

Digoxine 0,25 mg, tabletten voor honden en katten

Authorised

- Digoxin

Product identification

Medicine name:

Digoxine 0,25 mg, tabletten voor honden en katten

Active substance:

Digoxin

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Digoxin

0.25 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Withdrawal period by route of administration:

Oral use:

-

Dog

-

Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:QC01AA05

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

Authorisation status:Valid

Authorised in:Netherlands

Package description:Available only in DutchAvailable 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:Alfasan Nederland B.V.

Marketing authorisation date:11/05/1992

Manufacturing sites for batch release:KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 4559

Date of authorisation status change:

5/07/2018

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.

Source URL: <https://medicines.health.europa.eu/veterinary/600000067552>