

# 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, LLANGFNI, LL77 7XA, UNITED KINGDOM

**4: Legally registered address of authorisation holder** CROSS VETPHARM GROUP UK LIMITED, UNIT 2, BRYN CEFNI INDUSTRIAL PARK, LLANGFNI, 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, LLANGFNI, LL77 7XA, UNITED KINGDOM

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

[ 2.2.1.1 ] Aseptically prepared

[ 2.2.2 ] Non-sterile products