# 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 4543

**CP PHARMACEUTICALS LIMITED** 

CP PHARMACEUTICALS LIMITED, ASH ROAD NORTH, WREXHAM, LL13 9UF, UNITED KINGDOM

CP PHARMACEUTICALS LIMITED, ASH ROAD NORTH, WREXHAM, LL13 9UF, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

29/05/2025

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

### CP PHARMACEUTICALS LIMITED, ASH ROAD NORTH, WREXHAM, LL13 9UF, 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

### [ 1.1 ] Sterile Products

[ 1.1.1 ] Aseptically prepared (processing operations for the following dosage forms)

[1.1.1.2] Lyophilisates

[1.1.1.4] Small volume liquids

[ 1.1.2 ] Terminally Sterilised (processing operations for the following dosage forms)

[1.1.2.3] Small volume liquids

[1.1.3] Batch certification

#### [ 1.2 ] Non-sterile products

[1.2.1] Non-Sterile Products (processing operations for the following dosage forms)

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[ 1.2.1.17 ] Other non-sterile medicinal products **Nasal Sprays** [1.2.2] Batch certification [ 1.3 ] Biological medicinal products [ 1.3.1 ] Biological medicinal products [1.3.1.2] Immunological products [ 1.3.1.5 ] Biotechnology products [ 1.3.1.6 ] Human or animal extracted products [ 1.3.1.8 ] Other biological medicinal products manufacture of products based on botanical extracts and hormones [1.3.2] Batch certification [ 1.3.2.5 ] Biotechnology products [1.3.2.6] Human or animal extracted products [ 1.3.2.8 ] Other biological medicinal products manufacture of products based on botanical extracts and hormones [ 1.4 ] Other products or manufacturing activity [ 1.4.1 ] Manufacture of: [1.4.1.4] Products authorised under regulation 174 (supply in response to spread of pathogenic agents etc) [ 1.4.2 ] Sterilisation of active substances/excipients/finished products: [ 1.4.2.1 ] Filtration [ 1.4.2.2 ] Dry heat [ 1.4.2.3 ] Moist heat [ 1.5 ] Packaging [1.5.1] Primary packaging [ 1.5.1.17 ] Other non-sterile medicinal products Nasal spray [1.5.2] Secondary packaging [ 1.6 ] Quality control testing [ 1.6.1 ] Microbiological: sterility [ 1.6.2 ] Microbiological: non-sterility [1.6.3] Chemical/Physical [1.6.4] Biological Part 2 - IMPORTATION OF MEDICINAL PRODUCTS [ 2.1 ] Quality control testing of imported medicinal products [2.1.1] Microbiological: sterility [2.1.2] Microbiological: non-sterility [2.1.3] Chemical/Physical [2.1.4] Biological [ 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 [2.2.3.5] Biotechnology products

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[ 2.2.3.6 ] Human or animal extracted products

