

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
3: Address(es) of manufacturing site(s)	THE QMC PHARMACY MANUFACTURING UNIT, PHARMACY PRODUCTION, QUEENS MEDICAL CENTRE, DERBY ROAD, NOTTINGHAM, NG7 2UH, UNITED KINGDOM
4: Legally registered address of authorisation holder	NOTTINGHAM UNIVERSITY HOSPITALS NHS TRUST, QUEENS 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	07/12/2023
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

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 [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 activity

[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