

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA(IMP) 59823
<b>2: Name of authorisation holder</b>	SEDA CLINICAL MANUFACTURING SERVICES LTD.
<b>3: Address(es) of manufacturing site(s)</b>	SEDA CLINICAL MANUFACTURING SERVICES LTD., 5000 LAKESIDE, CHEADLE ROYAL BUSINESS PARK, CHEADLE, SK8 3AX, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	SEDA CLINICAL MANUFACTURING SERVICES LTD., 5000 LAKESIDE, CHEADLE ROYAL BUSINESS PARK, CHEADLE, SK8 3AX, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]
<b>7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation</b>	Confidential
<b>8: Authorisation Date</b>	20/02/2025
<b>9: Annexes attached</b>	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)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.2 ] Non-sterile investigational medicinal products</b> [ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms) [ 1.2.1.1 ] Capsules, hard shell [ 1.2.1.13 ] Tablets [ 1.2.2 ] Batch certification <b>[ 1.5 ] Packaging</b> [ 1.5.1 ] Primary packaging

[ 1.5.1.1 ] Capsules, hard shell

[ 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