Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA(IMP) 59823
2: Name of authorisation holder	SEDA CLINICAL MANUFACTURING SERVICES LTD.
3: Address(es) of manufacturing site(s)	SEDA CLINICAL MANUFACTURING SERVICES LTD., 5000 LAKESIDE, CHEADLE ROYAL BUSINESS PARK, CHEADLE, SK8 3AX, UNITED KINGDOM
4: Legally registered address of authorisation holder	SEDA CLINICAL MANUFACTURING SERVICES LTD., 5000 LAKESIDE, CHEADLE ROYAL BUSINESS PARK, CHEADLE, SK8 3AX, 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	22/01/2026
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

SEDA CLINICAL MANUFACTURING SERVICES LTD., 5000 LAKESIDE, CHEADLE ROYAL BUSINESS PARK, CHEADLE, SK8 3AX, UNITED KINGDOM

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

[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

[1.2.1.8] Other solid dosage forms

Special Requirements

Powders

[1.2.1.13] Tablets

MHRA-GMDP

[1.2.2] Batch certification

[1.5] Packaging

[1.5.1] Primary packaging

[1.5.1.1] Capsules, hard shell

[1.5.1.8] Other solid dosage forms

[1.5.1.11] Semi-solids

[1.5.1.13] Tablets

[1.5.2] Secondary packaging

[1.6] Quality control testing

[1.6.3] Chemical/Physical

MHRA-GMDP