

# Fasifree 10% w/v Oral Suspension

Authorised

- Triclabendazole

## Product identification

**Medicine name:**

Fasifree 10% w/v Oral Suspension

**Active substance:**

Triclabendazole

**Target species:**

Cattle

Sheep

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Triclabendazole

100.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**

Oral suspension

**Withdrawal period by route of administration:****Oral use:**

- Cattle

- Meat and offal. 56 day

- Milk. 48 hour

- **Sheep**

- Meat and offal. 55 day

- Milk. 1 year

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP52AC01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Ireland

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**Package description:**

1 litre high density polyethylene flat bottom back pack containers containing a white to off-white smooth suspension. The closures used for all pack sizes are 38 mm polypropylene standard caps with an induction heat seal liner. A 38 mm polypropylene spouted cap is also provided with each pack size for dispensing purposes.

2.5 litre high density polyethylene flat bottom back pack containers containing a white to off-white smooth suspension. The closures used for all pack sizes are 38 mm polypropylene standard caps with an induction heat seal liner. A 38 mm polypropylene spouted cap is also provided with each pack size for dispensing purposes.

5 litre high density polyethylene flat bottom back pack containers containing a white to off-white smooth suspension. The closures used for all pack sizes are 38 mm polypropylene standard caps with an induction heat seal liner. A 38 mm polypropylene spouted cap is also provided with each pack size for dispensing purposes.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Bimeda Animal Health Limited

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**Marketing authorisation date:**

22/02/2002

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**Manufacturing sites for batch release:**

Bimeda Animal Health Limited

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA22033/020/001

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**Date of authorisation status change:**

22/02/2002

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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**Source URL:** <https://medicines.health.europa.eu/veterinary/600000064300>