Procainii Chloridum 4 % + Adrenalinum 40 mg/ml Solution injectable

Authorised

- EPINEPHRINE BITARTRATE
- Procaine hydrochloride

Product identification

Medicine name:
Procainii Chloridum 4 % + Adrenalinum 40 mg/ml Solution injectable
Procainii Chloridum 4 % + Adrenalinum 40 mg/ml Injektionslösung

Active substance:
EPINEPHRINE BITARTRATE
Procaine hydrochloride

Target species:
Cattle
Sheep
Pig
Horse

Route of administration:
Epidural use
Intramuscular use
Subcutaneous use
Perineural use

Product details

Active substance and strength:
EPINEPHRINE BITARTRATE
0.04 milligram(s) / 1.00 millilitre(s)

Procaine hydrochloride
40.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Epidural use:

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

Intramuscular use:

- **Cattle**
  - Meat and offal. 3 day
- **Sheep**
  - Meat and offal. 3 day
- **Pig**
  - Meat and offal. 3 day
- **Horse**
  - Meat and offal. 3 day

Subcutaneous use:

Cattle

- Meat and offal. 3 day

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Sheep

- Meat and offal. 3 day

•

Pig

- Meat and offal. 3 day

•

Horse

- Meat and offal. 3 day

Perineural use:

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Cattle

- Meat and offal. 3 day

•

Sheep

- Meat and offal. 3 day

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Pig

- Meat and offal. 3 day

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Horse

- Meat and offal. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01BA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Luxembourg

Available in:
Luxembourg

Package description:
Procainii Chloridum 4 % + Adrenalinum 40 mg/ml Solution for injection Vial of 50 ml
Procainii Chloridum 4 % + Adrenalinum 40 mg/ml Solution for injection Vial of 100 ml
Procainii Chloridum 4 % + Adrenalinum 40 mg/ml Solution for injection Vial of 250 ml
Procainii Chloridum 4 % + Adrenalinum 40 mg/ml Solution for injection Vial of 500 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
24/01/1975

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Ministry Of Health And Social Security

Authorisation number:
V 188/18/05/1695

Date of authorisation status change:
14/11/2008

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.