PRILACTONE NEXT 50 MG CHEWABLE TABLETS FOR DOGS

Authorised

- Spironolactone

Product identification

Medicine name:
PRILACTONE NEXT 50 MG CHEWABLE TABLETS FOR DOGS
Prilactone Next 50 mg kauwtabletten voor honden

Active substance:
Spironolactone

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Spironolactone
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Withdrawal period by route of administration:

Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC03DA01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Cardboard box with 1 blister of 10 tablets
Cardboard box with 2 blisters of 10 tablets
Cardboard box with 3 blisters of 10 tablets
Cardboard box with 10 blisters of 10 tablets
Cardboard box with 18 blisters of 10 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
CEVA Sante Animale B.V.

Marketing authorisation date:
29/06/2012

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 109555

Date of authorisation status change:
14/04/2022

Reference member state:
France

Procedure number:
FR/V/0235/002

Concerned member states:
Austria Belgium Finland Germany Greece Ireland Italy Luxembourg Netherlands Poland Portugal Spain 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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