

Prednisolone 2,5% suspensie voor injectie voor honden

Authorised

- Prednisolone

Product identification

Medicine name:

Prednisolone 2,5% suspensie voor injectie voor honden

Active substance:

Prednisolone

Target species:

Dog

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Prednisolone

25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

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:

8/09/1992

Manufacturing sites for batch release:

KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 5946

Date of authorisation status change:

29/12/2021

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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