

Clindabactin 440 mg Chewable Tablets for Dogs

Authorised

- Clindamycin hydrochloride

Product identification

Medicine name:

Clindabactin 440 mg Chewable Tablets for Dogs
Clindabactin 440 mg, kramtomosios tabletės šunims

Active substance:

Clindamycin hydrochloride

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Clindamycin hydrochloride
440.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01FF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Lithuania

Package description:

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 8 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 6 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 4 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 3 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 25 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 1 blister of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 9 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 7 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 5 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 2 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 10 blisters of 10 tablets.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Dechra Regulatory B.V.

Marketing authorisation date:

19/05/2019

Manufacturing sites for batch release:

Lelypharma B.V.

Genera d.d.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/19/2530/001-012

Date of authorisation status change:

19/05/2019

Reference member state:

Netherlands

Procedure number:

NL/V/0317/003

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland
France Germany Hungary Iceland Ireland Italy Latvia Lithuania
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain
Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

RV2530.pdf

Clindabactin 55 220 en 440 mg - Puar.pdf