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DAIMETON 40 soluzione iniettabile al 40% per bovini, equini, suini, polli da carne, tacchini, cani e gatti

Authorised

- Sulfamonomethoxine sodium

Product identification

Medicine name:
DAIMETON 40 soluzione iniettabile al 40% per bovini, equini, suini, polli da carne, tacchini, cani e gatti

Active substance:
Sulfamonomethoxine sodium

Target species:
Turkey
Cattle
Chicken (broiler)
Dog
Cat
Pig
Horse

Route of administration:
Intramuscular use

Intravenous use

Product details

Active substance and strength:
Sulfamonomethoxine sodium
43.00 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Turkey**
  - Meat and offal. 28 day

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

- **Cattle**
  - Milk. 72 hour
  - Meat and offal. 11 day

- **Chicken (broiler)**
  - Meat and offal. 28 day

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

- **Pig**
  - Meat and offal. 16 day

Intravenous use:

- **Turkey**

- Meat and offal. 28 day

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

- **Cattle**
  - Milk. 72 hour
  - Meat and offal. 11 day
- **Chicken (broiler)**
  - Meat and offal. 28 day

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

- **Pig**
  - Meat and offal. 16 day
- **Horse**
  - Meat and offal. 35 day

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

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EQ18

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

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Authorisation status:
Valid

Authorised 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:
Izo S.r.l.

Marketing authorisation date:
23/04/1985

Manufacturing sites for batch release:
Izo S.r.l.

Responsible authority:
Ministry Of Health

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

Date of authorisation status change:
1/10/1996

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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