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Fatrovax RHD (--) - Suspension for injection

Authorised

- Rabbit haemorrhagic disease virus, type 1, capsid protein VP60a
- Rabbit haemorrhagic disease virus, type 2, capsid protein VP60ab

Product identification

Medicine name:
Fatrovax RHD (--) - Suspension for injection

Active substance:
Rabbit haemorrhagic disease virus, type 1, capsid protein VP60a

Rabbit haemorrhagic disease virus, type 2, capsid protein VP60ab

Target species:
Rabbit

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Rabbit haemorrhagic disease virus, type 1, capsid protein VP60a

Presentation_strength:≥1 RP Index:2

Rabbit haemorrhagic disease virus, type 2, capsid protein VP60ab

Presentation_strength:≥1 RP Index:3

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Rabbit**
  - All relevant tissues. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI08AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Bottle (PP), Package_size:1 bottle, Content:100 ml (200 doses)

Packaging:Bottle (PP), Package_size:1 bottle, Content:25 ml (50 doses)

Packaging:Syringe (glass), Package_size:5 pre-filled syringes of 1 dose, Content:0.5 ml (1 dose)

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:
Fatro S.p.A

Marketing authorisation date:
16/08/2021

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
16/08/2021

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

Documents

Combined File of all Documents

English (PDF)
Published on: 26/08/2024
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