

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

**1: Authorisation Number** UK MIA 8553

**2: Name of authorisation holder** DR REDDY'S LABORATORIES (UK) LIMITED

**3: Address(es) of manufacturing site(s)** DR. REDDY'S LABORATORIES (UK) LIMITED, 410 CAMBRIDGE SCIENCE PARK, MILTON ROAD, CAMBRIDGE, CB4 0PE, UNITED KINGDOM

**4: Legally registered address of authorisation holder** DR REDDY'S LABORATORIES (UK) LIMITED, 410 CAMBRIDGE SCIENCE PARK, MILTON ROAD, CAMBRIDGE, CB4 0PE, UNITED KINGDOM

**5: Scope of authorisation and dosage forms** ANNEX 1 and/ or ANNEX 2

**6: Legal Basis of authorisation** Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)

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

**8: Authorisation Date** 24/11/2025

**9: Annexes attached** 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

|                                                                                                                                                                                                                                                                                                                                    |
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| Human Medicinal Products                                                                                                                                                                                                                                                                                                           |
| Authorised Operations                                                                                                                                                                                                                                                                                                              |
| IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)                                                                                                                                                                                                                                                                            |
| Part 2 - IMPORTATION OF MEDICINAL PRODUCTS<br>[ 2.2 ] Batch certification of imported medicinal products<br>[ 2.2.1 ] Sterile Products<br>[ 2.2.1.1 ] Aseptically prepared<br>[ 2.2.1.2 ] Terminally sterilised<br>[ 2.2.2 ] Non-sterile products<br>[ 2.2.3 ] Biological medicinal products<br>[ 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