DRUGS CONTROL ADMINISTRATION
Government of Telangana

L. Dis. No.7179/E(J)/TS/2017 Dated: 05-09-2018

To
M/s.A.R.Life Sciences Private Limited
Plot No.33, Sri Venkateshwara Co-op Indl. Estate
Jeedimetla, Hyderabad-500055
Telangana State, India.
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Sir,

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder – Issue of World Health Organisation G.M.P. Certificate – Regarding.

Ref: 1. Your letter dated: 10.05.2017
2. Joint Inspection report dt:18.12.2017 & 19.12.2017.

-x-x-x-x-

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, Telangana State and CDSCO, Zonal Office, Hyderabad vide reference 2nd cited.

This Certificate is valid for a period of Three years from the date of issue.

Deputy Director Drugs Control Administration Government Of Telangana HYDERABAD

Yours faithfully,
05/09/18
Dr.B.VENKATESWARLU
Joint Director & Licensing Authority(I/c)

DRUGS CONTROL ADMINISTRATION
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L.Dis.No.7179/E(J)/TS/2017 Dated: 05-09-2018

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

1. SPARFLOXACIN
2. CLOPIDOGREL BISULPHATE USP
   (Clopidogrel Hydrogen Sulfate)

Manufacturer : M/s.A.R.Life Sciences Private Limited
Plot No.33, Sri Venkateshwara Co-op Indl. Estate, Jeedimetla, Hyderabad-500055
Telangana State, India.

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When applicable detailed : Placing the product on the market as above.

It is certified that the above products had been authorized to be placed on the market for use in the Country and exporting countries.

Drug Licence No. : 48/RR/AP/2003/B/CC, dated:20.09.2003 under Form – 25 valid upto 05.02.2022

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

The Unit M/s.A.R.Life Sciences Private Limited, Plot No.33, Sri Venkateshwara Co-op Indl.Estate, Jeedimetla, Hyderabad-500055, Telangana State, India was inspected jointly by Mrs.M.Shakuntala, Drugs Inspector, CDSCO, Hyderabad and Mrs.M.Sree Bindu, Drugs Inspector, Jeedimetla(Mfg) FAC, DCA, Hyderabad on 18.12.2017 & 19.12.2017.

(b) The manufacturer conforms to requirements for Good Manufacturing Practices in the manufacturer and Quality Control (As recommended by the World Health Organisation) in respect of 02 (Two) products to be sold or distributed with in the Country or origin (or to be exported).

This Certificate is valid for Three years from the date of issue.

Deputy Director Drugs Control Administration Government Of Telangana HYDERABAD

05.09.18

Dr.B.VENKATESWARLU
Joint Director & Licensing Authority(I/c)