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:

Malta

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:
24/02/2020

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

Responsible authority:
Ministry For Agriculture Fisheries And Animal Rights

Authorisation number:
VMA86

Date of authorisation status change:
24/02/2020

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