

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