File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600000048698

Qualimec 10 mg/ml solution for injection for cattle, sheep and pigs

Authorised

- Ivermectin

Product identification

Medicine name:
Qualimec 10 mg/ml solution for injection for cattle, sheep and pigs

Active substance:
Ivermectin

Target species:
Cattle
Sheep
Pig

Route of administration:
Subcutaneous use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - Meat and offal. 49 day
- **Sheep**
  - Meat and offal. 42 day
- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA01

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
Clear PET multidose container with bromobutyl rubber stopper and aluminium cap. Pack size 250 ml
HDPE multidose container with bromobutyl rubber stopper and aluminium cap.Pack size: 50 ml
HDPE multidose container with bromobutyl rubber stopper and aluminium cap.Pack size: 500 ml.
Clear PET multidose container with bromobutyl rubber stopper and aluminium cap. Pack size 500 ml

HDPE multidose container with bromobutyl rubber stopper and aluminium cap.Pack size: 200 ml
Clear PET multidose container with bromobutyl rubber stopper and aluminium cap. Pack size 50 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Eco Animal Health Europe Limited

Marketing authorisation date:
1/10/1999

Manufacturing sites for batch release:
Divasa Farmavic S.A.
Produlab Pharma B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22693/002/001

Date of authorisation status change:
1/10/1999

Reference member state:
Ireland

Procedure number:
IE/V/0145/001

Concerned member states:

France Germany Greece Hungary Italy Luxembourg Netherlands Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents