

Eravac (--)- Emulsion for injection

Authorised

- Rabbit haemorrhagic disease virus, type 2, strain V-1037, Inactivated

Product identification

Medicine name:

Eravac (--)- Emulsion for injection

Active substance:

Rabbit haemorrhagic disease virus, type 2, strain V-1037, Inactivated

Target species:

Rabbit

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Rabbit haemorrhagic disease virus, type 2, strain V-1037, Inactivated

Presentation_strength: $\geq 70\%$ Reference:Hse Index:0

Pharmaceutical form:

Emulsion 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:Vial (HDPE), Package_size:1 vial, Content:100 ml (200 doses)

Packaging:Vial (glass), Package_size:10 vials, Content:0.5 ml (1 dose)

Packaging:Vial (glass), Package_size:1 vial, Content:5 ml (10 doses)

Packaging:Vial (glass), Package_size:1 vial, Content:20 ml (40 doses)

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:

Laboratorios Hipra, S.A.

Marketing authorisation date:

22/09/2016

Manufacturing sites for batch release:

Laboratorios Hipra S.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: 8/01/2024

[Download](#)

ema-puar-eravac-v-4239-par-en.pdf

ema-puar-eravac-v-4239-var-ii-0005-g-en.pdf

ema-puar-eravac-v-4239-var-ii-0003-g-en.pdf