

CUNIPRAVAC-RHD ΕΝΕΣΙΜΟ ΓΑΛΑΚΤΩΜΑ

Not
authorised

- Rabbit haemorrhagic disease virus, type 1, strain 3116-AP, Inactivated

Product identification

Medicine name:

CUNIPRAVAC-RHD ΕΝΕΣΙΜΟ ΓΑΛΑΚΤΩΜΑ

Active substance:

Rabbit haemorrhagic disease virus, type 1, strain 3116-AP, Inactivated

Target species:

Rabbit

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Rabbit haemorrhagic disease virus, type 1, strain 3116-AP, Inactivated
0.05 gram(s) / 0.50 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:**Subcutaneous use:**

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Rabbit

- Meat and offal. 21 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI08AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Greece

Package description:

Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone)

Marketing authorisation holder:

Laboratorios Hipra S.A.

Marketing authorisation date:

11/04/1994

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

34643/93/K-2629/12-04-1994/K-0083401

Date of authorisation status change:

29/05/2025

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