Baytril vet. 100 mg/ml Stungulyf, lausn Handa nautgripum, sauðfé, geitum og svínum.

Authorised

- Enrofloxacin

Product identification

Medicine name:
Baytril vet. 100 mg/ml Stungulyf, lausn Handa nautgripum, sauðfé, geitum og svínum.

Active substance:
Enrofloxacin

Target species:
Cattle
Sheep
Goat
Pig

Route of administration:
Intravenous use
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Enrofloxacin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle**
  - Meat and offal. 5 day
  - Milk. 3 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 12 day
  - Milk. 4 day
- **Sheep**
  - Meat and offal. 4 day
  - Milk. 3 day
- **Goat**
  - Meat and offal. 6 day
  - Milk. 4 day

Intramuscular use:

- **Pig**
  - Meat and offal. 13 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
Elanco Animal Health GmbH

Marketing authorisation date:
13/10/2008

Manufacturing sites for batch release:
KVP Pharma+Veterinär Produkte GmbH

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/08/006/02

Date of authorisation status change:
17/01/2022

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

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.
