

PRONESTESIC 40 mg/ml + 0.036 mg/ml solution for injection for horses, cattle, swine and sheep

Authorised

- Procaine hydrochloride
- ADRENALINE TARTRATE PH. EUR.

Product identification

Medicine name:

PRONESTESIC 40 mg/ml + 0.036 mg/ml solution for injection for horses, cattle, swine and sheep

Pronestesic 40 mg/ml / 0,036 mg/ml oplossing voor injectie voor paard, rund, varken en schaap

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

•

Sheep

- Meat and offal. 0 day
- Milk. 0 day

•

Pig

- Meat and offal. 0 day

•

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:

Netherlands

Available in:

Netherlands

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:

22/03/2016

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 117141

Date of authorisation status change:

2/02/2022

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

Combined File of all Documents

English (PDF)

Published on: 24/07/2025

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Summary of Product Characteristics

Package Leaflet

Labelling

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