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	13/10/2026
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)

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

MHRA-GMDP MHRA-GMDP MHRA