

TETROXY LA

Oxytetracycline

INDICATIONS

Indicated for cattle, sheep and pigs.
Tetroxy La is licenced for the treatment and control of disease processes where oxy tetracycline susceptible bacteria are involved.

Tetroxy LA is a Long Acting Injection, one injection giving three days antibiotic cover.

If necessary the long acting treatment should be repeated after 72 hours.

BENEFITS

- Convenience, one treatment usually sufficient, reduces labour and number of injection sites
- Broad spectrum activity of oxytetracycline
- Minimal irritancy-well tolerated in cattle, sheep and pigs
- Ease of administration- 1ml/10kg
- Rapid initial blood levels followed by a long acting effect
- Ready to use, no mixing or refrigeration
- High syringability, injects well even at cold temperatures



PACKAGING

LIST NO.	UNIT PACKAGE	CASE SIZE
1TET017	100ml	12

See reverse side for Administration and Dosage.



TETROXY LA

Oxytetracycline hydrochloride

PRESENTATION

A clear yellow to amber aqueous solution for injection. Each ml contains 200mg Oxytetracycline (as oxytetracycline dihydrate).

TARGET SPECIES

Cattle, Sheep and Pigs.

INDICATIONS FOR USE

Tetroxy L.A. is indicated in the treatment and control of diseases caused by or associated with organisms sensitive to Oxytetracycline in cattle, sheep and pigs.

CONTRAINDICATIONS

Not recommended for cats, dogs, horses and donkeys.

SPECIAL WARNINGS FOR EACH TARGET SPECIES

Prolonged use of anti-infectives may result in super infection by non-susceptible organisms. Photodermatitis may occasionally occur after treatment under strong exposure to sunlight.

SPECIAL PRECAUTIONS FOR USE

- (i) Special precautions for use in animals
Following withdrawal of the first dose, use the product within 28 days.
Not recommended in cases of renal impairment (see also 4.4 above). Discard unused material.
- (ii) Special precautions to be taken by the person administering the veterinary medicinal product to animals.
Take care to avoid accidental self injection. In case of contact with eyes or skin, wash immediately with plenty of water as irritation may occur. Wash hands after use.

ADVERSE REACTIONS (FREQUENCY AND SERIOUSNESS)

Occasional local reaction of a transient nature may occur at the site of injection.

USE DURING PREGNANCY, LACTATION OR LAY

The use of Tetroxy L.A. during the period of tooth development including late pregnancy may lead to discoloration.

INTERACTION WITH OTHER MEDICINAL PRODUCTS AND OTHER FORMS OF INTERACTIONS

Do not dilute. It is unwise to administer bacteriostatic and bactericidal antibiotics concurrently.

AMOUNTS TO BE ADMINISTERED AND ADMINISTRATION ROUTE

Tetroxy L.A. is administered by deep intramuscular injection at the rate of 1 ml per 10kg bodyweight which is equivalent to 20mg Oxytetracycline per kg.
It is recommended that the following amounts of Tetroxy L.A. at one site should not be exceeded:
Cattle, Sheep and Pigs - 10 ml
Pigs under 10 kg - maximum dose of 1 ml
Effective blood levels are maintained for up to 72 hours
in cattle and 48 hours in pigs and sheep.
Because of the sustained blood levels attained at the above dosage rates with Tetroxy L.A., this is a single dose treatment.

OVERDOSE

Not applicable.

WITHDRAWAL PERIODS

Cattle (meat & offal): 39 days
Pigs (meat & offal): 40 days
Sheep (meat & offal): 28 days
Do not use in animals producing milk for human consumption.

INCOMPATIBILITIES (MAJOR)

Tetroxy L.A. should not be brought into contact with calcium solutions. Do not dilute.

SHELF LIFE

Shelf-life of the veterinary medicinal product as packaged for sale: 2 years.
Shelf-life after first opening the immediate packaging:
28 days.
Discard unused material.

SPECIAL PRECAUTIONS FOR STORAGE

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

SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED VETERINARY MEDICINAL PRODUCT OR WASTE MATERIALS DERIVED FROM USE OF SUCH PRODUCTS, IF APPROPRIATE

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal products should be disposed of in accordance with local requirements.

MARKETING AUTHORISATION HOLDER

Cross Vetpharm Group Limited, Broomhill Road, Tallaght, Dublin 24, Ireland

MARKETING AUTHORISATION NUMBER

VPA 10126/010/001

LEGAL CATEGORY

POM-V

PACKAGE QUANTITIES

100ML MULTIDOSE VIALS

A full product SPC is available on request from Bimeda or alternatively can be found on the VMD website

