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:
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 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

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

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