

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

**1: Authorisation Number** UK MIA 20395

**2: Name of authorisation holder** RELONCHEM LIMITED

**3: Address(es) of manufacturing site(s)** RELONCHEM LIMITED, CHESHIRE HOUSE, GORSEY LANE, WIDNES, WA8 0RP, UNITED KINGDOM

**4: Legally registered address of authorisation holder** RELONCHEM LIMITED, 27 OLD GLOUCESTER STREET, Array, LONDON, WC1N 3AX, 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** 03/09/2026

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

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**RELONCHEM LIMITED**, CHESHIRE HOUSE, GORSEY LANE, WIDNES, WA8 0RP, 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>Part 2 - IMPORTATION OF MEDICINAL PRODUCTS</b><br><b>[ 2.2 ] Batch certification of imported medicinal products</b><br>[ 2.2.2 ] Non-sterile products |