

Summary of Product Characteristics

1 NAME OF THE VETERINARY MEDICINAL PRODUCT

Alamycin LA Veterinary

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each ml contains:

Active substance

Oxytetracycline (as Oxytetracycline Dihydrate) 200 mg

For a full list of excipients see section 6.1.

3 PHARMACEUTICAL FORM

Solution for injection.

A clear amber solution

4 CLINICAL PARTICULARS

4.1 Target Species

Cattle, sheep, pigs.

4.2 Indications for use, specifying the target species

Antibiotic: treatment of sensitive microorganisms in Cattle, Sheep and Pigs.

In use for the following infections:

- Atrophic rhinitis caused by *Bordetella bronchiseptica*, *Mannheimia haemolytica* and *Pasteurella multocida*.
- Navel/joint ill caused by *Treuperella pyogenes*, *E. coli* or *Staphylococcus aureus*.
- Mastitis caused by *Corynebacterium pyogenes*, *E. coli*, *Staphylococcus aureus*, *Streptococcus agalactiae*, *Streptococcus uberis*.
- Metritis caused by *E. coli* or *Streptococcus pyogenes*.
- Pasteurellosis and infections of the respiratory tract caused by *Pasteurella haemolytica* and *Pasteurella multocida*.
- Septicaemia caused by *Salmonella dublin* and *Streptococcus pyogenes*.

Erysipelas caused by *Erysipelothrix rhusiopathiae*.

4.3 Contraindications

Do not use in Cats, Dogs and Horses

Do not use in animals suffering from hepatic or renal damage.

Do not use in animals with known hypersensitivity to Oxytetracycline.

4.4 Special warnings for each target species

None.

4.5 Special precautions for use

Special precautions for use in animals

Do not dilute Alamycin LA.

If concurrent treatment is administered, use a separate injection site.

Special Precautions to be taken by the Person Administering the Medicinal Product to Animals

In case of contact with eyes or skin, wash immediately with water as irritation may occur.

Wash hands after use.

Take care to avoid accidental injection.

4.6 Adverse reactions (frequency and seriousness)

Although well tolerated, occasionally a slight local reaction of a transient nature has been observed.

Any suspected adverse events should be reported to the Ministry of Health by clicking on the link below and using an online form:

<https://sideeffects.health.gov.il>

4.7 Use during pregnancy, lactation or lay

The use of tetracycline during the period of tooth and bone development, including late pregnancy may lead to tooth discoloration. Alamycin LA can be safely administered during lactation.

4.8 Interaction with other medicinal products and other forms of interaction

None known.

4.9 Amounts to be administered and administration route

The recommended dose rate is 20 mg/kg bodyweight (i.e. 1 ml per 10 kg bodyweight) administered by deep intramuscular injection.

Maximum recommended dose at any one site:

Cattle:	20 ml	
Pigs:	5.5 ml	
Sheep:	5 ml	
Piglets:	1 day	0.2 ml
	7 days	0.3 ml

14 days	0.4 ml
21 days	0.5 ml
over 21 days	1.0 ml/10 kg

4.10 Overdose (symptoms, emergency procedures, antidotes), if necessary

Not applicable.

4.11 Withdrawal Period(s)

Cattle:

Meat and offal: 35 days

Milk: 8 days

Pigs:

Meat and offal: 20 days

Sheep:

Meat and offal: 20 days

Milk: 8 days

5 PHARMACOLOGICAL or IMMUNOLOGICAL PROPERTIES

Pharmacotherapeutic group: Antibacterials for systemic use, tetracyclines

ATCvet Code: QJ01AA06

5.1 Pharmacodynamic properties

Oxytetracycline is a bacteriostatic antibiotic that inhibits protein synthesis in susceptible bacteria. Inside the cell it binds irreversibly to receptors on the 30S subunit of the bacterial ribosome where it interferes with the binding of the aminoacyl-transfer RNA to the receptor site on the messenger RNA ribosome complex. This effectively prevents the addition of amino acids to the elongating peptide chain, inhibiting protein synthesis. Alamycin LA is specifically formulated to provide a prolonged action resulting in sustained antibacterial activity.

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Sodium Formaldehyde Sulphoxylate

Magnesium Oxide Light

2-Pyrrolidone

Povidone K12

Monoethanolamine
Hydrochloric Acid
Water for Injections

6.2 Incompatibilities

Do not mix the product with other medicinal products.

6.3 Shelf-life

Shelf-life of the veterinary medicinal product as packaged for sale: 18 months

Shelf-life after first opening the immediate packaging: 28 days

6.4 Special precautions for storage

Do not store above 25°C. Protect from light.

6.5 Nature and composition of immediate packaging

Amber type II glass vials of 50 ml, and 100 ml with Chlorobutyl bungs and aluminium seals containing a clear solution for injection.

Not all pack sizes may be marketed.

6.6 Special precautions for the disposal of unused veterinary medicinal products or waste materials

Any unused product or waste materials should be disposed of in accordance with national requirements.

7 REGISTRATION HOLDER

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8 MANUFACTURER

Norbrook Laboratories LTD
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9 PRODUCT REGISTRATION NO>

080-95-92179-00

10 DATE OF REVISION OF THE TEXT

20.2.2021