

PAN TERRAMICINA

Authorised

- Oxytetracycline

Product identification

Medicine name:

PAN TERRAMICINA

Active substance:

Oxytetracycline

Target species:

Dog

Horse

Cat

Turkey

Chicken (broiler)

Chicken

Cattle

Route of administration:

Ocular use

Intramuscular use

Intravenous use

Subcutaneous use

Cutaneous use

Intrauterine use

Product details

Active substance and strength:

Oxytetracycline

30.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Ocular use:

-

Horse

- Meat and offal. 20 day

Usò non autorizzato in equidi che producono latte per il consumo umano

Intramuscular use:

-

Horse

- Meat and offal. 20 day

Usò non autorizzato in equidi che producono latte per il consumo umano

Intravenous use:

-

Horse

- Meat and offal. 20 day

Usò non autorizzato in equidi che producono latte per il consumo umano

Subcutaneous use:

-

Turkey

- Meat and offal. 15 day

Usò non autorizzato in uccelli che producono uova per il consumo umano

-

Chicken (broiler)

- Meat and offal. 15 day

Usò non autorizzato in uccelli che producono uova per il consumo umano

-

Chicken

- Meat and offal. 15 day

Usò non autorizzato in uccelli che producono uova per il consumo umano

-

Horse

- Meat and offal. 20 day

Usò non autorizzato in equidi che producono latte per il consumo umano

Cutaneous use:

-

Horse

- Meat and offal. 20 day

Usò non autorizzato in equidi che producono latte per il consumo umano

Intrauterine use:

-

Cattle

- Milk. 12 hour

- Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA06

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](#)

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:

Zoetis Italia S.r.l

Marketing authorisation date:

14/07/1960

Manufacturing sites for batch release:

Fareva Amboise

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

8/03/1989

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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