GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

L.Dis.No.2923/BM/2013 **Dated: 25-09-2013**

From:

P.Nagabhushanam,M.Pharm, B.L, MBA,
Director & Licensing Authority
O/o.the Director General,
Drugs and Copyrights.
Drugs Control Administration,
Vengalraonagar, Hyderabad – 500 038.

To

M/s.Sreepathi Pharmaceuticals Ltd,
Plot No.163,
Phase-V, I.D.A,.
Jeedimetla,
Hyderabad..
Andhra Pradesh, India.

Sirs,

Sub:- Drugs and Cosmetics Act, 1940 and Rules made thereunder – Issue of World Health Organisation Good Manufacturing Practice Certificate – Regarding.

Ref:- 1.Your application dt.30-01-2013
2. Joint Inspection team report dt.02-07-2013

With reference to your application cited, I forward herewith **WORLD HEALTH ORGANISATION GOOD MANUFACTURING PRACTICE** Certificate for the products mentioned in the Joint Inspection Report of the Officers of Drugs Control Administration, Andhra Pradesh.

This Certificate is valid for a period of Two years from the date of issue This certificate is meant for Export of drugs only.

DIRECTORATE OF DRUGS CONTROL ADMINISTRATION GOVERNMENT OF ANDHRA PRADESH HYDERABAD

Yours faithfully,

25-9-13

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

Office of the Director General
Drugs and Copyrights
Drugs Control Administration
Vengalrao Nagar, Hyderabad.

L.Dis.No.2923/BM/2013 Dated: 25-09-2013

LIST OF PRODUCTS APPROVED UNDER WHO GMP CERTIFICATION SCHEME FOR EXPORT PURPOSE

1. CIPROFLOXACIN HYDROCHLORIDE USP /Ph.Eur/BP
2. CIPROFLOXACIN USP
3. ENROFLOXACIN

Manufacturer	:	M/s. Sreepathi Pharmaceuticals Ltd, Plot No.163, Phase-V, I.D.A,. Jeedimetla, Hyderabad, Andhra Pradesh, Indai...
When applicable	:	Placing the product on the market as detailed.

It is certified that the above products have been authorized to be placed on the market for use in the Country.

Drug Licence No.	:	76/RR/AP/95/B/R dt.09-06-1986 Valid upto 31-12-2016 in Form No. 25

It is also certified that (a) the manufacturing plant in which the products are produced is subject to inspection at suitable intervals.

The Unit M/s. Sreepathi Pharmaceuticals Ltd, Plot No.163, Phase-V, I.D.A,Jeedimetla, Hyderabad..A.P. was jointly inspected by Sri.G.Dharmadata, Deputy Director, Drugs Control Administration, Hyderabad Sri.K.Raja Bhanu, Assistant Director, Drugs Control Administration, Hyderabad and Sri. E.Sambasiva Rao, Drugs Inspector, Jeddimetla Zone, Drugs Control Administration, Hyderabad, Andhra Pradesh, Hyderabad on 02-07-2013.

The manufacturer conforms to requirement for Good Manufacturing Practices in the manufacture and quality control (As recommended by the World Health Organization) in respect of products mentioned above (Three numbers) for Export in the International market.

This Certificate is valid for a period of Two years from the date of issue This certificate is meant for Export of drugs only.

DIRECTOR & LICENSING AUTHORITY,
DRUGS CONTROL ADMINISTRATION.

25-9-13

DIRECTORATE OF DRUGS CONTROL ADMINISTRATION GOVERNMENT OF ANDHRA PRADESH HYDERABAD