Profender 15 mg + 3 mg - Modified-release tablet

Authorised

- Praziquantel
- Emodepside

Product identification

Medicine name:
Profender 15 mg + 3 mg - Modified-release tablet

Active substance:
Praziquantel
Emodepside

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Praziquantel
15.00 milligram(s) / 1.00 Tablet
Emodepside

3.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Modified-release tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA51

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 , France , Germany , Portugal , Spain

Package description:
Packaging:Blister (Alu/Alu), Package_size:50 tablets
Packaging:Blister (Alu/Alu), Package_size:24 tablets
Packaging:Blister (Alu/Alu), Package_size:10 tablets
Packaging:Blister (Alu/Alu), Package_size:4 tablets
Packaging:Blister (Alu/Alu), Package_size:2 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Vetoquinol SA

Marketing authorisation date:

27/07/2005

Manufacturing sites for batch release:

Vetoquinol S.A.
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KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:
European Commission

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

Date of authorisation status change:
25/08/2008

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

Documents

Combined File of all Documents

English (PDF)

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