PRILACTONE NEXT 50 MG CHEWABLE TABLETS FOR DOGS

Authorised

- Spironolactone

Product identification

Medicine name:
PRILACTONE NEXT 50 MG CHEWABLE TABLETS FOR DOGS

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC03DA01

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

Authorisation status:
Valid

Authorised in:
Finland

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

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

Marketing authorisation date:
30/01/2013

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Finnish Medicines Agency

Authorisation number:
29734

Date of authorisation status change:
30/01/2013

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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