

Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA(IMP) 30013
2: Name of authorisation holder	PHARMARON BIOLOGICS (UK) LTD
3: Address(es) of manufacturing site(s)	PHARMARON BIOLOGICS (UK) LTD, NATIONAL BIOMANUFACTURING CENTRE, ESTUARY BANKS, ESTUARY COMMERCE PARK, SPEKE ROAD, LIVERPOOL, L24 8RB, UNITED KINGDOM
4: Legally registered address of authorisation holder	PHARMARON BIOLOGICS (UK) LTD, NATIONAL BIOMANUFACTURING CENTRE, ESTUARY BANKS, ESTUARY COMMERCE PARK, SPEKE ROAD, LIVERPOOL, L24 8RB, 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	01/09/2026
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

PHARMARON BIOLOGICS (UK) LTD, NATIONAL BIOMANUFACTURING CENTRE, ESTUARY BANKS, ESTUARY COMMERCE PARK, SPEKE ROAD, LIVERPOOL, L24 8RB, UNITED KINGDOM

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)
Part 1 - MANUFACTURING OPERATIONS [1.1] Sterile Investigational Medicinal Products [1.1.3] Batch certification [1.2] Non-sterile investigational medicinal products [1.2.2] Batch certification [1.3] Biological investigational medicinal products

- [1.3.1] Biological medicinal products
 - [1.3.1.2] Immunological products
 - [1.3.1.3] Cell therapy products
 - [1.3.1.4] Gene therapy products
 - [1.3.1.5] Biotechnology products
 - [1.3.1.6] Human or animal extracted products
 - [1.3.1.8] Other biological medicinal products
- Biological Active Starting Materials

[1.6] Quality control testing

- [1.6.2] Microbiological: non-sterility
- [1.6.3] Chemical/Physical
- [1.6.4] Biological

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

[2.2] Batch certification of imported medicinal products

- [2.2.1] Sterile Products
 - [2.2.1.1] Aseptically prepared
 - [2.2.1.2] Terminally sterilised
- [2.2.2] Non-sterile products
- [2.2.3] Biological medicinal products
 - [2.2.3.2] Immunological products
 - [2.2.3.3] Cell therapy products
 - [2.2.3.4] Gene therapy products
 - [2.2.3.5] Biotechnology products
 - [2.2.3.6] Human or animal extracted products
 - [2.2.3.8] Other biological medicinal products
- Biological Active Starting Materials