Bovilis Cryptium (--) - Emulsion for injection

Authorised

- Cryptosporidium parvum, glycoprotein gp40

Product identification

Medicine name:
Bovilis Cryptium (--) - Emulsion for injection

Active substance:
Cryptosporidium parvum, glycoprotein gp40

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Cryptosporidium parvum, glycoprotein gp40
1.00 unit(s) / 1.00 Vial

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AO02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (PET), Package_size:1 vial (5 doses), Content:10 ml

Packaging:Vial (glass), Package_size:1 vial (5 doses), Content:10 ml

Packaging:Vial (glass), Package_size:10 vial (10 x 1 doses), Content:2 ml

Packaging:Vial (PET), Package_size:1 vial (50 doses), Content:100 ml

Packaging:Vial (PET), Package_size:1 vial (20 doses), Content:40 ml

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application - new active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
23/11/2023

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
23/11/2023

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)
Published on: 19/12/2024
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