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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:
Germany

Available in:
Germany

Package description:
Cardboard box of 3 Aluminium - PVC/PE/PVDC blisters of 10 tablets
Cardboard box of 2 Aluminium - PVC/PE/PVDC blisters of 10 tablets
Cardboard box of 1 Aluminium - PVC/PE/PVDC blister of 10 tablets
Cardboard box containing 10 separate cardboard boxes, each containing 1 Aluminium - PVC/PE/PVDC blister of 10 tablets
Cardboard box of 10 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:
17/08/2017

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

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402374.00.00

Date of authorisation status change:
4/10/2022

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)

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Documents

Combined File of all Documents

English (PDF)
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