Cefalexine Kela 250 mg Tablet

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Cefalexine Kela 250 mg Tablet

Active substance:
Cefalexin monohydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
263.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01DB01

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

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
Cefalexine Kela 250 mg Tablet Box with 10 blisters of 10 tablets
Cefalexine Kela 250 mg Tablet Box with 1 blister of 10 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:
19/11/2007

Manufacturing sites for batch release:
KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V305252

Date of authorisation status change:
27/11/2018

Generic of:
600000085336

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

Documents

Package Leaflet

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Summary of Product Characteristics

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Labelling

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