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MYXOREN lyofilizát a rozpúšťadlo na injekčnú suspenziu

Authorised

- Myxoma virus, strain CAMP V-219, Live
- Myxoma virus, strain CAMP V-219, Live
- Myxoma virus, strain CAMP V-219, Live

Product identification

Medicine name:
MYXOREN lyofilizát a rozpúšťadlo na injekčnú suspenziu

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

Target species:
Rabbit
Rabbit
Rabbit

Route of administration:
Transdermal use
Intradermal use
Subcutaneous use

Product details

Active substance and strength:
Myxoma virus, strain CAMP V-219, Live
2.60 log10 50% tissue culture infectious dose / 1.00 Dose

Myxoma virus, strain CAMP V-219, Live
2.70 log10 50% tissue culture infectious dose / 1.00 Dose

Myxoma virus, strain CAMP V-219, Live
3.30 log10 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Transdermal use:

- **Rabbit**
  - All relevant tissues. 0 day

Intradermal use:

- **Rabbit**
  - All relevant tissues. 0 day

Subcutaneous use:

- **Rabbit**
  - Adipose tissue. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI08AD02

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

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Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak

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:
Bioveta a.s.

Marketing authorisation date:
2/06/2005

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/191/91-S

Date of authorisation status change:
2/06/2005

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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