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Dalmazin, 75 micrograms/ml, solution for injection for cattle, buffaloes and pigs

Authorised

- Cloprostenol

Product identification

Medicine name:
Dalmazin, 75 micrograms/ml, solution for injection for cattle, buffaloes and pigs

Active substance:
Cloprostenol

Target species:
Cattle (cow)
Pig (sow)
Buffalo (female)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Cloprostenol
75.00 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (cow)**
  - Meat and offal. 0 day
  - Milk. 0 hour
- **Pig (sow)**
  - Meat and offal. 1 day
- **Buffalo (female)**
  - Meat and offal. 1 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG02AD90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Cardboard box with 1 type II glass vial of 50 ml with a chlorobutyl type I rubber stopper and an aluminium overseal
Cardboard box with 1 HDPE multidose container of 100 ml with a chlorobutyl type I rubber stopper and an aluminium overseal
Cardboard box containing a 20 ml type II glass vial sealed with a chlorobutyl type I rubber stopper and aluminium overseal

Cardboard box containing 5 x 20 ml type II glass vials sealed with a chlorobutyl type I rubber stopper and aluminium overseal
Cardboard box containing 10 x 10 ml type II glass vials sealed with a chlorobutyl type I rubber stopper and aluminium overseal
Cardboard box containing 1 blister of 15 x 2 ml type I glass vials (15 vials) sealed with a chlorobutyl type I rubber stopper and aluminium overseal
Cardboard box containing 4 blister of 15 x 2 ml type I glass vials (60 vials) sealed with a chlorobutyl type I rubber stopper and aluminium overseal
Cardboard box containing 1 x 10 ml type II glass vial sealed with a chlorobutyl type I rubber stopper and aluminium overseal

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
4/12/2000

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
10/03/2026

Reference member state:
Italy

Procedure number:
IT/V/0105/001

Concerned member states:
Austria Belgium France Germany Ireland Luxembourg
United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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

Combined File of all Documents