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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 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

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

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Buffalo (female)

- Meat and offal. 1 day
- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

OG02AD90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

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 15×2 ml type I glass vials (60 vials) sealed with a chlorobutyl type I rubber stopper and aluminium overseal

Cardboard box containing 1 blister of 15×2 ml type I glass vials (15×15 vials) sealed with a chlorobutyl type I rubber stopper and aluminium overseal

Cardboard box containing $10 \times 10 \text{ ml}$ type II glass vials sealed with a chlorobutyl type I rubber stopper and aluminium overseal

Cardboard box containing 5 \times 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

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:

13/10/2000

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10836/001/001

| Date of authorisation status change: 13/10/2000 |
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| 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 |
| Combined File of all Documents |
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