

# SULFACOX T perorálny roztok

Authorised

- Trimethoprim
- Sulfadimidine sodium

## Product identification

**Medicine name:**

SULFACOX T perorálny roztok

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**Active substance:**

Trimethoprim

Sulfadimidine sodium

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**Target species:**

Cattle

Cattle (calf)

Sheep

Sheep (lamb)

Pig

Pig (piglet)

Dog

Poultry

Pheasant

Rabbit

Hare

Fur animals

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**Route of administration:**

In drinking water use

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## Product details

### **Active substance and strength:**

Trimethoprim

12.50 milligram(s) / 1.00 millilitre(s)

Sulfadimidine sodium

50.00 milligram(s) / 1.00 millilitre(s)

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### **Pharmaceutical form:**

Oral solution

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### **Withdrawal period by route of administration:**

#### **In drinking water use:**

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#### **Cattle**

- Meat and offal. 15 day

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#### **Cattle (calf)**

- Meat and offal. 15 day

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#### **Sheep**

- Meat and offal. 15 day

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#### **Sheep (lamb)**

- Meat and offal. 15 day

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#### **Pig**

- Meat and offal. 15 day

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#### **Pig (piglet)**

- Meat and offal. 15 day

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**Poultry**

- Meat and offal. 15 day

Do not use in laying hens producing eggs for human consumption.

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**Pheasant**

- Meat and offal. 15 day

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**Rabbit**

- Meat and offal. 15 day

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**Hare**

- Meat and offal. 15 day

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

QJ01EW03

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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:**

Slovakia

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**Package description:**

Available only in Slovak

Available only in Slovak

Available only in Slovak

Available only in Slovak

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Pharmagal spol. s r.o.

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**Marketing authorisation date:**

20/03/2001

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**Manufacturing sites for batch release:**

Pharmagal spol. s r.o.

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/017/01-S

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**Date of authorisation status change:**

20/03/2001

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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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