

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:**

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Cattle

- Meat and offal. 14 day

- Milk. 5 day

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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 500 ml 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 2.5 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 10 L 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