OFFICE OF THE DRUGS CONTROLLING & LICENSING AUTHORITY Directorate General of Medical Health & Family Welfare, Sahastradhara Road, Dehradun (Uttarakhand)

F.no.17P/1/37/2005/7691

Certificate of Good Manufacturing Practices

Date: 64

June 2016

Certificate no.:17P/1/37/2005/1320

On the basis of the inspection carried out on 02.06.2016 we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in Table 1.

1. Name & Address of site:

M/s Synokem Pharmaceuticals Ltd. Plot No. 35-36, Sector-6A, IIE, Ranipur, Haridwar, Uttarakhand.

2. Manufacturer's license number:

Form 25- 19/UA/2005 Form 28- 17/UA/SC/P-2005

3. Table 1:

Dosage form(s)	Category(ies)	Activity(ies)
Tablets	Non betalactum,	Manufacturing
Capsules (Hard Gelatin)	Non betalactum,	Manufacturing
Oral Liquid	Non beta lactum,	Manufacturing
Tablets(Hard Gelatin) /Capsule	Hormone	Manufacturing

The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

This certificate remains valid until 05.05.2018. It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

The firm is following Good Manufacturing Practices as per World Health Organization(WHO)TRS Guide Lines, in the Manufacturing & testing of the said categories of Products and Items in respect of which the Certificates of Pharmaceuticals products have been issued.

Address of certifying Authority:

Directorate General of Medical Health & Family Welfare, Sahastradhara Road, Dehradun (Uttarakhand) INDIA.

Name & function of responsible person:

Shri Tajber Singh

Drugs Licensing & Controlling Authority

Uttarakhand.

Email: drugcontroluk@gmail.com

Tel.no. NA

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Garhwal Mandal (Uttarakhand)