File downloaded on 2026-11-03
Source URL: https://medicines.health.europa.eu/veterinary/en/600000086392

Prednisolone 1% 10 mg/ml Oplossing voor injectie

Authorised

- Prednisolone

Product identification

Medicine name:
Prednisolone 1% 10 mg/ml Oplossing voor injectie

Active substance:
Prednisolone

Target species:
Dog

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Prednisolone
10.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB04

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Prednisolone 1% 25 ml Vial Solution for injection
Prednisolone 1% 50 ml Vial Solution for injection
Prednisolone 1% 100 ml Vial Solution for injection

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
V.M.D.

Marketing authorisation date:
1/01/1973

Manufacturing sites for batch release:
V.M.D.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V093387

Date of authorisation status change:
6/03/2024

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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