Procaine HCl 4 % + Adrenaline Oplossing voor injectie

Authorised

- EPINEPHRINE BITARTRATE
- Procaine hydrochloride

Product identification

Medicine name:
Procaine HCl 4 % + Adrenaline Oplossing voor injectie
Procaine HCl 4 % + Adrenaline Solution injectable
Procaine HCl 4 % + Adrenaline Injektionslösung

Active substance:
EPINEPHRINE BITARTRATE
Procaine hydrochloride

Target species:
Horse
Cattle

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

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

Infiltration:

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

Epidural use:

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

Subcutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 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:
Belgium

Package description:
Procaine HCl 4 % + Adrenaline 1 Vial with 250 ml Solution for injection

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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:
Prodivet Pharmaceuticals

Marketing authorisation date:
22/04/1977

Manufacturing sites for batch release:
Prodivet Pharmaceuticals

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V107791

Date of authorisation status change:
7/10/2016

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

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