

PESTORIN MORMYX

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

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

3.30 log₁₀ 50% cell culture infectious dose / 1.00 millilitre(s)

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

7.00 log₂ haemagglutination inhibiting unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Lyophilisate for suspension for injection

Withdrawal period by route of administration:**Subcutaneous use:**

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Rabbit

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI08AH01

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Romania

Package description:

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

Available only in Romanian

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Available only in Romanian

Available only in Romanian

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Bioveta a.s.

Marketing authorisation date:

25/06/2007

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

120311

Date of authorisation status change:

2/09/2025

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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