Prednisolone 5 mg Kela 5 mg Tablet

Authorised

- Prednisolone

Product identification

Medicine name:
Prednisolone 5 mg Kela 5 mg Tablet

Active substance:
Prednisolone

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Prednisolone
5.00 milligram(s) / 200000.00 milligram(s)

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 5 mg Kela 100 Tablets in Blister
Prednisolone 5 mg Kela 50 Tablets in Blister
Prednisolone 5 mg Kela 30 Tablets in Blister

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
23/12/1998

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V199516

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

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Labelling

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