## 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 3422

CROSS VETPHARM GROUP UK LIMITED

CROSS VETPHARM GROUP UK LIMITED, UNIT 2, BRYN CEFNI INDUSTRIAL PARK, LLANGEFNI, LL77 7XA, UNITED KINGDOM

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ANNEX 1 and/ or ANNEX 2

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

Confidential

15/07/2024

Annex 1 and/or Annex 2

## SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

CROSS VETPHARM GROUP UK LIMITED, UNIT 2, BRYN CEFNI INDUSTRIAL PARK, LLANGEFNI, LL77 7XA, UNITED KINGDOM

Human Medicinal Products	
Authorised Operations	
MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)	
Part 1 - MANUFACTURING OPERATIONS [ 1.4 ] Other products or manufacturing activity [ 1.4.1 ] Manufacture of: [ 1.4.1.3 ] Other Site of physical importation Part 2 - IMPORTATION OF MEDICINAL PRODUCTS [ 2.2 ] Batch certification of imported medicinal products	R
[ 2.2.1 ] Sterile Products	