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/2014/125

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

On basis of the inspection carried out in **28th – 29th October 2013** we certify that, as at the time of inspection, **Cure Medicines (India) Pvt. Ltd** at Plot NO. C-12 / 13, MIDC, Bhosari, Pune – 411 026, 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
Oral Liquid Dose Forms	**General Products:** Liquids, Syrups & Suspensions	All manufacturing activities

This certificate remains valid until the **31st day** of **October 2016.** 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
PHARMACY AND POISONS BOARD
MINISTRY OF HEALTH
P. O. Box 27663 - 00506, NAIROBI

19th June 2014

REGISTRAR ***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