Levapharm Injection 75 mg/ml

Authorised

- Levamisole hydrochloride

Product identification

Medicine name:
Levapharm Injection 75 mg/ml

Active substance:
Levamisole hydrochloride

Target species:
Cattle
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Levamisole hydrochloride
88.39 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 14 day

- **Sheep**
  - Meat and offal. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AE01

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

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Authorisation status:
Valid

Authorised in:
Ireland

Package description:
The veterinary medicinal product is supplied in glass vials in a cardboard box, sealed with bromobutyl bung and aluminium seals. Package size: 1 x 100 ml vial in a cardboard box
The veterinary medicinal product is supplied in glass vials in a cardboard box, sealed with bromobutyl bung and aluminium seals. Package size: 1 x 250 ml vial in a cardboard box
The veterinary medicinal product is supplied in glass vials in a cardboard box, sealed with bromobutyl bung and aluminium seals. Package size: 1 x 500 ml vial in a cardboard box
The veterinary medicinal product is supplied in glass vials in a cardboard box, sealed with bromobutyl bung and aluminium seals. Package size: 6 x 500 ml vials with 1 protective container in a cardboard box.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Chem-Pharm (Ballyvaughan) Limited

Marketing authorisation date:
1/10/1999

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10823/010/001

Date of authorisation status change:
1/10/1999

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

Documents

Summary of Product Characteristics