CattleMaster 4

Authorised

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

Product identification

Medicine name:
CattleMaster 4

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

Target species:
Cattle
Cattle

Route of administration:
Intramuscular use
Intramuscular use

Product details

Active substance and strength:

Bovine viral diarrhoea virus 1, strain 6309, Inactivated
2.00 log2 virus neutralising unit(s) / 2.00 millilitre(s)

Bovine viral diarrhoea virus 1, strain 5960, Inactivated
2.00 log2 virus neutralising unit(s) / 2.00 millilitre(s)

Bovine respiratory syncytial virus, strain 375, Live
4.10 log10 50% cell culture infectious dose / 2.00 millilitre(s)

Bovine parainfluenza virus 3, strain RLB103, Live
5.00 log10 50% cell culture infectious dose / 2.00 millilitre(s)

Bovine herpesvirus 1, strain RLB 106, Live
5.70 log10 50% cell culture infectious dose / 2.00 millilitre(s)

Pharmaceutical form:

Lyophilisate and suspension for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AH

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Zoetis Belgium

Marketing authorisation date:
31/07/2007

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
130157

Date of authorisation status change:
26/10/2023

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

Documents

Summary of Product Characteristics

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