

# 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

REGUMATE EQUIN 2,2 MG/ML SOLUTION BUVABLE POUR CHEVAUX

**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:**

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**Horse**

- Meat and offal. 9 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QG03DX90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Intervet

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**Marketing authorisation date:**

3/12/2004

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**Manufacturing sites for batch release:**

Intervet Productions S.A.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/8393965 3/2004

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**Date of authorisation status change:**

3/12/2009

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0155/001

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**Concerned member states:**

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

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## Documents

### Summary of Product Characteristics

English (PDF)

Published on: 3/11/2024

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### Package Leaflet and Labelling

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