

# PANACUR 100 mg/ml SUSPENSION ORAL

Authorised

- Fenbendazole

## Product identification

**Medicine name:**

PANACUR 100 mg/ml SUSPENSION ORAL

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**Active substance:**

Fenbendazole

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**Target species:**

Cattle

Sheep

Goat

Horse

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Fenbendazole

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Oral suspension

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

- Meat and offal. 8 day
- Milk. 156 hour

**• Sheep**

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

**• Goat**

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

**• Horse**

- Meat and offal. 14 day
- Milk. no withdrawal period

Leche: Su uso no está autorizado en équidos cuya leche se utiliza para consumo humano

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

QP52AC13

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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:**

Spain

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**Package description:**

Available only in Spanish

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Merck Sharp & Dohme Animal Health S.L.

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**Marketing authorisation date:**

12/05/1993

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**Manufacturing sites for batch release:**

Intervet Productions

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**Responsible authority:**

The Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

706 ESP

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**Date of authorisation status change:**

12/05/1993

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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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## Labelling

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## Package Leaflet

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