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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

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

Package Leaflet

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