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SADIMET

Authorised

- Sulfadimethoxine

Product identification

Medicine name:
SADIMET

Active substance:
Sulfadimethoxine

Target species:
Cattle
Sheep
Pig
Horse

Route of administration:
Oral use
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Sulfadimethoxine
330.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Oral use:

- **Cattle**
  - Meat and offal. 35 day
  - Milk. 672 hour
- **Sheep**
  - Meat and offal. 35 day
  - Milk. 192 hour
- **Pig**
  - Meat and offal. 35 day
- **Horse**
  - Meat and offal. 35 day

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

Intramuscular use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 264 hour
- **Sheep**
  - Meat and offal. 14 day
  - Milk. 264 hour
-

Pig

- Meat and offal. 14 day

•

Horse

- Meat and offal. 14 day

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

Intravenous use:

•

Cattle

- Meat and offal. 6 day
- Milk. 60 hour

•

Sheep

- Meat and offal. 10 day
- Milk. 192 hour

•

Pig

- Meat and offal. 10 day

•

Horse

- Meat and offal. 9 hour

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

Subcutaneous use:

•

Cattle

- Meat and offal. 14 day
- Milk. 264 hour

•

Sheep

- Meat and offal. 14 day
- Milk. 264 hour

- **Pig**
  - Meat and offal. 14 day

- **Horse**
  - Meat and offal. 14 day

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EQ09

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

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:
16/05/1990

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:
16/05/1990

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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