Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA 43685
2: Name of authorisation holder	WASDELL MANUFACTURING LIMITED
3: Address(es) of manufacturing site(s)	WASDELL MANUFACTURING LIMITED, UNIT 3 & 4, GATEWAY WEST, GREENFINCH WAY, NEWBURN, NEWCASTLE UPON TYNE, NE15 8NX, UNITED KINGDOM
4: Legally registered address of authorisation holder	WASDELL MANUFACTURING LIMITED, UNITS 1, 2, 3, 5, 6, 7 & 8 EURO WAY INDUSTRIAL ESTATE, BLAGROVE, SWINDON, SN5 8YW, 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	26/03/2026
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

WASDELL MANUFACTURING LIMITED, UNIT 3 & 4, GATEWAY WEST, GREENFINCH WAY, NEWBURN, NEWCASTLE UPON TYNE, NE15 8NX, 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

[1.2.1.6] Liquids for internal use

[1.2.2] Batch certification

[1.5] Packaging

MHRA-GMDP MHRA-GMDP MHRA

[1.5.1] Primary packaging

[1.5.1.5] Liquids for external use

[1.5.1.6] Liquids for internal use

[1.5.2] Secondary packaging

[1.6] Quality control testing

[1.6.2] Microbiological: non-sterility

[1.6.3] Chemical/Physical

MHRA-GMDP MHRA MHRA-GMDP MHRA