

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, Pharmacy, 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 in 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



(Tajber Singh)

Drug Controlling & Licensing (Mfg.)

Drug Controlling & Licensing Authority (Mfg.)
(Uttarakhand)

**OFFICE OF THE DRUG CONTROLLING & LICENSING AUTHORITY,
DIRECTORATE GENERAL OF MEDICAL HEALTH & FAMILY WELFARE,
SAHASTRADHARA ROAD, DEHRADUN (UTTARAKHAND)**

F.No. 17P/1/48/2010/ 8146

Dated: 08-05-2019

G.M.P. CERTIFICATE

This is to certify that M/s Verve Human Care Laboratories, Plot No. 15-A, Pharmacy, Selaqui, Distt. Dehradun Uttarakhand (India) has been licensed under Drug & Cosmetics Act 1940 & Rules there under. They are holding valid Drug manufacturing license no. 33/UA/2010 on form 25 & 34/UA/SC/P-2010 on form 28 valid up to 24-03-2020 & license No. 1/UA/X/2012 on form 26 F valid up to 07-08-2022 & 29/UA/X/SC/P-2014 on form 28B valid up to 04-04-2024 manufacture for sale of Drugs.

In view of report dt. 07-05-2019 of Inspector of Drugs, H.Q, it is certified that M/s Verve Human Care Laboratories, Plot No. 15-A, Pharmacy, Selaqui, Distt. Dehradun Uttarakhand (India) conforms to requirements of Rule 71, 74 & 76,78, and Good Manufacturing Practices as laid down under revised Schedule "M" of the Drugs & Cosmetics Rule 1945.

This certificate is issued to the firm on their request for submission to Govt. departments/Institutions/Overseas Authority.

This certificate is valid for a period up to Three year from the date of issue.




(Tajber Singh)
Drug Controlling &
Licensing Authority (Mfg.)
Uttarakhand.

(Tajber Singh)
Drug Controlling & Licensing Authority (Mfg.)
(Uttarakhand)

**OFFICE OF THE DRUG CONTROLLING & LICENSING AUTHORITY,
DIRECTORATE GENERAL OF MEDICAL HEALTH & FAMILY WELFARE,
SAHASTRADHARA ROAD, DEHRADUN (UTTARAKHAND)**

F.No. 17P/1/48/2010/ 8148

Dated: 08-05-2019

G.L.P. CERTIFICATE

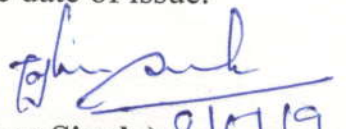
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In view of report dt. 07-05-2019 of Inspector of Drugs H.Q, it is certified that M/s Verve Human Care Laboratories, Plot No. 15-A, Pharmacy, Selaqui, Distt. Dehradun Uttarakhand (India) confirms to G.L.P. requirements of Good Laboratory Practices as laid down under Schedule "L-1" (150-E) of the Drugs & Cosmetics Rule 1945.

This certificate is issued to the firm on their request for submission to Govt. departments / Institutions / Overseas Authority.

This certificate is valid for a period up to three Year from the date of issue.




(Tajber Singh) 8/5/19
Drug Controlling &
Licensing Authority (Mfg.)
Uttarakhand.
Drug Controlling & Licensing Authority (Mfg.)
(Uttarakhand)

From,

Drugs Controller,
Director General Medical Health & Family Welfare
Sahastradhara Road, Dehradun,
Uttarakhand.

To,

M/s Verve Human Care Laboratories,
15-A, PharmaCity, Selaqui,
Dehradun, Uttarakhand,

No. 26/1/Drug/44/2019/ 15217

Dated 24/08/2021

Sub- Extension of Validity of COPP granted under WHO-GMP & COPP
certification scheme & WHO-GMP Certificate.

Sir,

In reference to your application dated 16-08-2021 regarding above noted subjects it is to inform you that the validity of COPP granted under WHO-GMP certification scheme & WHO-GMP Certificate is further extended from 28-08-2021 to 27-02-2022 for Six months.



(Tajber Singh)
Drugs Controller
Uttarakhand

No. 26/1/Drug/44/2019/

Of dated

Copy to- Dy. Drug Controller (North Zone) CDSCO, Govt. of India, Kamla Nehru Nagar, Hapur Road, Gaziabad, UP.

(Tajber Singh)
Drugs Controller
Uttarakhand