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PRILACTONE NEXT 50 MG CHEWABLE TABLETS FOR DOGS

Authorised

• Spironolactone

Product identification

Medicine name:

PRILACTONE NEXT 50 MG CHEWABLE TABLETS FOR DOGS

Prilactone vet 50 mg purutabletti

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 <u>French</u>				
Available only in <u>French</u>				
Available only in French				
Available only in <u>French</u> Available only in French				
Entitlement type:				
Entitlement type: Marketing Authorisation Legal basis of product authorisation:	2001/82/EC)			
Entitlement type: Marketing Authorisation Legal basis of product authorisation: Full application - Known active substance (Article 12(3) of Directive No 2 Marketing authorisation holder:	2001/82/EC)			
Entitlement type: Marketing Authorisation Legal basis of product authorisation: Full application - Known active substance (Article 12(3) of Directive No 2 Marketing authorisation holder: Ceva Sante Animale Marketing authorisation date:	2001/82/EC)			
Additional information Entitlement type: Marketing Authorisation Legal basis of product authorisation: Full application - Known active substance (Article 12(3) of Directive No 2 Marketing authorisation holder: Ceva Sante Animale Marketing authorisation date: 30/01/2013 Manufacturing sites for batch release: Ceva Sante Animale	2001/82/EC)			

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

Combined File of all Documents		