

YURVAC RHD (--) - Emulsion for injection

Authorised

- Rabbit haemorrhagic disease virus, type 2, capsid protein VP60

Product identification

Medicine name:

YURVAC RHD (--) - Emulsion for injection

Active substance:

Rabbit haemorrhagic disease virus, type 2, capsid protein VP60

Target species:

Rabbit

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Rabbit haemorrhagic disease virus, type 2, capsid protein VP60

Presentation_strength: ≥ 0.7 RP 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:

QI08AV

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:10 vials, Content:0.5 ml

Packaging:Vial (Glass), Package_size:1 vial, Content:5 ml

Packaging:Vial (PET), Package_size:1 vial, Content:100 ml

Packaging:Vial (PET), Package_size:1 vial, Content:20 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - new active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Laboratorios Hipra, S.A.

Marketing authorisation date:

11/09/2023

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:

11/09/2023

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

Documents

Combined File of all Documents

English (PDF)

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