



REPUBLIC OF KENYA
 MINISTRY OF HEALTH
 PHARMACY AND POISONS BOARD

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PHARMACY AND POISONS BOARD HOUSE
 LENANA ROAD
 P.O. Box 27663-00506
 NAIROBI

When replying please quote
PPB/GMP/F/2014/136

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

On basis of the inspection carried out on **6TH & 7TH May 2014** we certify that, as at the time of inspection, **Bharat Parenterals Limited** at 144 & 146, Vill. Haripura, Jarod- Samlaya road, Ta Savli, Dist. Vadodara - 391 520, Gujarat, India, Complies with current PPB requirements and WHO current Good Manufacturing Practice standards for Human dosage forms, categories and activities listed below.

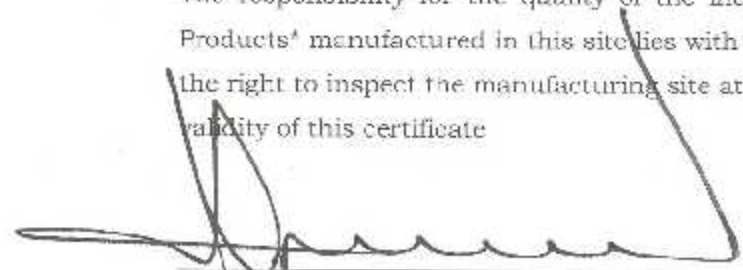
Dosage Form	Category	Activity
Oral Solid Dose Forms	General Products: Tablets & Hard Gelatin Capsules	All manufacturing activities
Sterile Preparations	General Products: SVP (Liquid injections) & Eye drops	All manufacturing activities

This certificate remains valid until the **31st day of May 2017**. The company has to apply for re-inspection **three months** before the expiry of this certificate if it intends to continue doing business in Kenya.

The certificate shall become invalid if;

1. The activities and/or categories certified herewith are changed.
2. The site is no longer considered to be in compliance with WHO cGMP.
3. The manufacturing site is changed.

The responsibility for the quality of the individual batches of the Pharmaceutical Products* manufactured in this site lies with the manufacturer and the PPB reserves the right to inspect the manufacturing site at any time it deems necessary within the validity of this certificate



REGISTRAR

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 MINISTRY OF HEALTH
 P. O. Box 27663 - 00506, NAIROBI
 11th August 2014

STAMP and DATE

*Pharmaceutical products: Any medicine intended for human use or veterinary product administered to food-producing animals, presented in its finished dosage form or as a starting material for use in such a dosage form, that is subject to control by pharmaceutical legislation in Kenya