GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

Office of the Director General,
Drugs and Copyrights,
Drugs Control Administration,
Chuttugunta, Guntur –522004.

L.Dis.No.324/Stores/2020 **Dated:25-02-2020**

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

S.No.	Product Name	Category
1.	GLUCOSAMINE HYDROCHLORIDE	USP
2.	GLUCOSAMINE SULFATE POTASSIUM CHLORIDE	USP
3.	GLUCOSAMINE SULFATE SODIUM CHLORIDE	USP
4.	GLUCOSAMINE HYDROCHLORIDE	Ph.Eur
5.	GLUCOSAMINE SULFATE POTASSIUM CHLORIDE	Ph.Eur
6.	GLUCOSAMINE SULFATE SODIUM CHLORIDE	Ph.Eur

Name of the Manufacturer : M/s.Andhra Medi Pharma India Pvt, Ltd, Sy.No.263, Veeravalli (V), Bapulapadu (M), Krishna District-521110, Andhra Pradesh, India.

When applicable : Placing the product on the market as detailed

Drug Licence No. : **10/KR/AP/2012/B/R dated 28-02-2012 in Form 25 valid up to 27-02-2022.**

It is also certified that

The manufacturing plant in which the products are produced is subject to inspection at suitable intervals.

The unit M/s. Andhra Medi Pharma India Pvt, Ltd, Sy.No.263, Veeravalli (V), Bapulapadu (M), Krishna District-521110, Andhra Pradesh, India. was inspected jointly by Mr.R. Dharmaraj, Assistant Drugs Controller (I), CDSCO, Hyderabad and Mr.Dr.J.Balu, MD, Drugs Inspector, Giduvada, Drugs Control Administration, on 12-12-2019

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 product mentioned above (Six numbers) for Export in the International market.

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

DIRECTORATE OF DRUGS CONTROL ADMINISTRATION GOVERNMENT OF ANDHRA PRADESH GUNTUR

25/2/2020

DIRECTOR
DRUGS CONTROL ADMINISTRATION

GOVERNMENT OF ANDHRA PRADESH
DRUGS CONTROL ADMINISTRATION

L.Dis.No.324/Stores/2020 **Dated:25-02-2020**

From:

M.B.R.Prasad, M,Pharm,
Director,
O/o.The Director General,
Drugs and Copyrights,
Drugs Control Administration,
Chuttugunta, Guntur – 522004.

To

M/s.Andhra Medi Pharma India Pvt.Ltd,
Sy.No. 263,
Veeravalli (V),
Bapulapadu Mandal,
Krishna District,
Andhra Pradesh, India.

Sirs,

Sub:- Drugs and Cosmetics Act, 1940 and rules made there under – Issue of World Health Organization G.M.P. Certificate – Reg.

Ref: 1.Your application dt.02-11-2019.
2. Joint Inspection Report dt. 12-12-2019
3. Ref: 5-6 (413 A3/2019/5918, dated; 29.11.2019 of the Deputy Drugs Controller (I), CDSCO, Hyderabad.
4. Lr.Rc.No. 327/DD/DCA/Eluru Region/2019, dated: 20-12-2019 of the Deputy Director, Eluru
5. Lr.Rc.No.JB/GDV/AMP/WHOGMP/2019 dt: 05-02-2020 of the Drugs Inspector, Gudivada

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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 and CDSCO, Hyderabad.

This Certificate is valid for a period of Three years from the date of issue and this certificate is meant for Export of Drugs only.

DIRECTORATE OF DRUGS CONTROL ADMINISTRATION GOVERNMENT OF ANDHRA PRADESH GUNTUR

Yours faithfully,

25/2/2020

DIRECTOR
DRUGS CONTROL ADMINISTRATION