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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 subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

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:
21/02/2011

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

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401283.01.00

Date of authorisation status change:
23/01/2017

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)

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Documents

Combined File of all Documents

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