activity
Oral Solid Dose Forms	General Products: Tablets & Hard gelatin Capsules Hormones: Tablets	All manufacturing activities
Oral Liquid Dosage Forms	General Products: Syrups, Suspensions & Liquids.	All manufacturing activities

This certificate remains valid until the 31st day of March 2019. The company has to apply for re-inspection three months before the expiry of this certificate if it intends to continue doing business in Kenya.

The certificate shall become invalid if;

- 1. The activities and/or categories certified herewith are changed.
- 2. The site is no longer considered to be in compliance with WHO cGMP.
- 3. The manufacturing site is changed.

The responsibility for the quality of the individual batches of the Pharmaceutical Products* manufactured in this site lies with the manufacturer and the PPB reserves the right to inspect the manufacturing site at any time it deems necessary within the validity of this certificate.

REGISTRAR

PHARMACY AND POISONS BOARD MINISTRY OF HEALTH
P. O. Box 27663 - 00506, NAIROBI

13th June 2016

REGISTRAR

STAMP and DATE

Pharmaceutical products: Any medicine intended for human use or veterinary product administered to food-producing animals, presented in its finished dosage form or as a starting material for use in such a dosage form, that is subject to control by pharmaceutical legislation in Kenya