Source URL: https://medicines.health.europa.eu/veterinary/en/600000038855

Dalmaprost 0.075 mg/ml solution for injection for cattle, pigs and horses

Authorised

R-Cloprostenol sodium

Product identification

Medicine name:

Dalmaprost 0.075 mg/ml solution for injection for cattle, pigs and horses Dalmazin SYNCH, 0,075 mg/mL, otopina za injekciju, za goveda, svinje i konje

Active substance:

R-Cloprostenol sodium

Target species:

Cattle (cow)

Pig (female)

Horse (mare)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

R-Cloprostenol sodium

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration: Intramuscular use:

Cattle (cow)

- Meat and offal. no withdrawal period withdrawal period is 0 days
- Milk. no withdrawal period withdrawal period is 0 hours

Pig (female)

- Meat and offal. 1 day

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Horse (mare)

- Meat and offal. 2 day
- Milk. no withdrawal period withdrawal period is 0 hours

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Croatia

Available in:

Croatia

Package description:

box containing 1 HDPE container of 100 ml box containing 1 vial of 20 ml box containing 1 vial of 10 ml box containing 60 vials of 2 ml

Additional information

box containing 15 vials of 2 ml

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

24/09/2019

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/19-01/590

Date of authorisation status change:

14/10/2025

Reference member state:

Spain

Procedure number:

ES/V/0305/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Norway Poland Portugal Romania Slovakia Slovenia Sweden 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: 12/04/2023 Download
Package Leaflet
Labelling
Combined File of all Documents

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