

# Bovilis Bovipast suspensie voor injectie voor runderen

Authorised

- Mannheimia haemolytica, serotype A1, strain M4/1, Inactivated
- Bovine respiratory syncytial virus, strain EV 908, Inactivated
- Bovine parainfluenza virus 3, strain SF-4, Inactivated

## Product identification

**Medicine name:**

Bovilis Bovipast suspensie voor injectie voor runderen

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**Active substance:**

Mannheimia haemolytica, serotype A1, strain M4/1, Inactivated

Bovine respiratory syncytial virus, strain EV 908, Inactivated

Bovine parainfluenza virus 3, strain SF-4, Inactivated

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**Target species:**

Cattle

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Mannheimia haemolytica, serotype A1, strain M4/1, Inactivated  
9.00 log10 unit(s) / 5.00 millilitre(s)

Bovine respiratory syncytial virus, strain EV 908, Inactivated  
1.00 virus neutralising unit(s) / 5.00 millilitre(s)

Bovine parainfluenza virus 3, strain SF-4, Inactivated  
1.00 haemagglutination inhibiting unit(s) / 5.00 millilitre(s)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Cattle**

- Milk. no withdrawal period nul dagen
  - Milk. no withdrawal period nul dagen
  - Meat and offal. no withdrawal period nul dagen
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI02AL04

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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:**

Netherlands

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**Available in:**

Netherlands

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**Package description:**

Available only in Dutch

Available only in Dutch

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Intervet Nederland B.V.

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**Marketing authorisation date:**

17/06/1999

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**Manufacturing sites for batch release:**

INTERVET INTERNATIONAL B.V.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 9260

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**Date of authorisation status change:**

1/07/2020

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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

Combined File of all Documents

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