Procamidor 20 mg/ml - Injektionslösung für Tiere

Authorised

- Procaine hydrochloride

Product identification

Medicine name:
Procamidor 20 mg/ml - Injektionslösung für Tiere
Procamidor vet. 20 mg/ml stungulyf, lausn

Active substance:
Procaine hydrochloride

Target species:
Cattle
Dog
Sheep
Pig
Cat
Horse

Route of administration:
Epidural use
Perineural use
Infiltration

Product details

Active substance and strength:
Procaine hydrochloride
20.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Epidural use:

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

Infiltration:

- **Cattle**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 hour
  - Meat and offal. 0 day

- **Horse**
  - Milk. 0 hour
  - Meat and offal. 0 day
- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01BA02

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

Authorisation status:
Valid

Authorised in:
Iceland

Available in:
Iceland

Package description:
Clear glass vial type II (Ph. Eur.) with bromobutyl rubber stopper type I (Ph.Eur.) and aluminium cap. Package sizes 10 x 100 ml
Clear glass vial type II (Ph. Eur.) with bromobutyl rubber stopper type I (Ph.Eur.) and aluminium cap. Package size: 1 x 100 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetviva Richter GmbH

Marketing authorisation date:
4/10/2013

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/13/012/01

Date of authorisation status change:
25/08/2017

Reference member state:
Austria

Procedure number:
AT/V/0011/001

Concerned member states:
Czechia Denmark Estonia Finland France Germany Iceland Italy Latvia Lithuania Netherlands Norway Portugal Romania Slovakia Slovenia Spain Sweden

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

Documents

Package Leaflet

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Labelling

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

Summary of Product Characteristics

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