

Prednisolon 5 mg, tabletten voor honden en katten

Authorised

- Prednisolone

Product identification

Medicine name:

Prednisolon 5 mg, tabletten voor honden en katten

Active substance:

Prednisolone

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Prednisolone

5.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

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:

Alfasan Nederland B.V.

Marketing authorisation date:

23/04/1992

Manufacturing sites for batch release:

KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 4749

Date of authorisation status change:

6/10/2021

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

Documents

Combined File of all Documents

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