

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 20/08/2025

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)
Part 1 - MANUFACTURING OPERATIONS [1.3] Biological investigational medicinal products [1.3.1] Biological medicinal products [1.3.1.2] Immunological products [1.6] Quality control testing [1.6.2] Microbiological: non-sterility [1.6.3] Chemical/Physical [1.6.4] Biological