Dimazon 50 mg/ml Oplossing voor injectie

Authorised

- Furosemide

Product identification

Medicine name:
Dimazon 50 mg/ml Oplossing voor injectie

Active substance:
Furosemide

Target species:
Cat
Dog
Cattle
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Furosemide
50.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Milk. 1 day
  - Meat and offal. 1 day
- **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:
Belgium

Available in:
Belgium

Package description:
Dimazon 1 Vial with 10 ml Solution for injection

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 International B.V.

Marketing authorisation date:
1/06/1972

Manufacturing sites for batch release:
Intervet International GmbH

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V067444

Date of authorisation status change:
17/07/2024

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

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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