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RABISIN, ≥ 2,09 log10 OD50 i ≥ 1 I.J., suspenzija za injekciju za pse, mačke, pitome vretice, konje, govedo i ovce

Authorised

- Rabies virus, strain G52, Inactivated

Product identification

Medicine name:
RABISIN, ≥ 2,09 log10 OD50 i ≥ 1 I.J., suspenzija za injekciju za pse, mačke, pitome vretice, konje, govedo i ovce

Active substance:
Rabies virus, strain G52, Inactivated

Target species:
Dog
Cat
Ferret
Cattle
Sheep
Horse

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Rabies virus, strain G52, Inactivated
2.09 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:
Croatia

Available in:
Croatia

Package description:
Available only in Croatian
Available only in Croatian
Available only in Croatian

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:
19/10/2018

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Croatian Veterinary Institute

Authorisation number:
UP/I-322-05/16-01/490

Date of authorisation status change:
17/01/2025

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

Documents

Summary of Product Characteristics

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