File downloaded on 2026-11-26
Source URL: https://medicines.health.europa.eu/veterinary/en/600000040113

OXFENIL 22,65 MG/ML SUSPENSION BUVABLE

Authorised

- Oxfendazole

Product identification

Medicine name:
OXFENIL 22,65 MG/ML SUSPENSION BUVABLE

Active substance:
Oxfendazole

Target species:
Cattle
Sheep
Goat

Route of administration:
Oral use

Product details

Active substance and strength:
Oxfendazole
22.65 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 15 day
  - Milk. 7 day
- **Sheep**
  - Meat and offal. 14 day
  - Milk. 8 day
- **Goat**
  - Meat and offal. 28 day
  - Milk. 14 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Available only in French

Available only in French

Available only in French

Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
24/09/1996

Manufacturing sites for batch release:
Virbac
FC France S.A.S.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/3743249 0/1996

Date of authorisation status change:
24/09/2011

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.