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

13/01/2025

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

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Human Medicinal Products	
Authorised Operations	
MANUFACTURING OPERATIONS (according to part 1)	
Part 1 - MANUFACTURING OPERATIONS	
[1.1] Sterile Products	
[1.1.3] Batch certification	<pre></pre>
[1.2] Non-sterile products	
[1.2.2] Batch certification	
[1.5] Packaging	
[1.5.2] Secondary packaging	