

# 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

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**Active substance:**

Rabbit haemorrhagic disease virus, type 1, capsid protein VP60a

Rabbit haemorrhagic disease virus, type 2, capsid protein VP60ab

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**Target species:**

Rabbit

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**Route of administration:**

Subcutaneous use

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## 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

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Rabbit**

- All relevant tissues. 0 day  
Zero days

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI08AA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Fatro S.p.A

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**Marketing authorisation date:**

16/08/2021

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

16/08/2021

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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