RILEXINE 200 MG/G ORAL PASTE

Authorised

- Cefalexin

Product identification

Medicine name:
RILEXINE 200 MG/G ORAL PASTE

Active substance:
Cefalexin

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin
200.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral paste

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01D

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Box with one aluminium sachet containing one graduated syringe of 4.2 g of oral paste (14 doses): each dose can treat 2 kg
Box with one aluminium sachet containing one graduated syringe of 10.5 g of oral paste (7 doses): each dose can treat 10 kg
Box with one aluminium sachet containing one graduated syringe of 21 g of oral paste (7 doses): each dose can treat 20 kg

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:
Virbac

Marketing authorisation date:
22/09/2005

Manufacturing sites for batch release:
Virbac

Responsible authority:
Ministry Of Health

Authorisation number:
102729

Date of authorisation status change:
22/09/2005

Reference member state:
Italy

Procedure number:
IT/V/0119/001

Concerned member states:
Czechia Greece Hungary Poland Slovakia Spain

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

Documents

Combined File of all Documents