Statement of Compliance

THIS IS TO CERTIFY THAT THE QUALITY MANAGEMENT SYSTEM OF

Maya Biotech Pvt. Ltd.

Site: Vill. Kondi. P.O. Thana, Baddi, Distt. Solan (H.P.) 173205

HAS BEEN FOUND TO CONFORM

Good Manufacturing Practices

Forty Sixth report of the WHO Committee on specifications for pharmaceutical preparations (WHO Technical Report series no. 970)

Scope: Manufacturing of Small Volume Parenterals (Liquid, Dry Injectables) General & Cephalosporin, Sterile Eye/Ear drops, Nasal drops, Nasal Sprays, Sublingual Oral Sprays,Contrast Media, Metered Dose Inhaler and Lyophillized Powder Injection

The product description, documents, assessment procedure and evaluations of the examination are presented in Site Master File retained at manufacturers place

First Date of Issue
01/04/2022

Date of Issue
01/04/2022

Certification Period
3 Years

Expiry Date
01/04/2025

Certificate Number
IND/02/5092022

Technical Manager

This Certificate is issued under the following conditions:

1. Product Liability rests with the Manufacturer, In case of damages caused by defective/non compliant products, the liability rests with the manufacturer.
2. The Certificate validity is conditioned by the positive results of the surveillance audits.
3. The Certificate remains valid until the manufacturing conditions, the quality systems or relevant legislation are changed.
4. Any significant changes in the design or process used to manufacture the product or revision to the guidelines or standard/s referred above may render this certificate invalid.
5. Before the above statement is put into use, the manufacturer has to ensure that all relevant guidelines and local legislations are complied with.
6. Product/s or product packaging shall not bear WHO Logo.