

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

**1: Authorisation Number**

UK MIA(IMP) 50416

**2: Name of authorisation holder**

STERLING DEESIDE LIMITED

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

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

**4: Legally registered address of authorisation holder**

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

23/01/2026

**9: Annexes attached**

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