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Prednisolone 20 mg Kela 20 mg Tablet

Authorised

- Prednisolone

Product identification

Medicine name:
Prednisolone 20 mg Kela 20 mg Tablet

Active substance:
Prednisolone

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Prednisolone
20.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD07BA04

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

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Prednisolone 20 mg Kela 100 Tablets in 10 blisters with 10 tablets
Prednisolone 20 mg Kela 50 Tablets in 5 blisters with 10 tablets
Prednisolone 20 mg Kela 30 Tablets in 3 blisters with 10 tablets

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
24/11/2008

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V329095

Date of authorisation status change:
20/01/2021

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

Documents

Summary of Product Characteristics

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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.