

# 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

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**Active substance:**

Doxycycline hyclate

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**Target species:**

Dog  
Cat

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Doxycycline hyclate  
239.40 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Tablet

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**Withdrawal period by route of administration:****Oral use:**

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**Dog**

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**Cat**

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**QJ01AA02

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

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**Authorisation status:**Valid

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**Authorised in:**Czechia

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ecuphar

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**Marketing authorisation date:**

3/10/2019

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**Manufacturing sites for batch release:**

Lelypharma B.V.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/079/19-C

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**Date of authorisation status change:**

3/10/2019

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0645/002

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**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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[www.adrreports.eu/vet](http://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

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

### Combined File of all Documents

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