# Genestran 75 micrograms/ml solution for injection for cattle, horses and pigs

Authorised

This information is not available for this product.

## Product identification

**Medicine name:**
Genestran 75 micrograms/ml solution for injection for cattle, horses and pigs
Genestran 75 microgramas/ml solução injetável para bovinos, equinos e suínos

**Active substance:**
This information is not available for this product.

**Target species:**
Cattle
Horse
Pig

**Route of administration:**
Intramuscular use

## Product details

**Active substance and strength:**
This information is not available for this product.

**Pharmaceutical form:**
Solution for injection

**Withdrawal period by route of administration:**

**Intramuscular use:**

- **Cattle**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 1 day

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

QG02AD90

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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:**

Portugal

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**Available in:**

Portugal

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**Package description:**

Colourless vial of type I glass containing 20 ml of solution for injection, with chlorobutyl rubber stopper and aluminium cap.Presentation: Cardboard box of 1 vial of 20 ml

Colourless vial of type I glass containing 50 ml of solution for injection, with chlorobutyl rubber stopper and aluminium cap.Presentation: Cardboard box of 1 vial of 50 ml

Colourless vial of type I glass containing 20 ml of solution for injection, with chlorobutyl rubber stopper and aluminium cap.Presentation: Cardboard box of 5 vials of 20 ml

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## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Well-established use application (Article 13a of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Animedica GmbH

**Marketing authorisation date:**
31/10/2011

**Manufacturing sites for batch release:**
Industrial Veterinaria S.A.
Animedica GmbH

**Responsible authority:**
Directorate General For Food And Veterinary

**Authorisation number:**
386/01/11RFVPT

**Date of authorisation status change:**
27/04/2022

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0228/001

**Concerned member states:**
Austria Belgium Czechia Estonia France Germany Iceland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain 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

This document does not exist in this language (English). You can find it in another language below.

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