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Bovilis BVD Suspension for Injection for Cattle

Authorised

- Bovine viral diarrhoea virus 1, strain C-86 (cytopathic), Inactivated

Product identification

Medicine name:
Bovilis BVD Suspension for Injection for Cattle

Active substance:
Bovine viral diarrhoea virus 1, strain C-86 (cytopathic), Inactivated

Target species:
Cattle
Cattle (for meat production)
Cattle (dairy cow)
Cattle (calf)
Cattle (heifer)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Bovine viral diarrhoea virus 1, strain C-86 (cytopathic), Inactivated

50.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Cattle (for meat production)**
  - Meat and offal. 0 day
- **Cattle (dairy cow)**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Cattle (calf)**
  - Meat and offal. 0 day
- **Cattle (heifer)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
(ID7) 100 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 100 millilitre(s)
(ID6) 50 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 50 millilitre(s)
(ID1) 10 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 10 millilitre(s)
(ID20) 2 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 2 millilitre(s)
(ID30) 2 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 2 millilitre(s)
(ID3) 20 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 20 millilitre(s)
(ID9) 250 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 250 millilitre(s)
(ID4) 20 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 20 millilitre(s)
(ID2) 10 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 10 millilitre(s)
(ID5) 50 millilitre(s): Box (Cardboard) with 1 Bottle (Glass) with 50 millilitre(s)
(ID8) 100 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 100 millilitre(s)
(ID10) 250 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:

25/06/1999

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 06376/3026

Date of authorisation status change:
10/06/2024

Reference member state:
Netherlands

Procedure number:
NL/V/0433/001

Concerned member states:
Austria Belgium France Germany Greece Ireland Italy Luxembourg Poland Portugal Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

Summary of Product Characteristics