

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:

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 Meat and offal
- Milk. 0 hour

-

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:

600000050164

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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

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Combined File of all Documents

Package Leaflet

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Labelling

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

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