Rispoval Pasteurella, lyofilisaat en suspendeervloeistof voor emulsie voor injectie voor runderen

Authorised

- Mannheimia haemolytica, serotype A1, strain NL 1009, capsular antigen
- Mannheimia haemolytica, serotype A1, strain NL 1009, leucotoxoid

Product identification

Medicine name:
Rispoval Pasteurella, lyofilisaat en suspendeervloeistof voor emulsie voor injectie voor runderen

Active substance:
Mannheimia haemolytica, serotype A1, strain NL 1009, capsular antigen
Mannheimia haemolytica, serotype A1, strain NL 1009, leucotoxoid

Target species:
Cattle

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Mannheimia haemolytica, serotype A1, strain NL 1009, capsular antigen
200.00 relative unit(s) / 2.00 millilitre(s)

Mannheimia haemolytica, serotype A1, strain NL 1009, leucotoxoid
345.00 relative unit(s) / 2.00 millilitre(s)

Pharmaceutical form:
Lyophilisate for suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. no withdrawal period nul dagen

Subcutaneous use:

- **Cattle**
  - Meat and offal. no withdrawal period nul dagen

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AB04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
5 doses of vaccine and solvent in glass vials (type I) with a rubber stopper and an aluminum cap (1 dose of 2 ml)
25 doses of vaccine and solvent in glass vials (type I) with a rubber stopper and an aluminum cap (1 dose of 2 ml)

1 dose of vaccine and solvent in glass vials (type I) with a rubber stopper and an aluminum cap (1 dose of 2 ml)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Zoetis B.V.

Marketing authorisation date:
28/04/1997

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 8661

Date of authorisation status change:
9/04/2018

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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