

Dimazon 50 mg/ml Injektionslösung für Rinder, Pferde, Hunde und Katzen

Authorised

- Furosemide

Product identification

Medicine name:

Dimazon 50 mg/ml Injektionslösung für Rinder, Pferde, Hunde und Katzen

Active substance:

Furosemide

Target species:

Dog
Cat
Cattle
Horse

Route of administration:

Intramuscular use
Intravenous use

Product details

Active substance and strength:

Furosemide

50.00 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

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Horse

- Meat and offal. 1 day
 - Milk. 24 hour
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC03CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria

Package description:

Available only in German

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet Ges.m.b.H.

Marketing authorisation date:

5/04/1973

Manufacturing sites for batch release:

Intervet International GmbH

Responsible authority:

Austrian Agency For Health And Food Safety

Authorisation number:

15262

Date of authorisation status change:

5/04/1973

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

Documents

Package Leaflet

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Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.