Optivermin 50 mg + 500 mg Tabletka

Not authorised

This information is not available for this product.

Product identification

Medicine name:
Optivermin 50 mg + 500 mg Tabletka

Active substance:
This information is not available for this product.

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
This information is not available for this product.

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AA51

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

Authorisation status:
Surrendered

Authorised in:
Poland

Package description:
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetoquinol Biowet Sp. z o.o.

Marketing authorisation date:
29/04/2004

Manufacturing sites for batch release:
Vetoquinol Biowet Sp. z o.o.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
1525

Date of authorisation status change:
29/04/2004

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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