Food and Drugs Authority

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FDA/DRID/DED/PIU/17/0188

30th June 2017

The Managing Director
Synokem Pharmaceuticals Ltd
Synokem House 14/486
Sunder Vihar
Paschim Vihn
New Delhi 110097
India

Dear Sir/Madam,

GOOD MANUFACTURING PRACTICE (GMP) CERTIFICATION

Pursuant to the provisions of **Section 131 of the Public Health Act, 2012, Act 851 of the Republic of Ghana,** this is to inform you that the Food and Drugs Authority, Ghana, after conducting Good Manufacturing Practices (GMP) audit at your facility sited at **Haridwar, India**, during the period of 25th- 26th May, 2017, has determined that the operations of the facility meet acceptable GMP standard. The facility is therefore licensed to manufacture dosage forms as covered in the certification for the Ghanaian market.

The Food and Drugs Authority hereby issues you with a GMP certificate with number: **FDA/GMP/004/06/17**, valid up to **April, 2022** and subject to renewal.

You are directed to act in accordance with the under listed terms and conditions of the certification.

1. The certificate should be renewed after its expiry.
2. Only product(s) manufactured in the audited facility can be distributed on the Ghanaian market.
3. Distribution of the product(s) covered by the certification in the Ghanaian market ceases after the expiration of the certification.

Yours faithfully,

DELESE A. A. DARKO (MRS)
AG. CHIEF EXECUTIVE OFFICER