

# Doxybactin 200 mg

Authorised

- Doxycycline hyclate

## Product identification

**Medicine name:**

Doxybactin 200 mg

Doxybactin vet 200 mg töflur handa hundum

**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

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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:**

Iceland

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**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

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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:**

Dechra Regulatory B.V.

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**Marketing authorisation date:**

2/08/2017

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**Manufacturing sites for batch release:**

Lelypharma B.V.

Genera d.d.

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**Responsible authority:**

Icelandic Medicines Agency

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**Authorisation number:**

IS/2/17/009/02

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**Date of authorisation status change:**

2/08/2017

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0218/003

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**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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