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

Active substance:
Trimethoprim

Sulfamethoxazole

Target species:
Pigs (for fattening)
Chicken (broiler)

Route of administration:
In drinking water use

Product details

Active substance and strength:
Trimethoprim
20.00 milligram(s) / 1.00 millilitre(s)

Sulfamethoxazole

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **Pigs (for fattening)**
  - Meat and offal. 8 day
- **Chicken (broiler)**
  - Meat and offal. 5 day

Not for use in birds producing eggs for human consumption

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EW11

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dopharma Research B.V.

Marketing authorisation date:
11/12/2017

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
11/12/2017

Reference member state:
Netherlands

Procedure number:
NL/V/0213/001

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

To consult adverse reactions on veterinary medicinal products please go to 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.
