

## T.S.-sol 20/100

Authorised

- Trimethoprim
- Sulfamethoxazole

## Product identification

### **Medicine name:**

T.S.-sol 20/100

T.S. Sol 20 mg/ml Oplossing voor gebruik in drinkwater

T.S. Sol 20 mg/ml Solution pour administration dans l'eau de boisson

T.S. Sol 20 mg/ml Lösung zum Eingeben über das Trinkwasser

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### **Active substance:**

Trimethoprim

Sulfamethoxazole

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### **Target species:**

Pig (for fattening)

Chicken (broiler)

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### **Route of administration:**

In drinking water use

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## Product details

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

Trimethoprim

20.00 milligram(s) / 1.00 millilitre(s)

Sulfamethoxazole

100.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for use in drinking water

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**Withdrawal period by route of administration:**

**In drinking water use:**

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**Pig (for fattening)**

- Meat and offal. 8 day

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**Chicken (broiler)**

- Meat and offal. 5 day

Not for use in birds producing eggs for human consumption

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

QJ01EW11

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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:**

Belgium

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**Available in:**

Belgium

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**Package description:**

High-density polyethylene bottles with low-density polyethylene screw cap containing 1 litre of product

High-density polyethylene jerrycan with high-density polyethylene screw cap containing 5 litres of product

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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:**

Dopharma Research B.V.

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**Marketing authorisation date:**

11/12/2017

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**Manufacturing sites for batch release:**

Dopharma B.V.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

11/12/2017

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0213/001

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**Concerned member states:**

Belgium Croatia France Germany Greece Hungary Italy Poland Romania  
Spain United Kingdom (Northern Ireland)

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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

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.