## Medicines and Healthcare products Regulatory Agency

### MANUFACTURER'S AUTHORISATION

1: Authorisation Number UK MIA(IMP) 19162

2: Name of authorisation holder NOTTINGHAM UNIVERSITY HOSPITALS NHS TRUST

THE QMC PHARMACY MANUFACTURING UNIT, PHARMACY

PRODUCTION, QUEENS MEDICAL CENTRE, DERBY ROAD,

NOTTINGHAM, NG7 2UH, UNITED KINGDOM

NOTTINGHAM UNIVERSITY HOSPITALS NHS TRUST, QUEENS

4: Legally registered address of authorisation holder MEDICAL CENTRE CAMPUS, DERBY ROAD, NOTTINGHAM, NG7

2UH, UNITED KINGDOM

5: Scope of authorisation and dosage forms ANNEX 1 and/ or ANNEX 2

6: Legal Basis of authorisation

Part 6 of The Medicines for Human Use (Clinical Trials) Regulations

2004 [SI 2004/1031]

7: Name of responsible officer of the competent authority

of the member state granting the manufacturing

authorisation

Confidential

8: Authorisation Date 17/06/2025

9: Annexes attached Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

3: Address(es) of manufacturing site(s)

#### Annex 2

Name and address of the site:

THE QMC PHARMACY MANUFACTURING UNIT, PHARMACY PRODUCTION, QUEENS MEDICAL CENTRE, DERBY ROAD, NOTTINGHAM, NG7 2UH, 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)

[1.1.1.1] Large volume liquids

[1.1.1.4] Small volume liquids

[1.1.1.6] Other aseptically prepared products

Cytoxics and Radiolabelled products

Issue Date: 17 Jun 2025

# [ 1.2 ] Non-sterile investigational medicinal products

[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)

[1.2.1.1] Capsules, hard shell

[ 1.2.1.5 ] Liquids for external use

[ 1.2.1.6 ] Liquids for internal use

[ 1.2.1.8 ] Other solid dosage forms

[1.2.1.11] Semi-solids

[1.2.1.12] Suppositories

[ 1.2.1.15 ] Other non-sterile medicinal products

Pessaries, multi and unit dose packs. Overlabelling of inhalers

### [ 1.3 ] Biological investigational medicinal products

[1.3.1] Biological medicinal products

[ 1.3.1.2 ] Immunological products

[ 1.3.1.5 ] Biotechnology products

[ 1.3.1.6 ] Human or animal extracted products

[ 1.3.1.8 ] Other biological medicinal products

Heparin Injection (using Product Licensed ingredients) and Monoclonal antibodies (from product licensed ingredients)

### [ 1.4 ] Other investigational medicinal products or manufacturing activitiy

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

[ 1.4.2.1 ] Filtration

### [ 1.5 ] Packaging

[1.5.1] Primary packaging

[1.5.1.2] Capsules, soft shell

[ 1.5.1.13 ] Tablets

[1.5.2] Secondary packaging

### [ 1.6 ] Quality control testing

[ 1.6.1 ] Microbiological: sterility

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

[1.6.4] Biological



Page 2 of 2 Issue Date: 17 Jun 2025