WORM STOP tablets for dogs

Authorised

- Fenbendazole
- Pyrantel embonate
- Praziquantel

Product identification

Medicine name:
WORM STOP tablets for dogs

Active substance:
Fenbendazole

Pyrantel embonate

Praziquantel

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Fenbendazole
200.00 milligram(s) / 1.00 Tablet

Pyrantel embonate

144.00 milligram(s) / 1.00 Tablet

Praziquantel
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA51

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

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Immediate packaging material: PVC/Alu blister or polyethylene container. Package size: Polyethylene container containing 200 tablets.
Immediate packaging material: PVC/Alu blister or polyethylene container. Package size: carton box containing PVC/Alu blister of 20x10 tablets
Immediate packaging material: PVC/Alu blister or polyethylene container. Package size: carton box containing PVC/Alu blister of 10x10 tablets
Immediate packaging material: PVC/Alu blister or polyethylene container. Package size: carton box containing PVC/Alu blister of 2x10 tablets
Immediate packaging material: PVC/Alu blister or polyethylene container. Package size: carton box containing PVC/Alu blister of 1x10 tablets
Immediate packaging material: PVC/Alu blister or polyethylene container. Package size: carton box containing PVC/Alu blister of 3x2 tablets
Immediate packaging material: PVC/Alu blister or polyethylene container. Package size: carton box containing PVC/Alu blister of 1x2 tablets

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Pharma World Pharmaceuticals Kft.

Marketing authorisation date:
29/01/2015

Manufacturing sites for batch release:
Pernix Pharma Kft.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3169 ESP

Date of authorisation status change:
25/08/2015

Reference member state:
Hungary

Procedure number:
HU/V/0121/001

Concerned member states:
Czechia Germany Italy Poland Portugal Romania Slovakia Spain

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

Documents

Package Leaflet

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

Labelling

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

Summary of Product Characteristics

English (PDF)
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Updated on: 13/03/2026
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