

Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA 8794
2: Name of authorisation holder	MICROPHARM LIMITED MICROPHARM LIMITED, UNITS B2, B1 & D1, ANTUR TEIFI BUSINESS PARK, NEWCASTLE EMLYN, SA38 9DB, UNITED KINGDOM
3: Address(es) of manufacturing site(s)	MICROPHARM LIMITED, STATION ROAD INDUSTRIAL ESTATE, NEWCASTLE EMLYN, SA38 9BY, UNITED KINGDOM MICROPHARM LIMITED, CNWCAU, CILGERRAN, CARDIGAN, SA43 2SN, UNITED KINGDOM
4: Legally registered address of authorisation holder	MICROPHARM LIMITED, STATION ROAD INDUSTRIAL ESTATE, NEWCASTLE EMLYN, SA38 9BY, 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	15/03/2024
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

MICROPHARM LIMITED, UNITS B2, B1 & D1, ANTUR TEIFI BUSINESS PARK, NEWCASTLE EMLYN, SA38 9DB, UNITED KINGDOM

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
Part 1 - MANUFACTURING OPERATIONS [1.1] Sterile Products [1.1.1] Aseptically prepared (processing operations for the following dosage forms) [1.1.1.4] Small volume liquids

- [1.1.1.6] Other aseptically prepared products
Viper Vet manufactured for trials in accordance with ATC

[1.2] Non-sterile products

- [1.2.1] Non-Sterile Products (processing operations for the following dosage forms)
 - [1.2.1.6] Liquids for internal use

[1.3] Biological medicinal products

- [1.3.1] Biological medicinal products
 - [1.3.1.2] Immunological products
 - [1.3.1.6] Human or animal extracted products

[1.4] Other products or manufacturing activity

- [1.4.2] Sterilisation of active substances/excipients/finished products:
 - [1.4.2.1] Filtration

[1.5] Packaging

- [1.5.1] Primary packaging
 - [1.5.1.6] Liquids for internal use
- [1.5.2] Secondary packaging

[1.6] Quality control testing

- [1.6.2] Microbiological: non-sterility
- [1.6.3] Chemical/Physical

SCOPE OF AUTHORISATION

Annex 1

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Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)
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<p>Part 1 - MANUFACTURING OPERATIONS</p> <p>[1.1] Sterile Products</p> <ul style="list-style-type: none"> [1.1.3] Batch certification <p>[1.3] Biological medicinal products</p> <ul style="list-style-type: none"> [1.3.2] Batch certification <ul style="list-style-type: none"> [1.3.2.2] Immunological products [1.3.2.6] Human or animal extracted products <p>[1.6] Quality control testing</p> <ul style="list-style-type: none"> [1.6.3] Chemical/Physical [1.6.4] Biological
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SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

MICROPHARM LIMITED, CNWCAU, CILGERRAN, CARDIGAN, SA43 2SN, UNITED KINGDOM

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
Part 1 - MANUFACTURING OPERATIONS [1.1] Sterile Products [1.1.3] Batch certification [1.3] Biological medicinal products [1.3.1] Biological medicinal products [1.3.1.2] Immunological products Special Requirements Immunoglobulin F(ab') ₂ [1.3.1.6] Human or animal extracted products [1.6] Quality control testing [1.6.2] Microbiological: non-sterility [1.6.3] Chemical/Physical [1.6.4] Biological