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Topimec Super solution for injection

Authorised

- Ivermectin
- Clorsulon

Product identification

Medicine name:
Topimec Super solution for injection

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:
Container material: High density polyethyleneContainer closure: Siliconised grey bromobutyl rubber stopperContainer volume: 500 ml
Container material: High density polyethyleneContainer closure: Siliconised grey bromobutyl rubber stopperContainer volume: 250 ml
Container material: High density polyethyleneContainer closure: Siliconised grey bromobutyl rubber stopperContainer volume: 50 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
14/10/2011

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/099/001

Date of authorisation status change:
14/10/2011

Reference member state:
Ireland

Procedure number:
IE/V/0286/001

Concerned member states:
Czechia France Greece Poland Portugal Spain

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

Documents

Combined File of all Documents

Summary of Product Characteristics