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Aderexa 12.5 mg/125 mg Chewable Tablets for Dogs Weighing at Least 5 kg

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
Aderexa 12.5 mg/125 mg Chewable Tablets for Dogs Weighing at Least 5 kg

Active substance:
Milbemycin oxime

Praziquantel

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Milbemycin oxime
12.50 milligram(s) / 1.00 Tablet

Praziquantel
125.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AB51

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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Package description:
Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil.Cardboard box with 12 blisters, each blister contains 4 tablets (total 48 tablets).
Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil.Cardboard box with 1 blister of 4 tablets.
Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil.Cardboard box with 1 blister of 2 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
5/04/2019

Manufacturing sites for batch release:
KRKA tovarna zdravil d.d. Novo mesto
TAD Pharma GmbH

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 01656/4178

Date of authorisation status change:
14/05/2024

Reference member state:
Ireland

Procedure number:
IE/V/0528/002

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics