# 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(IMP) 322 NORGINE LIMITED

NORGINE LIMITED, NEW ROAD, TIR-Y-BERTH, HENGOED, CF82 8SJ, UNITED KINGDOM

NORGINE LIMITED, ARC UXBRIDGE, BUILDING 01, SANDERSON ROAD, UXBRIDGE, UB8 1DH, UNITED

**KINGDOM** 

ANNEX 1 and/ or ANNEX 2

Part 6 of The Medicines for Human Use (Clinical Trials)

Regulations 2004 [SI 2004/1031]

Confidential

24/10/2025

Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

NORGINE LIMITED, NEW ROAD, TIR-Y-BERTH, HENGOED, CF82 8SJ, UNITED KINGDOM

**Human Investigational Medicinal Products** 

**Authorised Operations** 

MANUFACTURING OPERATIONS (according to part 1)

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

# Part 1 - MANUFACTURING OPERATIONS

# [ 1.1 ] Sterile Investigational Medicinal Products

[ 1.1.3 ] Batch certification

# [ 1.2 ] Non-sterile investigational medicinal products

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

[1.2.1.1] Capsules, hard shell

[ 1.2.1.5 ] Liquids for external use

[1.2.1.6] Liquids for internal use

[1.2.1.8] Other solid dosage forms

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[1.2.1.12] Suppositories [ 1.5 ] Packaging [1.5.1] Primary packaging [1.5.1.1] Capsules, hard shell [1.5.1.2] Capsules, soft shell [ 1.5.1.5 ] Liquids for external use [ 1.5.1.6 ] Liquids for internal use [ 1.5.1.8 ] Other solid dosage forms [1.5.1.11] Semi-solids [ 1.5.1.13 ] Tablets [1.5.1.15] Other non-sterile medicinal products Pessaries, Powders [1.5.2] Secondary packaging [ 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 [2.2.1] Sterile Products [2.2.1.1] Aseptically prepared [2.2.1.2] Terminally sterilised [2.2.2] Non-sterile products



Importation of QP-certified IMPs from a country on the approved country for import list

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[ 2.3 ] Other Importation Activities

[ 2.3.4 ] Other