

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

|                                                                                                                              |                                                                                                                                       |
|------------------------------------------------------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------|
| 1: Authorisation Number                                                                                                      | UK MIA 427                                                                                                                            |
| 2: Name of authorisation holder                                                                                              | ROSEMONT PHARMACEUTICALS LIMITED                                                                                                      |
| 3: Address(es) of manufacturing site(s)                                                                                      | ROSEMONT PHARMACEUTICALS LIMITED, ROSEMONT HOUSE,<br>YORKDALE INDUSTRIAL PARK, BRAITHWAITE STREET, LEEDS,<br>LS11 9XE, UNITED KINGDOM |
| 4: Legally registered address of authorisation holder                                                                        | ROSEMONT PHARMACEUTICALS LIMITED, ROSEMONT HOUSE,<br>YORKDALE INDUSTRIAL PARK, BRAITHWAITE STREET, LEEDS,<br>LS11 9XE, 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<br>2012/1916)                                                               |
| 7: Name of responsible officer of the competent authority<br>of the member state granting the manufacturing<br>authorisation | Confidential                                                                                                                          |
| 8: Authorisation Date                                                                                                        | 04/02/2026                                                                                                                            |
| 9: Annexes attached                                                                                                          | Annex 1 and/or Annex 2                                                                                                                |

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**ROSEMONT PHARMACEUTICALS LIMITED**, ROSEMONT HOUSE, YORKDALE INDUSTRIAL PARK, BRAITHWAITE STREET,  
LEEDS, LS11 9XE, UNITED KINGDOM

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| Human Medicinal Products                                                                                                                                                                                                                                                                                                                                                 |
| Authorised Operations                                                                                                                                                                                                                                                                                                                                                    |
| MANUFACTURING OPERATIONS (according to part 1)                                                                                                                                                                                                                                                                                                                           |
| <b>Part 1 - MANUFACTURING OPERATIONS</b><br><b>[ 1.2 ] Non-sterile products</b><br>[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)<br>[ 1.2.1.6 ] Liquids for internal use<br>[ 1.2.1.8 ] Other solid dosage forms<br>[ 1.2.1.17 ] Other non-sterile medicinal products<br>Manufacture of cytotoxic agents - Methotrexate disodium |

[ 1.2.2 ] Batch certification

**[ 1.5 ] Packaging**

[ 1.5.1 ] Primary packaging

[ 1.5.1.6 ] Liquids for internal use

[ 1.5.2 ] Secondary packaging

**[ 1.6 ] Quality control testing**

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical