# 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 20492

CD PHARMA LIMITED

TELETA HOUSE, 4 CAIRN COURT, EAST KILBRIDE, GLASGOW, G74 4NB, UNITED KINGDOM

CD PHARMA LIMITED, 4 CAIRN COURT, EAST KILBRIDE, GLASGOW, G74 4NB, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

17/07/2025

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

### Annex 1

Name and address of the site:

TELETA HOUSE, 4 CAIRN COURT, EAST KILBRIDE, GLASGOW, G74 4NB, UNITED KINGDOM

**Human Medicinal Products** 

**Authorised Operations** 

MANUFACTURING OPERATIONS (according to part 1)

## Part 1 - MANUFACTURING OPERATIONS

[ 1.1 ] Sterile Products

[1.1.3] Batch certification

[ 1.2 ] Non-sterile products

[ 1.2.2 ] Batch certification

[ 1.5 ] Packaging

[1.5.2] Secondary packaging

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