

Metronidavet, 500 mg tablets for dogs and cats

Authorised

- Metronidazole

Product identification

Medicine name:

Metronidavet, 500 mg tablets for dogs and cats

Active substance:

Metronidazole

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Metronidazole

500.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01XD01

QP51AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

Package description:

PVC - Aluminium - oPA (oriented polyamide) / heat seal lacquer - Aluminium blister packed in cardboard box. Cardboard box with 250 tablets (25 blisters of 10 tablets).

PVC - Aluminium - oPA (oriented polyamide) / heat seal lacquer - Aluminium blister packed in cardboard box. Cardboard box with 20 tablets (2 blisters of 10 tablets).

PVC - Aluminium - oPA (oriented polyamide) / heat seal lacquer - Aluminium blister packed in cardboard box. Cardboard box with 100 tablets (10 blisters of 10 tablets).

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Przedsiębiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Marketing authorisation date:

19/04/2021

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 126483

Date of authorisation status change:

27/01/2022

Reference member state:

Netherlands

Procedure number:

NL/V/0324/002

Concerned member states:

Austria Belgium Bulgaria Czechia France Greece Hungary Italy Lithuania
Poland Portugal Romania Spain

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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