Imaverol 100 mg/ml Concentrate for Cutaneous Emulsion

Authorised

- Enilconazole

Product identification

Medicine name:
Imaverol 100 mg/ml Concentrate for Cutaneous Emulsion

Active substance:
Enilconazole

Target species:
Cattle
Dog
Horse

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Enilconazole
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Cutaneous emulsion

Withdrawal period by route of administration:

Cutaneous use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD01AC90

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
1 litre pack size: Container: white, high density polyethylene bottle with transparent window containing 1 litre of concentrate. Closure: white high density polyethylene screw cap and seal insert lined with polyethylene Dosing device: low density polyethylene measuring cup
100 ml pack size: Container: amber Type III glass bottle containing 100ml of concentrate. Closure: tamper evident and child resistant polypropylene screw cap lines with LDPE Dosing device: polypropylene measuring cup

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Audevard

Marketing authorisation date:
30/09/2009

Manufacturing sites for batch release:
Mcgregor Cory Limited
Lusomedicamenta Sociedade Tecnica Farmaceutica S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10481/001/001

Date of authorisation status change:
30/09/2009

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

Documents

Summary of Product Characteristics