Regumate Equine 2.2 mg/ml oral solution for horses

Authorised

- Altrenogest

Product identification

Medicine name:
Regumate Equine 2.2 mg/ml oral solution for horses

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

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Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

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.

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 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 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.

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 (Ireland) Limited

Marketing authorisation date:
1/10/1997

Manufacturing sites for batch release:
Intervet Productions S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/124/001

Date of authorisation status change:
1/10/1997

Reference member state:
Ireland

Procedure number:
IE/V/0155/001

Concerned member states:
Austria Belgium Czechia Denmark Finland France Germany Greece Hungary Italy Luxembourg Netherlands Norway Portugal Slovakia United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents