

RABISIN suspenzija za injiciranje

Authorised

- Rabies virus, strain G52, Inactivated

Product identification

Medicine name:

RABISIN suspenzija za injiciranje

Active substance:

Rabies virus, strain G52, Inactivated

Target species:

Cattle
Sheep
Horse
Mink
Dog
Cat

Route of administration:

Subcutaneous use
Subcutaneous 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, milk. 0 day

Meso in organi živali namenjenih za prehrano ljudi: nič dni. Mleko živali, namenjeno za prehrano ljudi: nič dni.

-

Sheep

- Meat and offal, milk. 0 day

Meso in organi živali namenjenih za prehrano ljudi: nič dni. Mleko živali, namenjeno za prehrano ljudi: nič dni.

-

Horse

- Meat and offal, milk. 0 day

Meso in organi živali namenjenih za prehrano ljudi: nič dni. Mleko živali, namenjeno za prehrano ljudi: nič dni.

Subcutaneous use:

-

Cattle

- Meat and offal, milk. 0 day

Meso in organi živali namenjenih za prehrano ljudi: nič dni. Mleko živali, namenjeno za prehrano ljudi: nič dni.

-

Sheep

- Meat and offal, milk. 0 day

Meso in organi živali namenjenih za prehrano ljudi: nič dni. Mleko živali, namenjeno za prehrano ljudi: nič dni.

•

Horse

- Meat and offal, milk. 0 day

Meso in organi živali namenjenih za prehrano ljudi: nič dni. Mleko živali, namenjeno za prehrano ljudi: nič dni.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Package description:

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Available only in [Slovenian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

30/12/1999

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

NP/V/0290/001

Date of authorisation status change:

30/12/1999

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.