# Medicines and Healthcare products Regulatory Agency

# MANUFACTURER'S AUTHORISATION

1: Authorisation Number

2: Name of authorisation holder

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

4: Legally registered address of authorisation holder

5: Scope of authorisation and dosage forms

6: Legal Basis of authorisation

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

8: Authorisation Date

9: Annexes attached

**UK MIA 51841** 

**GMP MANUFACTURING LTD** 

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

GMP MANUFACTURING LTD, MARFLEET HOUSE,

VALLETTA STREET, HULL, HU9 5NP, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI

2012/1916)

Confidential

19/04/2024

Annex 1 and/or Annex 2

## SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

GMP MANUFACTURING LTD, 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)

#### Part 1 - MANUFACTURING OPERATIONS

## [ 1.5 ] Packaging

[ 1.5.1 ] Primary packaging

[1.5.1.5] Liquids for external use

[1.5.1.11] Semi-solids

[1.5.2] Secondary packaging

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

Issue Date: 19 Apr 2024

