

# Twinox 40 mg/10 mg chewable tablets for cats and dogs

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

## Product identification

**Medicine name:**

Twinox 40 mg/10 mg chewable tablets for cats and dogs

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**Active substance:**

Amoxicillin trihydrate

Potassium clavulanate

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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:**

Amoxicillin trihydrate

45.92 milligram(s) / 1.00 Tablet

Potassium clavulanate  
11.91 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 10 tablets. Carton contains 10 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 10 tablets. Carton contains 20 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 10 tablets. Carton contains 100 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 10 tablets. Carton contains 500 tablets.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) 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:**

V7001209.00.00

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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/0656/001

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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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