

NEOPRIDIMET soluzione orale, 200 mg + 40 mg/ml, per uso in acqua da bere per polli da carne e conigli

Authorised

- Trimethoprim
- Sulfadiazine

Product identification

Medicine name:

NEOPRIDIMET soluzione orale, 200 mg + 40 mg/ml, per uso in acqua da bere per polli da carne e conigli

Active substance:

Trimethoprim

Sulfadiazine

Target species:

Rabbit

Chicken (broiler)

Route of administration:

Oral use

Product details

Active substance and strength:

Trimethoprim

40.00 milligram(s) / 1.00 millilitre(s)

Sulfadiazine

200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for use in drinking water

Withdrawal period by route of administration:

Oral use:

-

Rabbit

- Meat and offal. 5 day

-

Chicken (broiler)

- Meat and offal. 5 day

Uso non consentito in galline ovaiole che producono uova destinate al consumo umano

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01EW10

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Available in:

Italy

Package description:

Available only in [Italian](#)

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:

Fatro S.p.A.

Marketing authorisation date:

9/10/2000

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

9/10/2010

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

Documents

Combined File of all Documents

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