# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

1: Authorisation Number

**UK MIA 3422** 

2: Name of authorisation holder

**CROSS VETPHARM GROUP UK LIMITED** 

3: Address(es) of manufacturing site(s)

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

4: Legally registered address of authorisation holder

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

15/07/2024

9: Annexes attached

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

[ 2.2.1 ] Sterile Products

[2.2.1.1] Aseptically prepared

[ 2.2.2 ] Non-sterile products

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