



Office of The Commissioner,  
Food & Drugs Administration M.S.  
Bandra – Kurla Complex,  
Bandra (E),  
Mumbai – 400 051  
Date : 5/5/2014

## 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.: **WHO-GMP/CERT/PD/15126/2014/11/6621**

On the basis of the inspection carried out on **16/09/2011** and **17/09/2011**, 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 of the Firm : **STERLING HEALTHCARE LTD.**  
Address : **GAT NO. 396, VILLAGE URSE, NEAR  
TALEGAON TOLL PLAZ MUMBAI- PUNE  
EXPRESSWAY, TAL. MAVAL, PUNE 410506  
MAHARASHTRA STATE, INDIA**
2. Licence No. : **PD195 In Form 25,  
PD116 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.

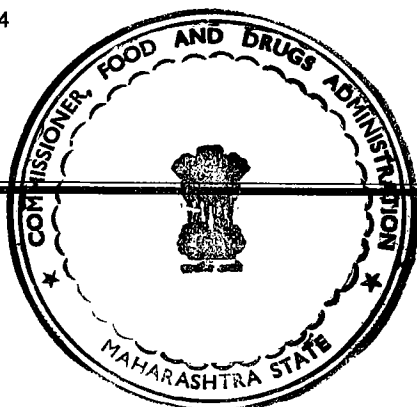
The Certificate was valid upto 06 Feb 2014. The validity of this certificate has been extended for the period of six months and now the validity of this Certificate remains 06 Aug 2014. It becomes invalid if the activities and / or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Address of certifying authority :  
Food & Drug Administration, M.S.  
Bandra-kurla Complex,  
Bandra (E), Mumbai – 400 051.  
Maharashtra, INDIA.  
Tel: +91-22-26592363/64  
Fax: +91-22-26591959  
1ETS1851512620140425H42

Name of the Authorised person : **O.S. SADHWANI**

Signature:

Stamp and Date : **Joint Commissioner (HQ) & Controlling  
Authority  
Food & Drug Administration, M.S.  
Bandra (E), Mumbai.  
Maharashtra State, India  
Date: 25 Apr 2014**



25 APR 2014

## 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) <sub>1</sub>	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) <sub>1</sub>	Category (ies)	Activity (ies)
Starting material (s) <sub>2</sub>		
Paracetamol	Analgesic	Synthesis, Purification, Packing, Labelling.

Use, whenever available. International Nonproprietary Names (INNs) or otherwise national nonproprietary names.

5. The certificate remains valid until the specified date. The certificate becomes invalid if the activities and/or categories certified are changed or if the site is no longer considered to be in compliance with GMP.
6. The requirements for good practices the manufacture and quality control of drugs referred to in the certificate are those included in Quality Assurance in Pharmaceuticals: a compendium of guidelines and related materials. Good manufacturing practices and inspection. Volume 2, 1999. World Health Organization, Geneva and subsequent updates.

