Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number UK MIA(IMP) 48101
2: Name of authorisation holder MAC RESEARCH LTD

MAC CLINICAL RESEARCH MANCHESTER-MAC CLINICAL RESEARCH FINANCE LIMITED, CITYLABS 1.0, NELSON STREET,

MANCHESTER, M13 9NQ, UNITED KINGDOM

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

MAC CLINICAL RESEARCH LIMITED - MAC CLINICAL RESEARCH

FINANCE LTD, MONARCH HOUSE, WAKEFIELD ROAD, LEEDS,

LS10 1DP, UNITED KINGDOM

4: Legally registered address of authorisation holder

1PW, 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 06/10/2025

9: Annexes attached Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

MAC CLINICAL RESEARCH MANCHESTER-MAC CLINICAL RESEARCH FINANCE LIMITED, CITYLABS 1.0, NELSON STREET, MANCHESTER, M13 9NQ, UNITED KINGDOM

Human Investigational Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

[1.5] Packaging

[1.5.1] Primary packaging

[1.5.1.1] Capsules, hard shell

Issue Date: 06 Oct 2025

- [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
 - SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

MAC CLINICAL RESEARCH LIMITED - MAC CLINICAL RESEARCH FINANCE LTD, MONARCH HOUSE, WAKEFIELD ROAD, LEEDS, LS10 1DP, 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

Manipulation of previously formulated products e.g. cytotoxics and monoclonal antibodies, GMOs and vaccines.

[1.1.3] Batch certification

[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
 - [1.2.1.6] Liquids for internal use
 - [1.2.1.8] Other solid dosage forms
 - [1.2.1.15] Other non-sterile medicinal products

Weighing and mixing of drug substance into individual subject dose containers

[1.2.2] 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

[1.5] Packaging

- [1.5.1] Primary packaging
 - [1.5.1.1] Capsules, hard shell
 - [1.5.1.2] Capsules, soft shell
 - [1.5.1.5] Liquids for external use
 - [1.5.1.6] Liquids for internal use
 - [1.5.1.8] Other solid dosage forms
 - [1.5.1.11] Semi-solids

Issue Date: 06 Oct 2025

[1.5.1.13] Tablets

[1.5.1.15] Other non-sterile medicinal products

Weighing and mixing of drug substance into individual subject dose containers

[1.5.2] Secondary packaging

[1.6] Quality control testing

[1.6.3] Chemical/Physical

