

Procamidor 20 mg/ml - Injektionslösung für Tiere

Authorised

- Procaine hydrochloride

Product identification

Medicine name:

Procamidor 20 mg/ml - Injektionslösung für Tiere

Procamidor 20 mg/ml injekcinis tirpalas

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:

Lithuania

Available in:

Lithuania

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:

3/11/2013

Manufacturing sites for batch release:

Vetviva Richter GmbH

Responsible authority:

State Food And Veterinary Service

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

19/09/2017

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

RV2191.pdf