

Tenaver 720 mg + 22,42 mg Tabletka

Authorised

- Niclosamide
- Levamisole hydrochloride

Product identification

Medicine name:

Tenaver 720 mg + 22,42 mg Tabletka

Active substance:

Niclosamide

Levamisole hydrochloride

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Niclosamide

720.00 milligram(s) / 1.00 Tablet

Levamisole hydrochloride

22.42 milligram(s) / 1.00 milligram(s)

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AE51

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

Available only in [Polish](#)

Available only in [Polish](#)

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:

Biowet Drwalew Sp. z o.o.

Marketing authorisation date:

3/12/1997

Manufacturing sites for batch release:

Drwalewskie Zakłady Przemysłu Bioweterynaryjnego S.A.

Responsible authority:

Authorisation number:

0420

Date of authorisation status change:

3/12/1997

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

Documents

Package Leaflet

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Summary of Product Characteristics

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

Labelling

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