

# 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:**

Austria

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**Available in:**

Austria

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**Package description:**

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

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Available only in [German](#)

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:**

8/09/2000

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### **Manufacturing sites for batch release:**

Fatro S.p.A.

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### **Responsible authority:**

Austrian Agency For Health And Food Safety

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### **Authorisation number:**

8-00470

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### **Date of authorisation status change:**

8/09/2000

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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

Package Leaflet

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

Summary of Product Characteristics

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