Clindabactin 220 mg Chewable Tablets for Dogs

Authorised

- Clindamycin hydrochloride

Product identification

Medicine name:
Clindabactin 220 mg Chewable Tablets for Dogs

Active substance:
Clindamycin hydrochloride

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Clindamycin hydrochloride
220.00 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:
Netherlands

Available in:
Netherlands

Package description:
Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 9 blisters of 10 tablets.
Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 8 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 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 2 blisters of 10 tablets.
Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 10 blisters of 10 tablets.
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 6 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 4 blisters of 10 tablets.

Aluminium - Polyamide/Aluminium/PVC blister. Cardboard box of 1 blister 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:
18/06/2019

Manufacturing sites for batch release:
Lelypharma B.V.
Genera d.d.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 122622

Date of authorisation status change:
19/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0317/002

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)

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Documents

Combined File of all Documents

English (PDF)
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