

Bovilis BVD Suspension for injection for cattle

Authorised

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

Product identification

Medicine name:

Bovilis BVD Suspension for injection for cattle

BOVILIS BVD

Active substance:

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

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Cattle

- Milk. no withdrawal period zero days
- Meat and offal. no withdrawal period zero days

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Cattle (for meat production)

- Meat and offal. no withdrawal period zero days

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Cattle (dairy cow)

- Milk. no withdrawal period zero days
- Meat and offal. no withdrawal period zero days

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Cattle (calf)

- Meat and offal. no withdrawal period zero days

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Cattle (heifer)

- Meat and offal. no withdrawal period zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Available in:

Italy

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:

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

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

27/10/1999

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Ministry Of Health

Authorisation number:

102883

Date of authorisation status change:

27/10/1999

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

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.