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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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