ULTRAPEN LA 300 mg/ml, injekcinė suspensija

Not authorised

- BENZYLPENICILLIN PROCAINE

Product identification

Medicine name:
ULTRAPEN LA 300 mg/ml, injekcinė suspensija

Active substance:
BENZYLPENICILLIN PROCAINE

Target species:
Cattle
Pig

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
BENZYLPENICILLIN 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
- **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