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

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

Intramuscular use:

- **Horse**
  - Meat and offal. 20 day

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

Intravenous use:

- **Horse**
  - Meat and offal. 20 day

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

Subcutaneous use:

- **Turkey**
  - Meat and offal. 15 day

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

- **Chicken (broiler)**
  - Meat and offal. 15 day

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

- **Chicken**
  - Meat and offal. 15 day

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

- **Horse**
  - Meat and offal. 20 day

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

Cutaneous use:

- **Horse**
  - Meat and offal. 20 day

Uso 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

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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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