Rafazole Oral Suspension

Not authorised

- Rafoxanide
- Levamisole hydrochloride

Product identification

Medicine name:
Rafazole Oral Suspension

Active substance:
Rafoxanide

Levamisole hydrochloride

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Rafoxanide
30.00 milligram(s) / 1.00 millilitre(s)

Levamisole hydrochloride
30.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 60 day

- **Sheep**
  - Meat and offal. 60 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AE51

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

Authorisation status:
Surrendered

Authorised in:
Ireland

Package description:
10 litre. High density polyethylene containers with polyethylene closures.
5 litre. High density polyethylene containers with polyethylene closures.
2.5 litre. High density polyethylene containers with polyethylene closures.
1 litre. High density polyethylene containers with polyethylene closures.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Informed consent (abridged application) - Council Directive 81/851/EEC

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
1/10/1989

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/012/001

Date of authorisation status change:
26/06/2024

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

Documents

Summary of Product Characteristics