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
Doxycare, 200mg, Tableta

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

Withdrawal period by route of administration:

Oral use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Czechia

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:
3/10/2019

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/079/19-C

Date of authorisation status change:
3/10/2019

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)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 13/04/2025
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Package Leaflet

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Labelling

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Combined File of all Documents