

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA(IMP) 50416
<b>2: Name of authorisation holder</b>	STERLING DEESIDE LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	STERLING DEESIDE LIMITED, UNIT 103, TENTH AVENUE, ZONE 3, DEESIDE INDUSTRIAL ESTATE, DEESIDE, CH5 2UA, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	STERLING DEESIDE LIMITED, UNIT 103, TENTH AVENUE, ZONE 3, DEESIDE INDUSTRIAL ESTATE, DEESIDE, CH5 2UA, 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>	23/01/2026
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

**STERLING DEESIDE LIMITED**, UNIT 103, TENTH AVENUE, ZONE 3, DEESIDE INDUSTRIAL ESTATE, DEESIDE, CH5 2UA,  
UNITED KINGDOM

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.3 ] Biological investigational medicinal products</b> [ 1.3.1 ] Biological medicinal products [ 1.3.1.2 ] Immunological products <b>[ 1.6 ] Quality control testing</b> [ 1.6.2 ] Microbiological: non-sterility [ 1.6.3 ] Chemical/Physical [ 1.6.4 ] Biological