OFFICE OF THE DRUG CONTROLLING & LICENSING AUTHORITY

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

F.No. 17P/1/48/2010/13642

Dated: 07 Sept 2018

Certificate of Good Manufacturing Practices

On the basis of the Joint Inspection carried out on 24.08.2018 & 25-08-2018 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 Verve Human Care Laboratories Plot No. 15-A, Pharmacity, Selaqui, Dehradun -248011, Uttarakhand INDIA

2. Manufacturer's license number:-

Form 25-33/UA/2010 Form 28-34/UA/SC/P-2010 Form 28B-29/UA/X/SC/P-2014 Form 25F-1/UA/X/2012

3. Table 1:

Dosage form (s)	Category(s)	Activity (ies)
Tablets	Non Betalactum	Manufacturing
Injectable	Non Betalactum	Manufacturing

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

This certificate remains valid until 28.08.2021. It becomes invalid if the activities and / or categories certified herewith are changed or if the site is no longer considered to be in completion with GMP.

The firm is following Good Manufacturing Practices as per World Health Organization (WHO) TRS Guidelines, in the manufacturing & testing of the said categories of Products and Items . respect of which the Certificate 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 Drug Controlling & Licensing (Mfg.) (Uttarakhand)

Email: drugcontroluk@gmail.com

Tel. no. NA

Fax No. 0135260874

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(Tajber Singh)

Drug Controlling & Livensing (Mfg.)

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