File downloaded on 2026-11-03
Source URL: https://medicines.health.europa.eu/veterinary/en/600000037028

Cestem flavoured tablets for medium and small dogs

Authorised

- Febantel
- Praziquantel
- Pyrantel embonate

Product identification

Medicine name:
Cestem flavoured tablets for medium and small dogs

Active substance:
Febantel

Praziquantel

Pyrantel embonate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Febantel

150.00 milligram(s) / 1.00 Tablet

Praziquantel
50.00 milligram(s) / 1.00 Tablet

Pyrantel embonate
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:
Greece

Available in:
Greece

Package description:
Polyamide-aluminium-PVC / aluminium blister packs. Box containing 52 blisters of 2 tablets.
Polyamide-aluminium-PVC / aluminium blister packs. Box containing 1 blister of 2 tablets.
Polyamide-aluminium-PVC / aluminium blister packs. Box containing 13 blisters of 8 tablets.
Polyamide-aluminium-PVC / aluminium blister packs. Box containing 2 blisters of 2 tablets.
Polyamide-aluminium-PVC / aluminium blister packs. Box containing 1 blister of 8 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Hellas LLC

Marketing authorisation date:
9/11/2010

Manufacturing sites for batch release:
Ceva Sante Animale
Ceva Sante Animale

Responsible authority:
National Organization For Medicines

Authorisation number:
6373/29-01-2016/K-0187901

Date of authorisation status change:
28/01/2016

Reference member state:
Netherlands

Procedure number:
NL/V/0269/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Estonia Germany Greece Hungary Italy Latvia Lithuania Luxembourg Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Package Leaflet

Combined File of all Documents

PuAR Cestem - 20230619.pdf

106871 PAR.pdf