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RABISIN vet, injeksjonsvæske, suspensjon Vaksine mot rabies til dyr (inaktivert)

Authorised

- Rabies virus, Inactivated

Product identification

Medicine name:
RABISIN vet, injeksjonsvæske, suspensjon Vaksine mot rabies til dyr (inaktivert)

Active substance:
Rabies virus, Inactivated

Target species:
Horse
Dog
Cat
Cattle
Sheep

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Rabies virus, Inactivated
2.09 log10 optical density50 / 1.00 international unit(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Horse**
  - All relevant tissues. 0 day
- **Cattle**
  - All relevant tissues. 0 day
  - Milk. 0 day
- **Sheep**
  - All relevant tissues. 0 day

Subcutaneous use:

- **Cattle**
  - All relevant tissues. 0 day
  - Milk. 0 day
- **Horse**
  - All relevant tissues. 0 day
- **Sheep**

- All relevant tissues. 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:
Norway

Available in:
Norway

Package description:
Available only in Norwegian

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 Denmark A/S

Marketing authorisation date:
22/10/2004

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
1997-00851

Date of authorisation status change:
27/08/2009

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

Documents

Summary of Product Characteristics

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Package Leaflet

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