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CATTLEMASTER 4

Authorised

- Bovine viral diarrhoea virus 1, strain 5960 (cythopathic), Inactivated
- Bovine parainfluenza virus 3, strain RLB103, Live
- Bovine viral diarrhoea virus 1, strain 6309 (non-cytopathic), 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 (cythopathic), Inactivated

Bovine parainfluenza virus 3, strain RLB103, Live

Bovine viral diarrhoea virus 1, strain 6309 (non-cytopathic), 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 (cythopathic), 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 (non-cytopathic), 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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