

# 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

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**Active substance:**

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

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**Target species:**

Cattle

Cattle (for meat production)

Cattle (dairy cow)

Cattle (calf)

Cattle (heifer)

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**Route of administration:**

Intramuscular use

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## 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)

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**Pharmaceutical form:**

Suspension for injection

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**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 ays

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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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI02AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Slovenia

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

5/11/2007

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

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**Authorisation number:**

MR/V/0043/001

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**Date of authorisation status change:**

5/11/2007

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0433/001

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**Concerned member states:**

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

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 28/01/2022

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## Labelling

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

## Package Leaflet

English (PDF)

Published on: 13/03/2026

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