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Doxycare Flavour 200 mg Tablets for Cats and Dogs

Authorised

- Doxycycline hyclate

Product identification

Medicine name:
Doxycare Flavour 200 mg Tablets for Cats and Dogs

Active substance:
Doxycycline hyclate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Doxycycline hyclate
239.40 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

Package description:
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 90 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 100 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 250 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 10 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 20 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 30 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 40 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 50 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 60 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 70 tablets.
OPA/Aluminium/PVC foil and Aluminium foil blister containing 10 tabletsCardboard box of 80 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:
Ecuphar

Marketing authorisation date:
18/01/2022

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402578.00.00

Date of authorisation status change:
18/01/2022

Reference member state:
Ireland

Procedure number:
IE/V/0645/002

Concerned member states:
Austria Belgium Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Spain Sweden United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics