

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01EQ18

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

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

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