File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/600000001499

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:≥ 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 (glass), Package_size:1 vial, Content:20 ml (40 doses)
Packaging:Vial (glass), Package_size:1 vial, Content:5 ml (10 doses)
Packaging:Vial (glass), Package_size:10 vials, Content:0.5 ml (1 dose)
Packaging:Vial (HDPE), Package_size:1 vial, Content:100 ml (200 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:
22/09/2016

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: 30/10/2026
Download

ema-puar-v4239-eravac-par-wepar-2026-10-23-en.pdf

ema-puar-v4239-eravac-var-ii-0003-g-wepar-2026-10-23-en.pdf

ema-puar-v4239-eravac-var-ii-0005-g-wepar-2026-10-23-en.pdf

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