Rilexine 600 mg Tablets for dogs

Authorised

- Cefalexin monohydrate

Product identification

Medicine name:
Rilexine 600 mg Tablets for dogs

Active substance:
Cefalexin monohydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Cefalexin monohydrate
631.04 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:
Ireland

Available in:
Ireland

Package description:
Blister packs consisting of blister aluminium - PVC/aluminium/OPA. Aluminium foil lid coated with lacquer.Cardboard Box with 30 blisters of 7 tablets.
Blister packs consisting of blister aluminium - PVC/aluminium/OPA. Aluminium foil lid coated with lacquer. Cardboard Box with 20 blisters of 7 tablets.
Blister packs consisting of blister aluminium - PVC/aluminium/OPA. Aluminium foil lid coated with lacquer.Cardboard Box with 2 blisters of 7 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent (abridged application) - Council Directive 81/851/EEC

Marketing authorisation holder:
Virbac

Marketing authorisation date:
21/09/2001

Manufacturing sites for batch release:
Virbac

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10988/018/003

Date of authorisation status change:
21/09/2001

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

Documents

Summary of Product Characteristics