

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

Authorised

- R-Cloprostenol sodium

Product identification

Medicine name:

Genestran 75 micrograms/ml solution for injection for cattle, horses and pigs
Genestran 75 mikrogramu/ml šķīdums injekcijām liellopiem, zirgiem un cūkām

Active substance:

R-Cloprostenol sodium

Target species:

Cattle
Horse
Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

R-Cloprostenol sodium
78.88 microgram(s) / 1.00 millilitre(s)

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
- Milk. 0 hour

-

Pig

- Meat and offal. 1 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

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

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:

23/12/2011

Manufacturing sites for batch release:

aniMedica GmbH

Industrial Veterinaria S.A.

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/MRP/11/0068

Date of authorisation status change:

25/12/2011

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.

Source URL: <https://medicines.health.europa.eu/veterinary/600000050187>