Parafend 2.265 % Oral Suspension

Authorised

- Oxfendazole

Product identification

Medicine name:
Parafend 2.265 % Oral Suspension

Active substance:
Oxfendazole

Target species:
Cattle
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Oxfendazole
2.27 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 5 day
- **Sheep**
  - Meat and offal. 10 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Parafend will be presented in 10 L multi-dose polyethylene containers with polyethylene closures.
Parafend will be presented in 5 L multi-dose polyethylene containers with polyethylene closures.
Parafend will be presented in 2.5 L multi-dose polyethylene containers with polyethylene closures.
Parafend will be presented in 1.0 L multi-dose polyethylene containers with polyethylene closures.
Parafend will be presented in 500 ml multi-dose 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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
10/02/1994

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/040/001

Date of authorisation status change:
10/02/1994

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

Documents

Summary of Product Characteristics