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Ultrapen LA 300 mg/ml Suspension for Injection

Authorised

- Benzylpenicillin procaine

Product identification

Medicine name:
Ultrapen LA 300 mg/ml Suspension for Injection

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CE09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

50 ml Grade II clear glass vial, complete with nitryl bung and aluminium cap.
100 ml Grade II clear glass vial, complete with nitryl bung and aluminium cap.

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:
12/12/1997

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/045/001

Date of authorisation status change:
12/12/1997

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

Documents

Summary of Product Characteristics

ie-puar-np-600000064329-ultrapen-la-300-mgml-suspension-for-injection-en.pdf