Medicines & Healthcare products Regulatory Agency

MHRA

MHRA
10 South Colonnade
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E14 4PU
United Kingdom

gov.uk/mhra

RESTRICTED – COMMERCIAL
Mr Graham Jones
FISHER CLINICAL SERVICES UK LIMITED
LANGHURSTWOOD ROAD
HORSHAM
RH12 4QD
UNITED KINGDOM

Medicines & Healthcare products Regulatory Agency

MHRA

Certificate No: UK MIA(IMP) 18693 Insp IMP 18693/11204-0017

Medicines and Healthcare products Regulatory Agency

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 (SI 2004/1031)

The competent authority of the United Kingdom confirms the following:

The manufacturer	FISHER CLINICAL SERVICES UK LIMITED
Site address	LANGHURSTWOOD ROAD HORSHAM RH12 4QD UNITED KINGDOM

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. MIA(IMP) 18693 in accordance with Art. 13 of Directive 2001/20/EC transposed in the following national legislation: The Medicines for Human Use (Clinical Trials) Regulations 2004 (SI 2004/1031).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 26/10/2021, it is considered that it complies with the principles and guidelines of Good Manufacturing Practice laid down in Regulation B17 of the Human Medicines Regulations 2012 (as amended).

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 only valid when presented with all pages and both parts 1 and 2.

The authenticity of this certificate may be verified in MHRA-GMDP database. If it does not appear please contact the issuing authority.

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Part 2

Human Investigational Medicinal Products for phase I, II, III clinical trials

1. MANUFACTURING OPERATIONS

1.1 Sterile products

1.1.3 Batch Certification

1.2 Non-sterile products

1.2.1 Non-sterile products (processing operations for the following dosage forms)

1.2.1.1 Capsules, hard shell

1.2.2 Batch Certification

1.3 Biological medicinal products

1.3.2 Batch certification (list of product types)

1.3.2.1 Blood products
1.3.2.2 Immunological products
1.3.2.3 Cell therapy products
1.3.2.4 Gene therapy products
1.3.2.5 Biotechnology products
1.3.2.6 Human or animal extracted products
1.3.2.8 Other biological medicinal products
IMPs

1.4 Other products or manufacturing activity

1.4.1 Manufacture of

1.4.1.1 Herbal products
1.4.1.2 Homeopathic products

1.5 Packaging

1.5.1 Primary packaging

1.5.1.1 Capsules, hard shell
1.5.1.2 Capsules, soft shell
1.5.1.8 Other solid dosage forms
1.5.1.13 Tablets

1.5.2 Secondary packaging

1.6 Quality control testing

Not Authorised

2. IMPORTATION OF MEDICINAL PRODUCTS

2.1 Quality control testing of imported medicinal products

Not Authorised

2.2 Batch certification of imported medicinal products

2.2.1 Sterile Products

2.2.1.1 Aseptically prepared products
2.2.1.2 Terminally sterilised products
2.2.2 Non-sterile products
2.2.3 Biological medicinal products

2.2.3.1 Blood products
2.2.3.2 Immunological products
2.2.3.3 Cell therapy products
2.2.3.4 Gene therapy products
2.2.3.5 Biotechnology products
2.2.3.6 Human or animal extracted products

2.3 Other importation activities

2.3.1 Site of Physical Importation
2.3.4 Other

Importation of QP certified IMPs from a country on the 'approved country for import list'

3. MANUFACTURING OPERATIONS

3.1 Manufacture of Active Substance by Chemical Synthesis
Not Authorised

3.2 Processing Activities of Active Substance from Natural Sources
Not Authorised

3.3 Manufacture of Active Substance using Biological Processes
Not Authorised

3.4 Manufacture of sterile active substance
Not Authorised

3.5 General Finishing Steps
Not Authorised

3.6 Quality Control Testing
Not Authorised

4 Other Activities
Not Authorised

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Any restrictions or clarifying remarks related to the scope of this certificate:

N/A

1. Building(s)/Area(s)

   N/A

2. Room(s)

   N/A

3. Line(s) Equipment(s)

   N/A

4. QC testing

   N/A

5. Medicinal Product(s)/IMP(s)

   N/A

Name of the authorised person of the Competent Authority of the United Kingdom

Dr A J Gray
Head of Inspectorate
inspectionplanning@mhra.gov.uk

Date: 03/11/2021