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FILAVAC VHD K C+V SUSPENSION FOR INJECTION FOR RABBITS

Not authorised

- Rabbit haemorrhagic disease virus, type 1, strain IM.507.SC.2011, Inactivated
- Rabbit haemorrhagic disease virus, type 2, strain LP.SV.2012, Inactivated

Product identification

Medicine name:
FILAVAC VHD K C+V SUSPENSION FOR INJECTION FOR RABBITS
Filavac VHD K C+V, PD90, Injekční suspenze

Active substance:
Rabbit haemorrhagic disease virus, type 1, strain IM.507.SC.2011, Inactivated
Rabbit haemorrhagic disease virus, type 2, strain LP.SV.2012, Inactivated

Target species:
Rabbit

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Rabbit haemorrhagic disease virus, type 1, strain IM.507.SC.2011, Inactivated
1.00 90% protective dose / 1.00 Dose

Rabbit haemorrhagic disease virus, type 2, strain LP.SV.2012, Inactivated
1.00 90% protective dose / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Rabbit**
  - All relevant tissues. no withdrawal period — Withdrawal period is 0 days

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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI08AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Surrendered

Authorised in:

Czechia

Package description:

Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Filavie

Marketing authorisation date:
15/03/2018

Manufacturing sites for batch release:
Filavie

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/009/18-C

Date of authorisation status change:
24/06/2025

Reference member state:
France

Procedure number:
FR/V/0315/001

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.