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Ultrapen LA, 300mg/ml, Injekční suspenze

Authorised

- Benzylpenicillin (procaine) monohydrate

Product identification

Medicine name:
Ultrapen LA, 300mg/ml, Injekční suspenze

Active substance:
Benzylpenicillin (procaine) monohydrate

Target species:
Cattle
Pig

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:
Benzylpenicillin (procaine) monohydrate
300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 10 day

Intramuscular use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 5 day
- **Pig**
  - Meat and offal. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CE09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

Available only in Czech

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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
30/10/2000

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/062/00-C

Date of authorisation status change:
30/10/2000

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.