KOLIBIN RC NEO, injekcinė emulsija galvijams

Authorised

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

Product identification

Medicine name:
KOLIBIN RC NEO, injekcinė emulsija galvijams

Active substance:
Bovine rotavirus A, type G6P1, strain TM-91, Inactivated

Bovine coronavirus, strain C-197, Inactivated

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

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

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

Target species:
Cattle (heifer)
Cattle (pregnant cow)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Bovine rotavirus A, type G6P1, strain TM-91, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Bovine coronavirus, strain C-197, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Escherichia coli, serotype O8:K35 (fimbrial adhesin F5), Inactivated
1.00 relative potency / 2.00 millilitre(s)

Escherichia coli, serotype O9:K35 (fimbrial adhesin F5), Inactivated
1.00 relative potency / 2.00 millilitre(s)

Escherichia coli, serotype O101:K30 (fimbrial adhesin F5), Inactivated
1.00 relative potency / 2.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI02AL01

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

Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
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Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetmarket UAB

Marketing authorisation date:
20/12/2021

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/21/2692/001-004

Date of authorisation status change:
20/12/2021

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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