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:

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