



Office of The Commissioner,  
Food & Drugs Administration M.S.  
Bandra - Kurla Complex,  
Bandra (E),  
Mumbai - 400 051  
Date :

## CERTIFICATE OF GOOD MANUFACTURING PRACTICES

This Certificate conforms to the format recommended by the World Health Organization.  
(General instructions and explanatory notes attached).

Certificate No.: **NEW-WHO-GMP/CERT/NKD/91974/2020/11/31704**

On the basis of the inspection carried out on 20/01/2020, 21/01/2020 & 21/04/2020, 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.

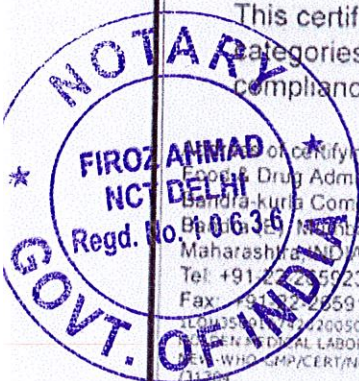
- Name of the Firm : **HOLDEN MEDICAL LABORATORIES PVT. LTD.**  
Address : **PLOT NO. C-35, C-36 AND C-37, M.I.D.C. MALEGAON, SINNAR, NASHIK 422113 MAHARASHTRA STATE, INDIA**
- Licence No. : **NKD34 In Form 25, NKD19 In Form 28**

Table 1

Sr.No.	Dosage Form(s)	Categor(ies)	Activity(ies)
1	Capsules	General ( Other than Cephalosporins, Penicillin, Cytotoxic, Hormones )	Production, Filling, Packing, labelling, Quality Control, Quality Assurance
2	Tablets	General ( Other than Cephalosporins, Penicillin, Cytotoxic, Hormones )	Production, Filling, Packing, labelling, Quality Control, Quality Assurance

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 May 2023. 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.



Authorizing authority :  
Food & Drug Administration, M.S.  
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HOLDEN MEDICAL LABORATORIES PVT. LTD.  
NEW-WHO-GMP/CERT/NKD/91974/2020/11/31704

Name of the Authorised person : J.B. MANTRI

**ATTESTED**

Signature :

Stamp and Date :

Joint Commissioner (HQ) & Controlling Authority  
Food & Drug Administration, M.S.  
Bandra (E), Mumbai  
Maharashtra State, India  
Date: 06 May 2020

**NOTARY PUBLIC  
DELHI (INDIA)**

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**ATTESTED**



Authorized Signatory  
Millennium India International  
Chamber of Commerce  
Industry & Agriculture  
New Delhi, (INDIA)

**GAURAV RAVISH**  
Addl. Director

## Explanatory notes

1. This certificate which is in the format recommended by WHO, certifies the status of the site listed in point 1 of the certificate.
2. The certification number should be traceable within the regulatory authority issuing the certificate.
3. Where the regulatory authority issues a licence for the site, this number should be specified record "not applicable" in cases where there is no legal framework for the issuing of a licence.
4. Table 1  
List the dosage forms, starting materials, categories and activities. Examples are given below.

### Example - 1

Pharmaceutical Product (s)1	Category (ies)	Activity (ies)
Dosage form (s)		
Tablets	Cytotoxic	Packaging
	Hormone	Production, Packaging, Quality control.
Injectables	Penicillin	Repackaging & Labelling.
	Cefalosporin	Aseptic preparation, Packaging, Labelling.

### Example - 2.

Pharmaceutical Product (s)1	Category (ies)	Activity (ies)
Starting material (s)2		
Paracetamol	Analgesic	Synthesis, Purification

Use, whenever available. International nonproprietary names.

5. The certificate remains valid until activities and/or categories certified compliance with GMP.
6. The requirements for good practices the certificate are those included in guidelines and related materials, 1999, World Health Organization, Geneva.



GAURAV VARMA  
Aqbl. Director