File downloaded on 2026-11-25
Source URL: https://medicines.health.europa.eu/veterinary/en/600000055135

Ultrapen LA, 300 mg/ml süstesuspensioon veistele ja sigadele

Authorised

- Benzylpenicillin procaine

Product identification

Medicine name:
Ultrapen LA, 300 mg/ml süstesuspensioon veistele ja sigadele

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. 8 day piim: 8 päeva (192 tundi)
- **Pig**
  - Meat and offal. 10 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 13 day
  - Milk. no withdrawal period

Subkutaanseks kasutamiseks mittelakteerivatel veistel

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CE09

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Estonia

Package description:
Available only in Estonian
Available only in Estonian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
25/03/2004

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

Responsible authority:
State Agency Of Medicines

Authorisation number:
1227

Date of authorisation status change:
25/03/2004

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

Documents

Summary of Product Characteristics

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