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:

Poland

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:
30/09/2020

Manufacturing sites for batch release:
Pharmagal Bio spol. s r.o.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3008

Date of authorisation status change:
30/09/2020

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

Summary of Product Characteristics

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

Labelling

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

Package Leaflet

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