Doxybactin 200 mg

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
Doxybactin 200 mg

Active substance:
Doxycycline hyclate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Doxycycline hyclate
230.83 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA02

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

Authorisation status:
Valid

Authorised in:
Cyprus

Available in:
Cyprus

Package description:
Cardboard box of 10 Aluminium - PVC/PE/PVDC blisters of 10 tablets
Cardboard box containing 10 separate cardboard boxes, each containing 1 Aluminium - PVC/PE/PVDC blister of 10 tablets
Cardboard box of 1 Aluminium - PVC/PE/PVDC blister of 10 tablets
Cardboard box of 2 Aluminium - PVC/PE/PVDC blisters of 10 tablets
Cardboard box of 3 Aluminium - PVC/PE/PVDC 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:
23/07/2017

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

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00626V

Date of authorisation status change:
11/01/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0218/003

Concerned member states:
Austria Belgium Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece 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

English (PDF)
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