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RABISIN injekčná suspenzia

Authorised

- Rabies virus, strain G52, Inactivated

Product identification

Medicine name:
RABISIN injekčná suspenzia

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
1.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**
  - All relevant tissues. 0 day — not applicable
- **Cat**
  - All relevant tissues. 0 day — not applicable
- **Ferret**
  - All relevant tissues. 0 day — not applicable
- **Cattle**
  - All relevant tissues. 0 day — zero days
- **Sheep**
  - All relevant tissues. 0 day — zero days

Intramuscular use:

- **Dog**
  - All relevant tissues. 0 day — not applicable
- **Cat**
  - All relevant tissues. 0 day — not applicable
-

Ferret

- All relevant tissues. 0 day — not applicable

- **Cattle**
  - All relevant tissues. 0 day — zero days

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

- **Sheep**
  - All relevant tissues. 0 day — zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI07AA02

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

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
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:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
14/11/2004

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/242/92-S

Date of authorisation status change:
14/11/2004

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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