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Procamidor Duo 40 mg/ml + 0,036 mg/ml Injektionslösung für Tiere

Authorised

- Procaine hydrochloride
- Adrenaline hydrogen tartrate

Product identification

Medicine name:
Procamidor Duo 40 mg/ml + 0,036 mg/ml Injektionslösung für Tiere

Active substance:
Procaine hydrochloride
Adrenaline hydrogen tartrate

Target species:
Cattle
Sheep
Horse
Pig

Route of administration:
Perineural use
Subcutaneous use

Product details

Active substance and strength:

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

Adrenaline hydrogen tartrate
0.04 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Perineural use:

- **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

Subcutaneous use:

- **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:
QN01BA52

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

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Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
Amber glass vial type II (Ph. Eur.) with coated or uncoated bromobutyl rubber stopper type I (Ph.Eur.) and aluminium cap in a cardboard box. Cardboard box with 1 vial of 100 ml
Amber glass vial type II (Ph. Eur.) with coated or uncoated bromobutyl rubber stopper type I (Ph.Eur.) and aluminium cap in a cardboard box. Cardboard box with 1 vial of

250 ml
Amber glass vial type II (Ph. Eur.) with coated or uncoated bromobutyl rubber stopper type I (Ph.Eur.) and aluminium cap in a cardboard box. Cardboard box with 5 vials of 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:
1/07/2019

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Danish Medicines Agency

Authorisation number:
60986

Date of authorisation status change:
1/07/2019

Reference member state:
Austria

Procedure number:
AT/V/0018/001

Concerned member states:
Belgium Bulgaria Croatia Czechia Denmark Estonia Finland Germany

Greece Hungary Iceland Ireland Italy Latvia Lithuania Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 3/04/2025
Updated on: 13/03/2026
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