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Rispoval Lactovac C vakcina A.U.V.

Not authorised

- Escherichia coli, serotype O9:K35 (fimbrial adhesin F5 and F41), strain S1091/83, Inactivated
- Bovine coronavirus, strain 800, Inactivated
- Bovine rotavirus A, strain Holland, Inactivated
- Bovine rotavirus A, strain 1005/78, Inactivated

Product identification

Medicine name:
Rispoval Lactovac C vakcina A.U.V.

Active substance:
Escherichia coli, serotype O9:K35 (fimbrial adhesin F5 and F41), strain S1091/83, Inactivated

Bovine coronavirus, strain 800, Inactivated

Bovine rotavirus A, strain Holland, Inactivated

Bovine rotavirus A, strain 1005/78, Inactivated

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Escherichia coli, serotype O9:K35 (fimbrial adhesin F5 and F41), strain S1091/83, Inactivated
250.00 haemagglutinating units / 1.00 unit(s)/dose

Bovine coronavirus, strain 800, Inactivated
5.80 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

Bovine rotavirus A, strain Holland, Inactivated
7.00 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

Bovine rotavirus A, strain 1005/78, Inactivated
7.40 log10 50% tissue culture infectious dose / 1.00 unit(s)/dose

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Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Hungary

Package description:

Available only in Hungarian

Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Hungary Kft.

Marketing authorisation date:
25/02/1998

Manufacturing sites for batch release:
Zoetis Belgium SA

Responsible authority:
Directorate Of Veterinary Medicinal Products

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

Date of authorisation status change:
25/02/1998

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