PANACUR 25 mg/ml SUSPENSION ORAL

Authorised

- Fenbendazole

Product identification

Medicine name:
PANACUR 25 mg/ml SUSPENSION ORAL

Active substance:
Fenbendazole

Target species:
Sheep

Route of administration:
Oral use

Product details

Active substance and strength:
Fenbendazole
25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Sheep**
  - Meat and offal. 16 day
  - Milk. no withdrawal period Leche: 8 días (192 horas)

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC13

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:
12/05/1993

Manufacturing sites for batch release:
Intervet Productions S.A.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
693 ESP

Date of authorisation status change:
12/05/1993

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

Documents

Summary of Product Characteristics

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Labelling

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Package Leaflet

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