Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

- 1: Authorisation Number
- 2: Name of authorisation holder
- 3: Address(es) of manufacturing site(s)
- 4: Legally registered address of authorisation holder
- 5: Scope of authorisation and dosage forms
- 6: Legal Basis of authorisation
- 7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation
- 8: Authorisation Date
- 9: Annexes attached

UK MIA(IMP) 44301 CURIUM PHARMA UK LIMITED

CURIUM PHARMA UK LIMITED, 5TH FLOOR, UNIVERSITY COLLEGE HOSPITAL, 235 EUSTON ROAD, LONDON, NW1 2BU, UNITED KINGDOM

CURIUM PHARMA UK LIMITED, 5TH FLOOR, UNIVERSITY COLLEGE HOSPITAL, 235 EUSTON ROAD, LONDON, NW1 2BU, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]

Confidential

18/09/2023

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2 Name and address of the site:

CURIUM PHARMA UK LIMITED, 5TH FLOOR, UNIVERSITY COLLEGE HOSPITAL, 235 EUSTON ROAD, LONDON, NW1 2BU, UNITED KINGDOM

Human Investigational Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

[1.1] Sterile Investigational Medicinal Products

[1.1.1] Aseptically prepared (processing operations for the following dosage forms)

[1.1.1.4] Small volume liquids

Special Requirements

Radiopharmaceuticals

[1.1.3] Batch certification

[1.4] Other investigational medicinal products or manufacturing activitiy

- [1.4.2] Sterilisation of active substances/excipients/finished products:
 - [1.4.2.1] Filtration

[1.6] Quality control testing

- [1.6.1] Microbiological: sterility
- [1.6.2] Microbiological: non-sterility
- [1.6.3] Chemical/Physical