

# 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

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**Active substance:**

Cloprostenol

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**Target species:**

Cattle (cow)

Pig (sow)

Buffalo (female)

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

Cloprostenol

75.00 microgram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Cattle (cow)**

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

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**Pig (sow)**

- Meat and offal. 1 day

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**Buffalo (female)**

- Meat and offal. 1 day
  - Milk. 0 hour
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QG02AD90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Italy

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**Package description:**

Cardboard box containing 1 x 10 ml type II glass vial 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 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 10 x 10 ml type II glass vials 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 a 20 ml type II glass vial sealed with a chlorobutyl type I rubber stopper and aluminium overseal

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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Fatro S.p.A.

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**Marketing authorisation date:**

17/12/1991

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**Manufacturing sites for batch release:**

Fatro S.p.A.

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

101725

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**Date of authorisation status change:**

29/06/2010

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**Reference member state:**

Italy

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**Procedure number:**

IT/V/0105/001

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**Concerned member states:**

Austria Belgium France Germany Ireland Luxembourg

United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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