

Nobivac Rabies Zawiesina do wstrzykiwań

Authorised

- Rabies virus, strain Pasteur RIV, Inactivated

Product identification

Medicine name:

Nobivac Rabies Zawiesina do wstrzykiwań

Active substance:

Rabies virus, strain Pasteur RIV, Inactivated

Target species:

Cattle

Sheep

Goat

Horse

Cat

Ferret

Fox

Dog

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Rabies virus, strain Pasteur RIV, Inactivated
2.00 international unit(s) / 1.00 international unit(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cattle

- Meat and offal. 0 day

-

Sheep

- Meat and offal. 0 day

-

Goat

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

-

Ferret

- All relevant tissues. no withdrawal period Not applicable.

-

Fox

- All relevant tissues. no withdrawal period Not applicable.

Intramuscular use:

-

Cattle

- Meat and offal. 0 day

-

Sheep

- Meat and offal. 0 day

-

Goat

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

-

Ferret

- All relevant tissues. no withdrawal period Not applicable.

-

Fox

- All relevant tissues. no withdrawal period Not applicable.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Available in:

Poland

Package description:

Available only in [Polish](#)

Available only in [Polish](#)

Available only in [Polish](#)

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:

Intervet International B.V.

Marketing authorisation date:

17/02/1998

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

0472

Date of authorisation status change:

17/02/1998

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

Documents

Labelling

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.

Summary of Product Characteristics

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