Oxytetracycline 200 mg/ml L.A. solution for injection for cattle

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:
Oxytetracycline 200 mg/ml L.A. solution for injection for cattle

Active substance:
Oxytetracycline hydrochloride

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Oxytetracycline hydrochloride
215.86 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 10 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA06

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
Brown glass vial 100 ml, type II, with a bromobutyl rubber closure, sealed with an aluminium cap containing 100 ml ofproduct.The vials are packed in a polystyrene box, 12 vials of 100 ml per box, with 12 package leaflets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent (abridged application) - Council Directive 81/851/EEC

Marketing authorisation holder:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
1/10/1988

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10981/005/001

Date of authorisation status change:
1/10/1988

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

Documents

Summary of Product Characteristics

Source URL: *https://medicines.health.europa.eu/veterinary/600000064413*