

Tensurin 20 mg kauwtabletten voor honden

Authorised

- Phenylpropanolamine hydrochloride

Product identification

Medicine name:

Tensurin 20 mg kauwtabletten voor honden

Active substance:

Phenylpropanolamine hydrochloride

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Phenylpropanolamine hydrochloride
20.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QG04BX91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

Package description:

Available only in [Dutch](#)

Available only in [Dutch](#)

Available only in [Dutch](#)

Available only in [Dutch](#)

Available only in [Dutch](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Ast Farma B.V.

Marketing authorisation date:

19/04/2007

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 10179

Date of authorisation status change:

10/01/2019

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

Documents

Combined File of all Documents

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