WORM STOP tablets for dogs

Authorised

- Fenbendazole
- Pyrantel embonate
- Praziquantel

Product identification

Medicine name:
WORM STOP tablets for dogs
WORM STOP comprimidos para cães

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:
Portugal

Package description:
Immediate packaging material: PVC/Alu blister or polyethylene container. Package size: carton box containing PVC/Alu blister of 1x2 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 1x10 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 10x10 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: Polyethylene container containing 200 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:
2/10/2014

Manufacturing sites for batch release:
Pernix Pharma Kft.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
845/01/14RFVPT

Date of authorisation status change:
3/06/2025

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

Combined File of all Documents

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