

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

**1: Authorisation Number**

UK MIA 51841

**2: Name of authorisation holder**

GMP MANUFACTURING LTD

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

GMP MANUFACTURING LTD, PARK ROYAL HOUSE AND MARFLEET HOUSE, VALLETTA STREET, HULL, HU9 5NP, UNITED KINGDOM

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

GMP MANUFACTURING LTD, MARFLEET HOUSE, VALLETTA STREET, HULL, HU9 5NP, UNITED KINGDOM

**5: Scope of authorisation and dosage forms**

ANNEX 1 and/ or ANNEX 2

**6: Legal Basis of authorisation**

Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)

**7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation**

Confidential

**8: Authorisation Date**

22/01/2026

**9: Annexes attached**

Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**GMP MANUFACTURING LTD, PARK ROYAL HOUSE AND MARFLEET HOUSE, VALLETTA STREET, HULL, HU9 5NP, UNITED KINGDOM**

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)
<b>Part 1 - MANUFACTURING OPERATIONS</b>
<b>[ 1.2 ] Non-sterile products</b>
[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)
[ 1.2.1.5 ] Liquids for external use
[ 1.2.1.11 ] Semi-solids
[ 1.2.2 ] Batch certification
<b>[ 1.5 ] Packaging</b>
[ 1.5.1 ] Primary packaging

[ 1.5.1.5 ] Liquids for external use

[ 1.5.1.11 ] Semi-solids

[ 1.5.2 ] Secondary packaging

**[ 1.6 ] Quality control testing**

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

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

MHRA-GMDP

MHRA

MHRA-GMDP

MHRA