Prednoral 100 mg tabletten met smaakstof voor honden

Authorised

- Prednisolone

Product identification

Medicine name:
Prednoral 100 mg tabletten met smaakstof voor honden

Active substance:
Prednisolone

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Prednisolone
100.00 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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ast Farma B.V.

Marketing authorisation date:
9/03/2010

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 104350

Date of authorisation status change:

16/04/2018

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.