**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000035635

# 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 Castorex NEO Zawiesina do wstrzykiwań

#### **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:**

# 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 <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>