

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA 3422
<b>2: Name of authorisation holder</b>	CROSS VETPHARM GROUP UK LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	CROSS VETPHARM GROUP UK LIMITED, UNIT 2, BRYN CEFNI INDUSTRIAL PARK, LLANGEFNI, LL77 7XA, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	CROSS VETPHARM GROUP UK LIMITED, UNIT 2, BRYN CEFNI INDUSTRIAL PARK, LLANGEFNI, LL77 7XA, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)
<b>7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation</b>	Confidential
<b>8: Authorisation Date</b>	01/08/2022
<b>9: Annexes attached</b>	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)
<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