Procamidor Duo 40 mg/ml + 0.036 mg/ml solution for injection

Authorised

- Procaine hydrochloride
- EPINEPHRINE BITARTRATE

Product identification

Medicine name:
Procamidor Duo 40 mg/ml + 0,036 mg/ml Injektionslösung für Tiere
Procamidor Duo 40 mg/ml + 0.036 mg/ml solution for injection

Active substance:
Procaine hydrochloride
EPINEPHRINE BITARTRATE

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)

EPINEPHRINE BITARTRATE
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

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Sheep

- Milk. 0 hour

- Meat and offal. 0 day

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Horse

- Milk. 0 hour

- Meat and offal. 0 day

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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:
Ireland

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:
30/08/2019

Manufacturing sites for batch release:
Vetviva Richter GmbH

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA23462/011/001

Date of authorisation status change:
30/08/2019

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