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

Authorised

• R-Cloprostenol sodium

Product identification

Medicine name:

Galapan 75 micrograms/ml solution for injection for cattle, horses and pigs GALAPAN 75 microgramos/ml SOLUCIÓN INYECTABLE PARA BOVINO, CABALLOS Y PORCINO

Active substance:

R-Cloprostenol sodium

Target species:

Cattle

Horse

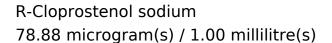
Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:



Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration: Intramuscular use:

Cattle

- Meat and offal. 1 day Meat and offal
- Milk. 0 hour

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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:

Spain

Package description:

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

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

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed Consent application (Article 21 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Industrial Veterinaria S.A.

Marketing authorisation date:

3/07/2025

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

Industrial Veterinaria S.A.

aniMedica GmbH

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

4422 ESP

Date of authorisation status change:

4/07/2025

Reference member state:

Ireland

Procedure number:

IE/V/0630/001
Concerned member states: Spain
Informed consent reference: 60000050164 60000050164
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet
Documents
Summary of Product Characteristics
English (PDF) Published on: 24/08/2025 Download
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Package Leaflet
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