

# TUDOMAX 10 MG/G, POWDER FOR USE IN DRINKING WATER/MILK

Authorised

- Bromhexine hydrochloride

## Product identification

**Medicine name:**

TUDOMAX 10 MG/G, POWDER FOR USE IN DRINKING WATER/MILK

TUDOMAX 10 mg/g, powder for use in drinking water/milk

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**Active substance:**

Bromhexine hydrochloride

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**Target species:**

Turkey

Pig

Cattle (calf)

Duck

Chicken

Chicken (broiler)

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Bromhexine hydrochloride  
10.98 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Powder for use in drinking water/milk

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**Withdrawal period by route of administration:**

**Oral use:**

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**Turkey**

- Meat and offal. 0 day
- Eggs. no withdrawal period

Not for use in birds producing eggs for consumption, during and 4 weeks before the laying phase.

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**Pig**

- Meat and offal. 0 day

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**Cattle (calf)**

- Meat and offal. 2 day
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

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**Duck**

- Meat and offal. 0 day
- Eggs. no withdrawal period

Not for use in birds producing eggs for consumption, during and 4 weeks before the laying phase.

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**Chicken**

- Eggs. no withdrawal period

Not for use in birds producing eggs for consumption, during and 4 weeks before the laying phase.

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**Chicken (broiler)**

- Meat and offal. 0 day

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

QR05CB02

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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:**

Bulgaria

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**Package description:**

Available only in French

Available only in French

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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:**

S P Veterinaria S.A.

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**Marketing authorisation date:**

13/02/2017

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**Manufacturing sites for batch release:**

S P Veterinaria S.A.

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-2709

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**Date of authorisation status change:**

13/02/2017

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**Reference member state:**

France

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**Procedure number:**

FR/V/0295/001

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**Concerned member states:**

Bulgaria Cyprus Greece Hungary Ireland Italy Malta Poland Portugal  
Romania Spain United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

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

## Package Leaflet

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

## Labelling

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