Medicines and Healthcare products Regulatory Agency

CERTIFICATE NUMBER : UK API 27830 Insp GMP 27830/119738-0012

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 : EUROFINS SELCIA LIMITED

Site address : EUROFINS SELCIA LIMITED, FYFIELD BUSINESS AND RESEARCH PARK, FYFIELD ROAD, ONGAR, CM5 0GS, UNITED KINGDOM

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 22/11/2021 , 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 :

- [1000018046] ACTIVE SUBSTANCES FOR CLINICAL TRIALS

3. MANUFACTURING OPERATIONS - ACTIVE SUBSTANCES

ACTIVE SUBSTANCES FOR CLINICAL TRIALS

3.1 Manufacture of Active Substance by Chemical Synthesis

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	3.1.1 Manufacture Of Active Substance Intermediates
	3.1.2 Manufacture Of Crude Active Substance
	3.1.3 Salt Formation/Purification steps (eg. Crystallisation) Small scale crystallisation
	3.1.4 Other Carbon-14 labelling of APIs for Clinical Trials
3.5	General Finishing Steps
	3.5.2 Primary Packaging
3.6	Quality Control Testing
	3.6.1 Physical / Chemical testing
4	Other Activities
	Carbon-14 labelling of APIs for Clinical Trials

Any restrictions related to the scope of this certificate:

Building	Room	Line/equipment	QC Testing	Products
	Manufacture in GMP suite (Rooms 131/132, 133, 134), QC Testing Room 013			Preparation and 14C labelling of APIs for Clinical trials

17/01/2022	Name and signature of the authorised person of the Competent Authority of United Kingdom Confidential Medicines and Healthcare products Regulatory Agency Tel : Confidential

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