

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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