RABISIN MULTI, injekcinė suspensija

Authorised

- Rabies virus, strain G52, Inactivated

Product identification

Medicine name:
RABISIN MULTI, injekcinė suspensija

Active substance:
Rabies virus, strain G52, Inactivated

Target species:
Dog
Cat
Cattle
Sheep
Ferret
Horse

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Rabies virus, strain G52, Inactivated
1.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day

Intramuscular use:

- **Horse**
  - Meat and offal. 0 day
- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Lithuania

Package description:
Available only in Lithuanian

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:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
4/12/1995

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/11/2029/001

Date of authorisation status change:
29/02/2016

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

Documents

RV2029.pdf