

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](#)

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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

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