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 4509 ECOLAB LIMITED

ECOLAB LIMITED, UNIT 1, WERNDDU COURT,

ECOLAB LIMITED, PO BOX 11, WINNINGTON AVENUE,

NORTHWICH, CW8 4DX, UNITED KINGDOM

CAERPHILLY, CF83 3SG, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012

(SI 2012/1916)

Confidential

24/10/2022

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

ECOLAB LIMITED, UNIT 1, WERNDDU COURT, CAERPHILLY, CF83 3SG, UNITED KINGDOM

Human Medicinal Products

Authorised Operations

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

[2.2] Batch certification of imported medicinal products

[2.2.2] Non-sterile products

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