# 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 8553

DR REDDY'S LABORATORIES (UK) LIMITED

DR. REDDY'S LABORATORIES (UK) LIMITED, 410 CAMBRIDGE SCIENCE PARK, MILTON ROAD, CAMBRIDGE, CB4 0PE, UNITED KINGDOM

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ANNEX 1 and/ or ANNEX 2

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

Confidential

19/06/2025

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

DR. REDDY'S LABORATORIES (UK) LIMITED, 410 CAMBRIDGE SCIENCE PARK, MILTON ROAD, CAMBRIDGE, CB4 0PE, UNITED KINGDOM

Human Medicinal Products

Authorised Operations

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

### 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

[2.2.3] Biological medicinal products

[2.2.3.5] Biotechnology products

#### [2.2.3.8] Other biological medicinal products

Monoclonal antibody, Fusion protein, Recombinant protein

## [ 2.3 ] Other Importation Activities

- [ 2.3.2 ] Importation of Intermediate which undergoes further processing
- [2.3.4] Products authorised under regulation 174 (supply in response to spread of pathogenic agents etc)
  - Importation of finished bulk for final packaging