Medicines and Healthcare products Regulatory Agency

CERTIFICATE NUMBER : UK API 49859 Insp GMP 49859/122060-0004

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER*(1),(2)*

Part 1

Issued following an inspection in accordance with :
Regulation 331A of The Human Medicines Regulations 2012 (SI 2012/1916)

The competent authority of United Kingdom confirms the following :

The Manufacturer : PFIZER HEALTHCARE INDIA PRIVATE LIMITED

Site address : PFIZER HEALTHCARE INDIA PRIVATE LIMITED, L-8 & L-9, MIDC INDUSTRIAL AREA, WALUJ, AURANGABAD, IN-431136, INDIA

Is an active substance manufacturer that has been inspected in accordance with Regulation 327 of The Human Medicines Regulations 2012 (SI 2012/1916).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 27/06/2018 , it is considered that it complies with

- The principles of GMP for active substances referred to in Regulation B17 and C17 of the Human Medicines Regulations 2012 (SI 2012/1916)

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field. This certificate is valid only when presented with all pages and both Parts 1 and 2. The authenticity of this certificate may be verified in MHRA-GMDP. If it does not appear, please contact the issuing authority.

- *(1) Guidance on the interpretation of this template can be found in the Help menu of MHRA-GMDP database.*
- *(2) These requirements fulfil the GMP recommendations of WHO.*

Part 2

Human Medicinal Products

Manufacture of active substance. Names of substances subject to inspection :

- [1000002970] MEROPENEM
- [1000004383] CILASTATIN
- [1000008812] IMIPENEM

3. MANUFACTURING OPERATIONS - ACTIVE SUBSTANCES

MEROPENEM

3.5	General Finishing Steps
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
	3.6.2 Microbiological testing (excluding sterility testing)
	3.6.3 Microbiological testing (including sterility testing)

CILASTATIN

3.5	General Finishing Steps
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
	3.6.2 Microbiological testing (excluding sterility testing)
	3.6.3 Microbiological testing (including sterility testing)

IMIPENEM

3.5	General Finishing Steps
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
	3.6.2 Microbiological testing (excluding sterility testing)
	3.6.3 Microbiological testing (including sterility testing)

Any restrictions related to the scope of this certificate:

Building	Room	Line/equipment	QC Testing	Products
Areas of the facility for the manufacture of sterile APIs (Block 32) and supporting QC & storage areas were included in the scope of this inspection.				Sterile APIs for EU supply: Meropenem and Imipenem/Cilastatin were covered by this inspection.

22/10/2018	Name and signature of the authorised person of the Competent Authority of United Kingdom Confidential Medicines and Healthcare products Regulatory Agency Tel : Confidential