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ALAMYCIN LA 300

Authorised

- Oxytetracycline

Product identification

Medicine name:
ALAMYCIN LA 300

Active substance:
Oxytetracycline

Target species:
Cattle
Sheep
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Oxytetracycline
300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 8 day
  - Meat and offal. 28 day
  - Meat and offal. 35 day giorni (alto dosaggio)
- **Cattle**
  - Milk. 8 day
  - Meat and offal. 28 day
  - Meat and offal. 35 day giorni (alto dosaggio)
- **Cattle**
  - Milk. 8 day
  - Meat and offal. 28 day
  - Meat and offal. 35 day giorni (alto dosaggio)
- **Sheep**
  - Milk. 8 day
  - Meat and offal. 28 day
  - Meat and offal. 28 day giorni (alto dosaggio)
- **Sheep**
  - Milk. 8 day
  - Meat and offal. 28 day
  - Meat and offal. 28 day giorni (alto dosaggio)

- **Sheep**
  - Milk. 8 day
  - Meat and offal. 28 day
  - Meat and offal. 28 day giorni (alto dosaggio)
- **Pig**
  - Meat and offal. 14 day
  - Meat and offal. 28 day giorni (alto dosaggio)
- **Pig**
  - Meat and offal. 14 day
  - Meat and offal. 28 day giorni (alto dosaggio)

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA06

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

Authorisation status:
Valid

Authorised in:
Italy

Available 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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
16/01/2003

Manufacturing sites for batch release:
Norbrook Laboratories Limited

Responsible authority:
Ministry Of Health

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

Date of authorisation status change:
16/01/2003

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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