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

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

Target species:
Cattle

Route of administration:
Intramuscular use

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 log10 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 log10 50% cell culture infectious dose / 1.00 unit(s)

Bovine herpesvirus 1, strain RLB 106, Live
5.70 log10 50% cell culture infectious dose / 1.00 unit(s)

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AH

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Italy

Available in:
Italy

Package description:
Available only in Italian
Available only in Italian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Italia S.r.l.

Marketing authorisation date:
21/04/1995

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
21/04/1995

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

Documents

Combined File of all Documents

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