File downloaded on 2026-11-06
Source URL: https://medicines.health.europa.eu/veterinary/en/700000137866

MYXOREN, liofilizatas ir skiediklis injekcinei suspensijai

Authorised

- Myxoma virus, strain CAMP V-219, Live

Product identification

Medicine name:
MYXOREN, liofilizatas ir skiediklis injekcinei suspensijai

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

Target species:
Rabbit

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Myxoma virus, strain CAMP V-219, Live
125893.00 50% tissue culture infectious dose / 1.00 dose

Pharmaceutical form:
Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Rabbit**
  - Not specified. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI08AD02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Lithuania

Package description:

Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian
Available only in Lithuanian

Available only in Lithuanian

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:
19/06/2002

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/02/1428/001-019

Date of authorisation status change:
19/06/2007

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.