Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number

2: Name of authorisation holder

3: Address(es) of manufacturing site(s)

4: Legally registered address of authorisation holder

5: Scope of authorisation and dosage forms

6: Legal Basis of authorisation

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

8: Authorisation Date

9: Annexes attached

UK MIA 51718

CLYDESDALE PHARMA LTD

CLYDESDALE PHARMA LTD, UNIT 3-5 CAMPBELL COURT, BRAMLEY, TADLEY, RG26 5EG, UNITED KINGDOM

CLYDESDALE PHARMA LTD, UNIT 3-5 CAMPBELL COURT,

BRAMLEY, TADLEY, RG26 5EG, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI

2012/1916)

Confidential

22/03/2024

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

Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

Part 1 - MANUFACTURING OPERATIONS

[1.2] Non-sterile products

[1.2.1] Non-Sterile Products (processing operations for the following dosage forms)

[1.2.1.1] Capsules, hard shell

[1.2.1.2] Capsules, soft shell

[1.2.1.3] Chewing gums

[1.2.1.5] Liquids for external use

[1.2.1.6] Liquids for internal use

[1.2.1.8] Other solid dosage forms

[1.2.1.9] Pressurised preparations

Issue Date: 22 Mar 2024

[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



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