Procox 18 mg/ml + 0.9 mg/ml - Oral suspension

Authorised

- Emodepside
- Toltrazuril

Product identification

Medicine name:
Procox 18 mg/ml + 0.9 mg/ml - Oral suspension

Active substance:
Emodepside

Toltrazuril

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Emodepside
0.09 gram(s) / 1.00 Bottle

Toltrazuril

1.80 gram(s) / 1.00 Bottle

Pharmaceutical form:
Oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AX60

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

Authorisation status:
Valid

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:
Austria , Belgium , Bulgaria , Czechia , Finland , France , Germany , Greece , Hungary , Italy , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Spain

Package description:
Packaging:Bottle (glass), Package_size:1 bottle, Content:20 ml
Packaging:Bottle (glass), Package_size:1 bottle, Content:7.5 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol SA

Marketing authorisation date:

20/04/2011

Manufacturing sites for batch release:
Vetoquinol Biowet Sp. z o.o.
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
20/04/2011

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)
Published on: 1/02/2024
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