

Castorex NEO suspension for injection for rabbits

Authorised

- Rabbit haemorrhagic disease virus, type 2, strain F/12B, Inactivated

Product identification

Medicine name:

Castorex NEO suspension for injection for rabbits

Active substance:

Rabbit haemorrhagic disease virus, type 2, strain F/12B, Inactivated

Target species:

Rabbit

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Rabbit haemorrhagic disease virus, type 2, strain F/12B, Inactivated
0.30 enzyme-linked immunosorbent assay unit / 500.00 microlitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Subcutaneous use:**

-

Rabbit

- All relevant tissues. 0 day
zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI08AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Paper carton containing glass vial made of neutral borosilicate glass with high hydrolytic resistance (Type 1) closed with a rubber plug suitable for parenteral preparations and an aluminium cap. (1x40 doses)

Paper carton containing glass vial made of neutral borosilicate glass with high hydrolytic resistance (Type 1) closed with a rubber plug suitable for parenteral preparations and an aluminium cap. (1x20 doses)

Paper carton containing glass vial made of neutral borosilicate glass with high hydrolytic resistance (Type 1) closed with a rubber plug suitable for parenteral preparations and an aluminium cap. (1x10 doses)

Paper carton containing glass vial made of neutral borosilicate glass with high hydrolytic resistance (Type 1) closed with a rubber plug suitable for parenteral preparations and an aluminium cap. (10x1 dose)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Pharmagal Bio spol. s r.o.

Marketing authorisation date:

31/03/2020

Manufacturing sites for batch release:

Pharmagal Bio spol. s r.o.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/005/DC/20-S

Date of authorisation status change:

31/03/2020

Reference member state:

Slovakia

Procedure number:

SK/V/0109/001

Concerned member states:

Czechia Hungary Malta Poland

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

Documents

Combined File of all Documents

English (PDF)

Published on: 15/12/2021

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