

ULTRAPEN LA 300 mg/ml, injekciné suspensija

Not
authorised

- BENZYL PENICILLIN PROCAINE

Product identification

Medicine name:

ULTRAPEN LA 300 mg/ml, injekciné suspensija

Active substance:

BENZYL PENICILLIN PROCAINE

Target species:

Cattle

Pig

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

BENZYL PENICILLIN PROCAINE

300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Meat and offal. 21 day
- Milk. 5 day

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Pig

- Meat and offal. 7 day

Subcutaneous use:

-

Cattle

- Meat and offal. 10 day Subcutaneous use for non-lactating cattle.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CE09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:

Surrendered

Authorised in:

Lithuania

Package description:

Available only in Lithuanian

Available only in Lithuanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:

Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:

11/09/2000

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/00/1153/001-002

Date of authorisation status change:

25/01/2011

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

Documents

RV1153.pdf