Romefen vet. 100 mg/ml stungulyf, lausn

Authorised

- Ketoprofen

Product identification

Medicine name:
Romefen vet. 100 mg/ml stungulyf, lausn

Active substance:
Ketoprofen

Target species:
Cattle
Pig
Horse

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Ketoprofen
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 4 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 4 day

Intravenous use:

- **Cattle**
  - Meat and offal. 1 day
  - Milk. 0 hour
- **Horse**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE03

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

Authorisation status:
Valid

Authorised in:
Iceland

Available in:
Iceland

Package description:
Available only in Icelandic
Available only in Icelandic
Available only in Icelandic
Available only in Icelandic
Available only in Icelandic
Available only in Icelandic

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
1/10/1995

Manufacturing sites for batch release:
Merial
Ceva Sante Animale

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
930245

Date of authorisation status change:
13/07/2011

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

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