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Ivomec Super Injection for Cattle

Authorised

- Ivermectin
- Clorsulon

Product identification

Medicine name:
Ivomec Super Injection for Cattle

Active substance:
Ivermectin

Clorsulon

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Ivermectin
10.00 milligram(s) / 1.00 millilitre(s)

Clorsulon
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 66 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Multiple-dose rubber-capped polyethylene bottle of 50 ml containing a sterile non-aqueous solution.
Multiple-dose rubber-capped polyethylene bottle of 200 ml containing a sterile non-aqueous solution.
Multiple-dose rubber-capped polyethylene bottle of 500 ml containing a sterile non-aqueous solution.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
1/10/1999

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10454/068/001

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