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Lutalyse

Authorised

- Dinoprost trometamol

Product identification

Medicine name:
Lutalyse

Active substance:
Dinoprost trometamol

Target species:
Cattle
Horse
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Dinoprost trometamol
6.71 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 2 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 1 day
- **Pig**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG02AD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

30 ml multi-dose, Type I, glass vial with red chlorobutyl rubber stopper and aluminium overseal.Pack sizes: 30 ml vial.
10 ml multi-dose, Type I, glass vials with red chlorobutyl rubber stopper and aluminium overseal.Pack sizes: 5 x 10 ml vials.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Zoetis Belgium S.A.

Marketing authorisation date:
18/10/2013

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10387/042/001

Date of authorisation status change:
18/10/2013

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

Documents

Summary of Product Characteristics