# Rabisin

Authorised

• Rabies virus, strain G52, Inactivated

# **Product identification**

#### **Medicine name:**

Rabisin

#### **Active substance:**

Rabies virus, strain G52, Inactivated

# **Target species:**

Cattle

Sheep

Horse

Ferret

Mink

Polecat

Dog

Cat

#### **Route of administration:**

Intramuscular use

Subcutaneous use

# **Product details**

# **Active substance and strength:**

Rabies virus, strain G52, Inactivated

# **Pharmaceutical form:**

Suspension for injection

# Withdrawal period by route of administration: Intramuscular use:

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#### Cattle

- Meat and offal. 0 day
- Milk. 0 hour

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# **Sheep**

- Meat and offal. 0 day
- Milk. 0 hour

•

## Horse

- Meat and offal. 0 day
- Milk. 0 hour

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#### **Ferret**

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#### Mink

•

# **Polecat**

## **Subcutaneous use:**

•

# **Cattle**

- Meat and offal. 0 day
- Milk. 0 hour

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# Dog

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# **Sheep**

- Meat and offal. 0 day
- Milk. 0 hour

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#### Cat

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#### **Ferret**

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#### Mink

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#### **Polecat**

# Anatomical therapeutic chemical veterinary (ATCvet) codes:

**QI07AA02** 

## Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

## **Authorisation status:**

Valid

#### **Authorised in:**

Ireland

### Package description:

Type I glass vials with butyl-elastomer closure.Package sizes:Bottle (glass) of 1 dose of suspension, box of 1 bottle

Type I glass vials with butyl-elastomer closure. Package sizes: Bottle (glass) of 1 dose of suspension, box of 10 bottles

Type I glass vials with butyl-elastomer closure. Package sizes: Bottle (glass) of 1 dose of suspension, box of 100 bottles

# 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 Vetmedica GmbH
Marketing authorisation date: 10/08/2004
Manufacturing sites for batch release: Boehringer Ingelheim Animal Health France
Responsible authority: Health Products Regulatory Authority
Authorisation number: VPA10454/074/001
Date of authorisation status change:

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

# **Documents**

Summary of Product Characteristics

**Source URL:** https://medicines.health.europa.eu/veterinary/600000080509