

Dimazon 50mg/ml ενέσιμο διάλυμα

Not
authorised

- Furosemide

Product identification

Medicine name:

Dimazon 50mg/ml ενέσιμο διάλυμα

Active substance:

Furosemide

Target species:

Cat
Cattle
Dog
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:**Intramuscular use:**

-

Cattle

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

-

Horse

- Meat and offal. 1 day

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:

Surrendered

Authorised in:

Cyprus

Package description:

Available only in Greek

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 International B.V.

Marketing authorisation date:

26/10/1972

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

3421

Date of authorisation status change:

21/03/2012

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

Documents

Summary of Product Characteristics

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