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 56177

ECOLAB MANUFACTURING UK LIMITED

ECOLAB MANUFACTURING UK LTD, BAGLAN ENERGY PARK, BRUNEL WAY, BRITON FERRY, NEATH, SA11 2GA, UNITED KINGDOM

ECOLAB MANUFACTURING UK LIMITED, BAGLAN ENERGY PARK, BRUNEL WAY, BRITON FERRY, NEATH, SA11 2GA, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)

Confidential

15/05/2025

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

ECOLAB MANUFACTURING UK LTD, BAGLA	N ENERGY PARK,	BRUNEL	WAY, BRIT	ON FERRY,	NEATH, SA11 2GA	, UNITED
KINGDOM						

Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

[1.2] Non-sterile products

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

[1.2.1.5] Liquids for external use

[1.2.2] Batch certification

[1.5] Packaging

[1.5.1] Primary packaging

[1.5.1.5] Liquids for external use

[1.5.2] Secondary packaging

[1.6] Quality control testing

[1.6.2] Microbiological: non-sterility

[1.6.3] Chemical/Physical