

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

**1: Authorisation Number** UK MIA 45001

**2: Name of authorisation holder** QMED PHARMACEUTICALS LIMITED

**3: Address(es) of manufacturing site(s)** QMED PHARMACEUTICALS LIMITED, UNIT 4, THORNHAM GROVE INDUSTRIAL ESTATE, LONDON, E15 1DN, UNITED KINGDOM

**4: Legally registered address of authorisation holder** QMED PHARMACEUTICALS LIMITED, 168 PEMBROKE ROAD, ILFORD, IG3 8PG, 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** 27/02/2026

**9: Annexes attached** Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**QMED PHARMACEUTICALS LIMITED**, UNIT 4, THORNHAM GROVE INDUSTRIAL ESTATE, LONDON, E15 1DN, UNITED KINGDOM

|                                                                                                                                                                                                                                                                                                                                |
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| Human Medicinal Products                                                                                                                                                                                                                                                                                                       |
| Authorised Operations                                                                                                                                                                                                                                                                                                          |
| MANUFACTURING OPERATIONS (according to part 1)<br>IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)                                                                                                                                                                                                                      |
| <b>Part 1 - MANUFACTURING OPERATIONS</b><br><b>[ 1.2 ] Non-sterile products</b><br>[ 1.2.2 ] Batch certification<br><b>[ 1.5 ] Packaging</b><br>[ 1.5.2 ] Secondary packaging<br>Part 2 - IMPORTATION OF MEDICINAL PRODUCTS<br><b>[ 2.2 ] Batch certification of imported medicinal products</b><br>[ 2.2.1 ] Sterile Products |

[ 2.2.1.1 ] Aseptically prepared

[ 2.2.1.2 ] Terminally sterilised

[ 2.2.2 ] Non-sterile products

**[ 2.3 ] Other Importation Activities**

[ 2.3.1 ] Site of Physical Importation