Bovilis BVD Suspension for injection for cattle

Not authorised

- Bovine viral diarrhoea virus 1, strain C-86, Inactivated

Product identification

Medicine name:
Bovilis BVD Suspension for injection for cattle
Bovilis BVD injektionsvæske, suspension

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

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

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Bovine viral diarrhoea virus 1, strain C-86, 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 cattle)**
  - Meat and offal. 0 day
  - Milk. 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:
Surrendered

Authorised in:
Denmark

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
3/08/2007

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
39194

Date of authorisation status change:
24/07/2023

Reference member state:
Germany

Procedure number:
DE/V/0211/001

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 28/01/2022
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