

Pronestestic 40 mg/ml + 0.036 mg/ml solution for injection for horses, cattle, pigs and sheep

Authorised

- Procaine hydrochloride
- ADRENALINE TARTRATE PH. EUR.

Product identification

Medicine name:

Pronestestic 40 mg/ml + 0.036 mg/ml solution for injection for horses, cattle, pigs and sheep

Active substance:

Procaine hydrochloride
ADRENALINE TARTRATE PH. EUR.

Target species:

Cattle
Sheep
Pig
Horse

Route of administration:

Perineural use
Subcutaneous use

Product details

Active substance and strength:

Procaine hydrochloride

40.00 milligram(s) / 1.00 millilitre(s)

ADRENALINE TARTRATE PH. EUR.

0.04 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Perineural use:

-

Cattle

- Meat and offal. 0 day

- Milk. 0 day

-

Sheep

- Meat and offal. 0 day

- Milk. 0 day

-

Pig

- Meat and offal. 0 day

-

Horse

- Meat and offal. 0 day

- Milk. 0 day

Subcutaneous use:

-

Cattle

- Meat and offal. 0 day

- Milk. 0 day

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Sheep

- Meat and offal. 0 day

- Milk. 0 day

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Pig

- Meat and offal. 0 day

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Horse

- Meat and offal. 0 day

- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01BA52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

box containing 10 vials of 100 ml

box containing 1 vial of 250 ml

box containing 1 vial of 100 ml

box containing 1 vial of 50 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:

Fatro S.p.A.

Marketing authorisation date:

15/04/2016

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10836/004/001

Date of authorisation status change:

15/04/2016

Reference member state:

Spain

Procedure number:

ES/V/0238/001

Concerned member states:

Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France
Germany Greece Iceland Ireland Italy Latvia Lithuania Luxembourg
Netherlands Norway Poland Portugal Slovakia Slovenia 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

Package Leaflet

Labelling

Combined File of all Documents

eu-PUAR-esv0238001-dcp-pronestestic-40-mg-ml-+-0.036-mg-ml-solution-for-injection-for-horses--cattle--swine-and-sheep-en.pdf