Rokovac Neo Plus

Authorised

- Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), Inactivated
- Escherichia coli, serotype K85:987P (fimbrial adhesin F6), Inactivated
- Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), Inactivated
- Escherichia coli, serotype O147:K88 (fimbrial adhesin F4), Inactivated
- Porcine rotavirus A, strain OSU 6, Inactivated

Product identification

Medicine name:
Rokovac Neo Plus
Роковак Нео Плюс

Active substance:
Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), Inactivated
Escherichia coli, serotype K85:987P (fimbrial adhesin F6), Inactivated
Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), Inactivated
Escherichia coli, serotype O147:K88 (fimbrial adhesin F4), Inactivated
Porcine rotavirus A, strain OSU 6, Inactivated

Target species:
Pig (pregnant sow)
Pig (young female)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), Inactivated
1.00 relative potency / 1.00 Dose

Escherichia coli, serotype K85:987P (fimbrial adhesin F6), Inactivated
1.00 relative potency / 1.00 Dose

Escherichia coli, serotype O101:K99 (fimbrial adhesin F5), Inactivated
1.00 relative potency / 1.00 Dose

Escherichia coli, serotype O147:K88 (fimbrial adhesin F4), Inactivated
1.00 relative potency / 1.00 Dose

Porcine rotavirus A, strain OSU 6, Inactivated
1.00 relative potency / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AL02

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
14/10/2020

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-3016

Date of authorisation status change:
28/05/2023

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

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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