Clindabuc vet. 75 mg tabletti

Not authorised

- Clindamycin

Product identification

Medicine name:
Clindabuc vet. 75 mg tabletti

Active substance:
Clindamycin

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Clindamycin
75.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Dog**

- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FF01

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

Authorisation status:
Surrendered

Authorised in:
Finland

Package description:
Available only in Finnish
Available only in Finnish
Available only in Finnish
Available only in Finnish
Available only in Finnish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic (abridged application) - art 10(1)

Marketing authorisation holder:
CEVA Sante Animale B.V.

Marketing authorisation date:
31/08/1998

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:

Finnish Medicines Agency

Authorisation number:
12953

Date of authorisation status change:
12/12/2023

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.
