衛 生 福 利 部
MINISTRY OF HEALTH AND WELFARE

CERTIFICATE OF GOOD MANUFACTURING PRACTICE

Issue Date: May 08, 2020

Issued following an inspection in accordance with Article 57 of the Pharmaceutical Affairs Law and relevant Regulations of the Republic of China (Taiwan).

The competent authority of the Republic of China confirms the following:
The manufacturer: Egyptian International Pharmaceutical Industries Co.
Site address: Tenth of Ramadan City Industrial area B1, Egypt
Approval number: 111-0583(PMF-I0367)

is the manufacturer of medicinal products for human use that has been inspected with the following pharmaceutical dosage forms:
-Sterile products: Sterile Semi-solid dosage forms: sterile semi-solid dosage forms (terminal sterilization) (not include specifically toxic and hazardous substances).

From the knowledge gained during GMP inspection performed on November 26-30, 2017 and dossiers assessment concluded on April 21, 2020, it is considered that the manufacturer complies with the Pharmaceutical Inspection Convention/Co-operation Scheme Guide to Good Manufacturing Practice (PIC/S GMP) for medicinal products.

This certificate is valid until April 22, 2022.
This certificate may be revoked at anytime as warranted.

Signed by

Shou-Mei Wu

Shou-Mei Wu, Ph.D.
Director-General
Food and Drug Administration
（http://www.fda.gov.tw/TC/index.aspx）

Under the delegated authority of
Shih-Chung Chen, D.D.S.
Minister
Ministry of Health and Welfare
Republic of China (Taiwan)

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