File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/600000041875

PESTORIN MORMYX liofilizāts un suspensija suspensijas injekcijām pagatavošanai trušiem

Authorised

- Myxoma virus, strain CAMP V-219, Live
- Rabbit haemorrhagic disease virus, type 1, strain CAMP V-351, Inactivated

Product identification

Medicine name:
PESTORIN MORMYX liofilizāts un suspensija suspensijas injekcijām pagatavošanai trušiem

Active substance:
Myxoma virus, strain CAMP V-219, Live

Rabbit haemorrhagic disease virus, type 1, strain CAMP V-351, Inactivated

Target species:
Rabbit

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Myxoma virus, strain CAMP V-219, Live
630957.00 50% tissue culture infectious dose / 1.00 unit(s)

Rabbit haemorrhagic disease virus, type 1, strain CAMP V-351, Inactivated
1024.00 unit(s) / 1.00 unit(s)

Pharmaceutical form:
Lyophilisate and suspension for suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Rabbit**
  - Meat and offal. 7 day

Iespējama lokāla audu reakcija, kura izzūd 7 dienu laikā.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI08AH01

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian

Available only in Latvian
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
24/05/2001

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/01/1335

Date of authorisation status change:
24/05/2001

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

Documents

Combined File of all Documents

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