

# Ultrapen LA 300 mg/ml suspensija injekcijām liellopiem un cūkām

Authorised

- Benzylpenicillin procaine

## Product identification

**Medicine name:**

Ultrapen LA 300 mg/ml suspensija injekcijām liellopiem un cūkām

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**Active substance:**

Benzylpenicillin procaine

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**Target species:**

Cattle

Pig

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Benzylpenicillin procaine

300.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Cattle**

- Milk. 132 hour
- Meat and offal. 23 day Ja ārstēšanas ilgums ir 3 dienas.
- Meat and offal. 25 day Ja ārstēšanas ilgums ir 4-7 dienas.

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**Pig**

- Meat and offal. 10 day Ja ārstēšanas ilgums ir 3 dienas.
- Meat and offal. 12 day Ja ārstēšanas ilgums ir 4-7 dienas.

**Subcutaneous use:**

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**Cattle**

- Meat and offal. 13 day Ja ārstēšanas ilgums ir 3 dienas.
- Milk. no withdrawal period

Nav reģistrēts subkutānai lietošanai dzīvniekiem, kuru pienu paredzēts izmantot cilvēku uzturā.

- Meat and offal. 15 day Ja ārstēšanas ilgums ir 4-7 dienas.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01CE09

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Latvia

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**Package description:**

Available only in Latvian

Available only in Latvian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Norbrook Laboratories (Ireland) Limited

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**Marketing authorisation date:**

3/12/1998

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**Manufacturing sites for batch release:**

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

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**Responsible authority:**

Food And Veterinary Service

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**Authorisation number:**

V/NRP/98/0939

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**Date of authorisation status change:**

3/12/1998

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

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### Labelling

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

### Package Leaflet

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