

KOLIBIN RC NEO, emulsione oleosa iniettabile per bovini

Authorised

- Escherichia coli, serotype O101:K30 (fimbrial adhesin F5), Inactivated
- Bovine coronavirus, strain C-197, Inactivated
- Escherichia coli, serotype O9:K35 (fimbrial adhesin F5), Inactivated
- Bovine rotavirus A, type G6P1, strain TM-91, Inactivated
- Escherichia coli, serotype O8:K35 (fimbrial adhesin F5), Inactivated

Product identification

Medicine name:

KOLIBIN RC NEO, emulsione oleosa iniettabile per bovini

Active substance:

Escherichia coli, serotype O101:K30 (fimbrial adhesin F5), Inactivated

Bovine coronavirus, strain C-197, Inactivated

Escherichia coli, serotype O9:K35 (fimbrial adhesin F5), Inactivated

Bovine rotavirus A, type G6P1, strain TM-91, Inactivated

Escherichia coli, serotype O8:K35 (fimbrial adhesin F5), Inactivated

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Escherichia coli, serotype O101:K30 (fimbrial adhesin F5), Inactivated

1.00 relative potency / 1.00 Dose

Bovine coronavirus, strain C-197, Inactivated

1.00 relative potency / 1.00 Dose

Escherichia coli, serotype O9:K35 (fimbrial adhesin F5), Inactivated

1.00 relative potency / 1.00 Dose

Bovine rotavirus A, type G6P1, strain TM-91, Inactivated

1.00 relative potency / 1.00 Dose

Escherichia coli, serotype O8:K35 (fimbrial adhesin F5), Inactivated

1.00 relative potency / 1.00 Dose

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

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

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Cattle

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

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

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:

Bio 98 S.r.l.

Marketing authorisation date:

20/10/2015

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

20/10/2015

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.