

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

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 + 0,036 mg)/ml Roztwór do wstrzykiwań

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

-

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

Authorisation status:

Valid

Authorised in:

Poland

Available in:

Poland

Package description:

Available only in Polish

Available only in Polish

Available only in Polish

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:

14/10/2019

Manufacturing sites for batch release:

Vetviva Richter GmbH

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

2915

Date of authorisation status change:

14/10/2019

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