LIDOCAINE 2% - lidocaine injection
MWI/VetOne

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Lidocaine 2% Injection Local Anesthetic

CAUTION: Federal law restricts this drug to use by or on the order of licensed veterinarian.

For animal use only.

Keep out of reach of children.

INDICATIONS

Lidocaine is a potent local anesthetic for producing epidural, nerve conduction and infiltration anesthesia.

CONTRAINDICATIONS

Lidocaine is contraindicated in animals with a known hypersensitivity to the drug.

PRECAUTIONS

Lidocaine is usually well tolerated. Nevertheless, as with all local anesthetics, untoward effects may occur due to hypersensitivity, faulty technique, overdosage and inadvertent intravascular or subarachnoid injection. In case of respiratory arrest, immediate resuscitation with oxygen is indicated.

CONTAINS

Each mL of sterile aqueous solution contains:
Lidocaine Hydrochloride 2.0%
Propylene Glycol 5.2%
Sodium Chloride 0.5%
Sodium Lactate 0.5%

with Methylparaben 0.15%, Sodium Metabisulfite 0.10%,
Propylparaben 0.03%, Disodium Edetate 0.001% as preservatives.

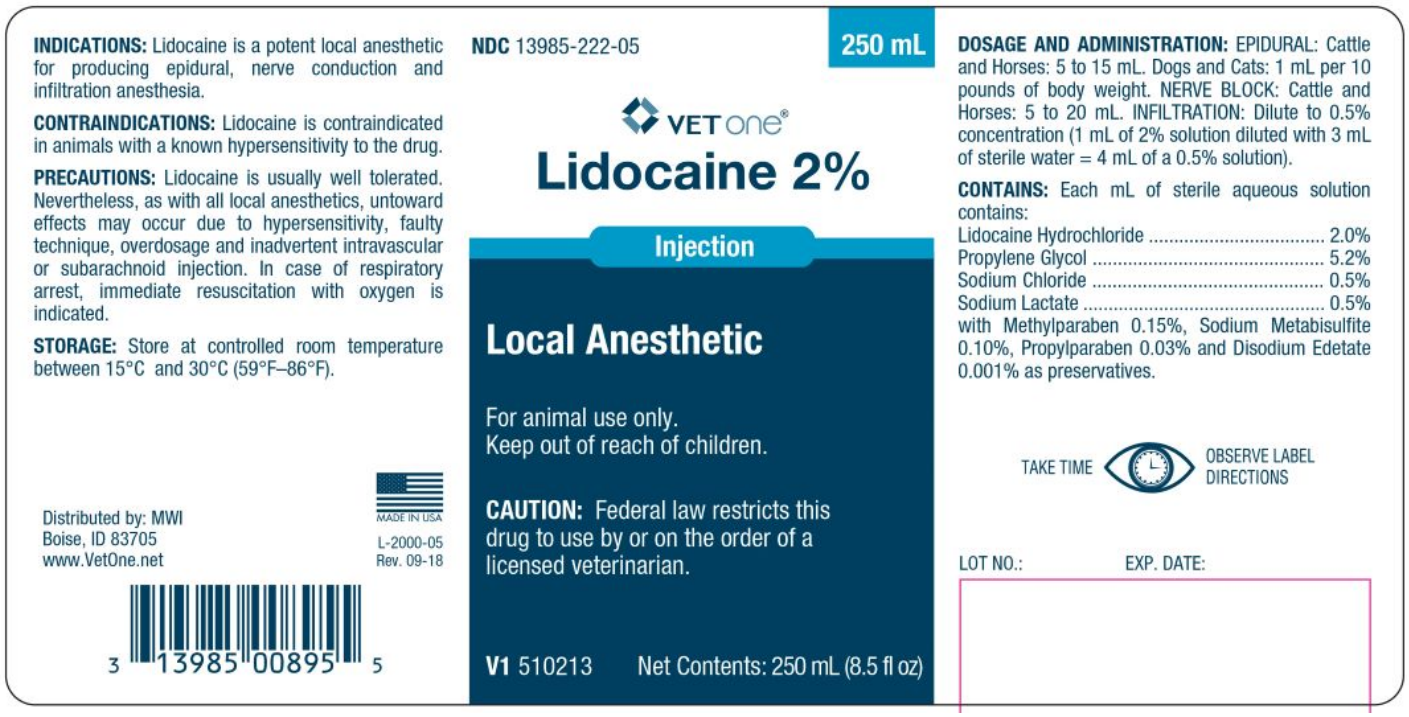
DOSAGE AND ADMINISTRATION

EPIDURAL: Cattle and Horses: 5 to 15 mL. Dogs and Cats: 1 mL per 10 pounds of body weight.
NERVE BLOCK: Cattle and Horses: 5 to 20 mL
INFILTRATION: Dilute to 0.5% concentration (1 mL of 2% solution diluted with 3 mL of sterile water = 4 mL of a 0.5% solution).

STORAGE

Store at controlled room temperature between 15º and 30ºC (59º-86ºF)

TAKE TIME OBSERVE LABEL DIRECTIONS



LIDOCAINE 2%

lidocaine injection

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:13985-222
Route of Administration	EPIDURAL, INFILTRATION		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.02 g in 1 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:13985-222-04	100 mL in 1 BOTTLE		
2	NDC:13985-222-05	250 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/02/2011	

Labeler - MWI/VetOne (019926120)