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, 75ug/ml, Injekční roztok

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD90

Legal status of supply:

This information is not available for this product.

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

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:
18/05/2011

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/043/11-C

Date of authorisation status change:
18/05/2011

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

English (PDF)
Published on: 11/02/2022
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Package Leaflet

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Labelling

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