

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:

Germany

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:

6/11/2014

Manufacturing sites for batch release:

Pernix Pharma Kft.

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402146.00.00

Date of authorisation status change:

29/01/2019

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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Summary of Product Characteristics