

# CATTLEMASTER 4

Authorised

- Bovine viral diarrhoea virus 1, strain 5960, Inactivated
- Bovine parainfluenza virus 3, strain RLB103, Live
- Bovine viral diarrhoea virus 1, strain 6309, Inactivated
- Bovine respiratory syncytial virus, strain 375, Live
- Bovine herpesvirus 1, strain RLB 106, Live

## Product identification

**Medicine name:**

CATTLEMASTER 4

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**Active substance:**

Bovine viral diarrhoea virus 1, strain 5960, Inactivated

Bovine parainfluenza virus 3, strain RLB103, Live

Bovine viral diarrhoea virus 1, strain 6309, Inactivated

Bovine respiratory syncytial virus, strain 375, Live

Bovine herpesvirus 1, strain RLB 106, Live

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**Target species:**

Cattle

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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 5960, Inactivated  
0.31 millilitre(s) / 1.00 unit(s)

Bovine parainfluenza virus 3, strain RLB103, Live  
5.00 log<sub>10</sub> 50% cell culture infectious dose / 1.00 unit(s)

Bovine viral diarrhoea virus 1, strain 6309, Inactivated  
0.62 millilitre(s) / 1.00 unit(s)

Bovine respiratory syncytial virus, strain 375, Live  
4.10 log<sub>10</sub> 50% cell culture infectious dose / 1.00 unit(s)

Bovine herpesvirus 1, strain RLB 106, Live  
5.70 log<sub>10</sub> 50% cell culture infectious dose / 1.00 unit(s)

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### **Pharmaceutical form:**

Lyophilisate and solvent for suspension for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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#### **Cattle**

- Milk. 0 hour
  - Meat and offal. 0 hour
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### **Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI02AH

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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:**

Italy

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**Available in:**

Italy

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**Package description:**

Available only in [Italian](#)

Available only in [Italian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Italia S.r.l.

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**Marketing authorisation date:**

21/04/1995

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**Manufacturing sites for batch release:**

Zoetis Belgium SA

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

21/04/1995

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## Documents

Combined File of all Documents

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