PAMIZOLE L

Authorised

- Levamisole
- Levamisole hydrochloride

Product identification

Medicine name:
PAMIZOLE L

Active substance:
Levamisole
Levamisole hydrochloride

Target species:
Cattle
Sheep
Pig
Buffalo (male)

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Levamisole

75.00 milligram(s) / 1.00 millilitre(s)

Levamisole hydrochloride
88.40 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. no withdrawal period

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

- **Sheep**
  - Meat and offal. no withdrawal period

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

- **Pig**
  - Meat and offal. no withdrawal period

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

- **Buffalo (male)**
  - Meat and offal. no withdrawal period

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

Subcutaneous use:

- **Cattle**
  - Meat and offal. no withdrawal period

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

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Sheep

- Meat and offal. no withdrawal period

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

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Buffalo (male)

- Meat and offal. no withdrawal period

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AE01

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:
Fatro S.p.A.

Marketing authorisation date:
2/01/1987

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:
2/01/1987

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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