Wellplus Flavoured Tablets

Authorised

- Febantel
- Pyrantel embonate
- Praziquantel

Product identification

Medicine name:
Wellplus Flavoured Tablets
Wellplus tabletten met smaakstof voor honden

Active substance:
Febantel

Pyrantel embonate

Praziquantel

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Febantel
150.00 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

Withdrawal period by route of administration:

Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA30

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
box containing 300 tablets (30 blisters)
box containing 100 tablets (10 blisters)
box containing 50 tablets (5 blisters)
box containing 20 tablets (2 blisters)
box containing 10 tablets (1 blister)
box containing 2 tablets (1 blister)

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:

Divasa Farmavic S.A.

Marketing authorisation date:
27/11/2013

Manufacturing sites for batch release:
Divasa Farmavic S.A.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 112912

Date of authorisation status change:
22/02/2022

Reference member state:
Spain

Procedure number:
ES/V/0163/001

Concerned member states:
Belgium Bulgaria Germany Greece Ireland Netherlands Poland Portugal Slovakia United Kingdom (Northern Ireland)

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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