Dimazon 50 mg/ml Solution for Injection

Authorised

- Furosemide

Product identification

Medicine name:
Dimazon 50 mg/ml Solution for Injection

Active substance:
Furosemide

Target species:
Dog
Cat
Cattle
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Furosemide
59.23 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Meat and offal. 1 day
  - Milk. 24 hour
- **Horse**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC03CA01

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
Clear type I tubular glass container sealed with grey type I bromobutyl rubber stopper and aluminum cap with a filling volume of 10 ml.

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:
Intervet (Ireland) Limited

Marketing authorisation date:
1/10/1995

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10996/109/001

Date of authorisation status change:
1/10/1995

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

Documents

Summary of Product Characteristics