Procaine HCl 2 % Oplossing voor injectie

Not authorised

- Procaine hydrochloride

Product identification

Medicine name:
Procaine HCl 2 % Oplossing voor injectie
Procaine HCl 2 % Solution injectable
Procaine HCl 2 % Injektionslösung

Active substance:
Procaine hydrochloride

Target species:
Sheep
Cattle
Pig
Dog
Horse
Cat

Route of administration:
Epidural use
Subcutaneous use
Perineural use

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:

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

Subcutaneous use:

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

- **Pig**
  - Meat and offal. 0 day
- **Sheep**
  - 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

Authorisation status:
Surrendered

Authorised in:
Belgium

Package description:
Procaine HCl 12 Vials with 500 ml of Solution for injection
Procaine HCl 12 Vials with 250 ml of Solution for injection
Procaine HCl 12 Vials with 100 ml of Solution for injection
Procaine HCl 12 Vials with 50 ml of Solution for injection
Procaine HCl 1 Vial with 500 ml of Solution for injection
Procaine HCl 1 Vial with 250 ml of Solution for injection
Procaine HCl 1 Vial with 100 ml of Solution for injection
Procaine HCl 1 Vial with 50 ml of Solution for injection

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:
V.M.D.

Marketing authorisation date:
26/09/1974

Manufacturing sites for batch release:
VMD N.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V104361

Date of authorisation status change:
5/03/2024

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

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