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 20504 CROSS HEALTHCARE LIMITED

CROSS HEALTHCARE LIMITED, UNIT 2A, BANDEATH INDUSTRIAL ESTATE, THROSK, STIRLING, FK7 7NP, UNITED KINGDOM

CROSS HEALTHCARE LIMITED, UNIT 2A, BANDEATH INDUSTRIAL ESTATE, THROSK, STIRLING, FK7 7NP, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

21/01/2025

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

CROSS HEALTHCARE LIMITED, UNIT 2A, BANDEATH INDUSTRIAL ESTATE, THROSK, STIRLING, FK7 7NP, UNITED KINGDOM

Human Medicinal Products	·
Authorised Operations	
Part 1 - MANUFACTURING OPERATIONS	~
[1.5] Packaging [1.5.2] Secondary packaging	
	NHK