

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

**1: Authorisation Number** UK MIA 51718

**2: Name of authorisation holder** CLYDESDALE PHARMA LTD

**3: Address(es) of manufacturing site(s)** CLYDESDALE PHARMA LTD, UNIT 3-5 CAMPBELL COURT, BRAMLEY, TADLEY, RG26 5EG, UNITED KINGDOM

**4: Legally registered address of authorisation holder** CLYDESDALE PHARMA LTD, UNIT 3-5 CAMPBELL COURT, BRAMLEY, TADLEY, RG26 5EG, 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** 03/07/2025

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

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**CLYDESDALE PHARMA LTD**, UNIT 3-5 CAMPBELL COURT, BRAMLEY, TADLEY, RG26 5EG, 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>[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)<br>[ 1.2.1.1 ] Capsules, hard shell<br>[ 1.2.1.2 ] Capsules, soft shell<br>[ 1.2.1.3 ] Chewing gums<br>[ 1.2.1.5 ] Liquids for external use<br>[ 1.2.1.6 ] Liquids for internal use |

[ 1.2.1.8 ] Other solid dosage forms

[ 1.2.1.9 ] Pressurised preparations

[ 1.2.1.11 ] Semi-solids

[ 1.2.1.12 ] Suppositories

[ 1.2.1.13 ] Tablets

[ 1.2.1.14 ] Transdermal patches

[ 1.2.2 ] Batch certification

**[ 1.6 ] Quality control testing**

[ 1.6.3 ] Chemical/Physical

**Part 2 - IMPORTATION OF MEDICINAL PRODUCTS**

**[ 2.1 ] Quality control testing of imported medicinal products**

[ 2.1.3 ] Chemical/Physical

**[ 2.2 ] Batch certification of imported medicinal products**

[ 2.2.2 ] Non-sterile products

**[ 2.3 ] Other Importation Activities**

[ 2.3.1 ] Site of Physical Importation