

Regumate Equine 2.2 mg/ml oral solution for horses

Not
authorised

- Altrenogest

Product identification

Medicine name:

Regumate Equine 2.2 mg/ml oral solution for horses

Regumate Equine 2,2 mg/ml belsőleges oldat lovaknak A.U.V.

Active substance:

Altrenogest

Target species:

Horse

Route of administration:

Oral use

Product details

Active substance and strength:

Altrenogest

2.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

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

-

Horse

- Meat and offal. 9 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:QG03DX90

Legal status of supply:Veterinary medicinal product subject to veterinary prescription

Authorisation status:Surrendered

Authorised in:Hungary

Package description:

The product is packaged in brown opaque 150 ml polyethylene bottles sealed by an aluminium protective cap and closed by a plastic screwed-on stopper. The 150 ml is provided with a luer lock cap which, when screwed on the bottle neck, allows the user to safely and accurately withdraw the product with a syringe that can directly adjust on the luer lock cap.

The product is packaged in brown opaque 250 ml polyethylene bottles sealed by an aluminium protective cap and closed by a plastic screwed-on stopper. The 250 ml bottle is equipped with a 12.5 measuring compartment.

The product is packaged in a brown opaque 300 ml polyethylene bottle sealed by an aluminium protective cap and closed by a plastic screwed-on stopper. The 300 ml bottle is provided with a luer lock cap which, when screwed on the bottle neck, allows the user to safely and accurately withdraw the product with a syringe that can directly adjust on the luer lock cap.

The product is packaged in a brown opaque 1000 ml polyethylene bottle sealed by an aluminium protective cap and closed by a plastic screwed-on stopper. The 1000 ml bottle is provided with a luer lock cap which, when screwed on the bottle neck, allows the user to safely and accurately withdraw the product with a syringe that can directly adjust on the luer lock cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

2/08/2005

Manufacturing sites for batch release:

Intervet Productions S.A.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

2664/X/10 MgSzH ÁTI

Date of authorisation status change:

28/09/2023

Reference member state:

Ireland

Procedure number:

IE/V/0155/001

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

Documents

Summary of Product Characteristics

English (PDF)

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