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Endogard Plus XL Tablets for dogs

Authorised

- Febantel
- Pyrantel embonate
- Praziquantel

Product identification

Medicine name:
Endogard Plus XL Tablets for dogs

Active substance:
Febantel

Pyrantel embonate

Praziquantel

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Febantel
525.00 milligram(s) / 1.00 Tablet

Pyrantel embonate

504.00 milligram(s) / 1.00 Tablet

Praziquantel
175.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC55

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Latvia

Available in:
Latvia

Package description:
Print and perforated Alu-Alu blister: 102 tablets (17 blisters with 6 tablets), in a box.
Print and perforated Alu-Alu blister: 100 tablets (10 blisters with 10 tablets), in a box.
Print and perforated Alu-Alu blister: 60 tablets (6 blisters with 10 tablets), in a box.
Print and perforated Alu-Alu blister: 60 tablets (10 blisters with 6 tablets), in a box.
Print and perforated Alu-Alu blister: 50 tablets (5 blisters with 10 tablets), in a box
Print and perforated Alu-Alu blister: 30 tablets (5 blisters with 6 tablets), in a box.
Print and perforated Alu-Alu blister: 30 tablets (3 blisters with 10 tablets), in a box.
Print and perforated Alu-Alu blister: 24 tablets (4 blisters with 6 tablets), in a box.
Print and perforated Alu-Alu blister: 12 tablets (2 blisters with 6 tablets), in a box.
Print and perforated Alu-Alu blister: 10 tablets (1 blister with 10 tablets), in a box.
Print and perforated Alu-Alu blister: 4 tablets (2 blisters with 2 tablets), in a box.
Print and perforated Alu-Alu blister: 2 tablets (1 blister with 2 tablets), in a box.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
22/03/2011

Manufacturing sites for batch release:
Virbac
Krka-Farma d.o.o.
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/DCP/11/0003

Date of authorisation status change:
22/03/2011

Reference member state:
Ireland

Procedure number:
IE/V/0539/002

Concerned member states:
Austria Belgium Czechia Denmark Estonia Germany Greece Hungary Italy Latvia Lithuania Netherlands Poland Portugal 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

Combined File of all Documents

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