

# Twinox 400 mg/100 mg chewable tablets for dogs

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

## Product identification

**Medicine name:**

Twinox 400 mg/100 mg chewable tablets for dogs

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**Active substance:**

Amoxicillin trihydrate

Potassium clavulanate

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Amoxicillin trihydrate

459.22 milligram(s) / 1.00 Tablet

Potassium clavulanate

119.13 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Chewable tablet

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

QJ01CR02

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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:**

Germany

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**Available in:**

Germany

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**Package description:**

Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating. Blister contains 6 tablets. Carton contains 12 tablets.

Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating. Blister contains 6 tablets. Carton contains 60 tablets.

Blister formed of an aluminium foil which consists of aluminum layer coated with OPA (Oriented Polyamide) film on one side and PE with desiccant on the other side and an aluminium sealing foil which consists of aluminium layer and PE coating. Blister contains 6 tablets. Carton contains 300 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:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

14/06/2021

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

V7001211.00.00

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**Date of authorisation status change:**

14/06/2021

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0656/003

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**Concerned member states:**

Austria Belgium France Germany Italy Netherlands Portugal

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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