

Nobivac Rabies ενέσιμο

Authorised

εναιώρημα για βοοειδή, άλογα,
σκύλους, γάτες και *Mustela furo*
(‘φέρετ’ ή οικιακό βρωμοκούναβο
ή εξημερωμένο βρωμοκούναβο)

- Rabies virus, strain Pasteur RIV, Inactivated

Product identification

Medicine name:

Nobivac Rabies ενέσιμο εναιώρημα για βοοειδή, άλογα, σκύλους, γάτες και *Mustela furo* (‘φέρετ’ ή οικιακό βρωμοκούναβο ή εξημερωμένο βρωμοκούναβο)

Active substance:

Rabies virus, strain Pasteur RIV, Inactivated

Target species:

Cattle

Dog

Horse

Cat

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Rabies virus, strain Pasteur RIV, Inactivated
100.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cattle

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

Intramuscular use:

-

Cattle

- Meat and offal. 0 day

-

Horse

- Meat and offal. 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:

Cyprus

Available in:

Cyprus

Package description:

Available only in Greek

Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

2/11/1995

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

16314

Date of authorisation status change:

1/11/2010

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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