# Prazil N Orale 200 mg/ml + 40 mg/ml, soluzione orale per bovini, suini, broilers, tacchini e conigli

Authorised

- Sulfadimethoxine
- Trimethoprim

## Product identification

**Medicine name:**
Prazil N Orale 200 mg/ml + 40 mg/ml, soluzione orale per bovini, suini, broilers, tacchini e conigli

**Active substance:**
Sulfadimethoxine
Trimethoprim

**Target species:**
Cattle (calf)
Pig
Pig (for fattening)
Turkey
Chicken (broiler)
Rabbit

**Route of administration:**
In drinking water/milk use
In drinking water use

## Product details

**Active substance and strength:**
Sulfadimethoxine
20.00 gram(s) / 100.00 millilitre(s)

Trimethoprim
4.00 gram(s) / 100.00 millilitre(s)

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**Pharmaceutical form:**
Oral solution

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**Withdrawal period by route of administration:**
**In drinking water/milk use:**

- **Cattle (calf)**
  - Meat and offal. 20 day

Uso non autorizzato in uccelli che producono uova per consumo umano

- **Pig**
  - Meat and offal. 8 day

Uso non autorizzato in uccelli che producono uova per consumo umano

- **Pig (for fattening)**
  - Meat and offal. 8 day

Uso non autorizzato in uccelli che producono uova per consumo umano

**In drinking water use:**

- **Turkey**
  - Meat and offal. 9 day

Uso non autorizzato in uccelli che producono uova per consumo umano

- **Chicken (broiler)**
  - Meat and offal. 4 day

Uso non autorizzato in uccelli che producono uova per consumo umano

- **Rabbit**
  - Meat and offal. 6 day

Uso non autorizzato in uccelli che producono uova per consumo umano

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01EW09

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

**Authorisation status:**
Valid

**Authorised in:**
Italy

**Package description:**
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Full application (Article 12(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**

Dopharma Research B.V.

**Marketing authorisation date:**
14/09/1979

**Manufacturing sites for batch release:**
Boehringer Ingelheim Animal Health France

**Responsible authority:**
Ministry Of Health

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

**Date of authorisation status change:**
8/05/2002

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Combined File of all Documents

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