TIGER DELAYSPRAY- lidocaine spray
Magverz INC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

British Bull Dog

Lidocaine HCI 10 % ... External Anesthetic

Purpose

External Anesthetic

Keep out of Reach of Children

WARNINGS: Keep out of reach of children.

For External use only

Indication and Usage

temporarily reduces sensitivity of the penis, which helps to delay ejaculation in cases of over-rapid or premature ejaculation (coming to a climax too quickly)

Inactive Ingredients

Ethanol,Propyelene glycol, Vitamin E and Essence

Warning

Allergy alert: do not use this product if you or your partner are allergic (sensitive) to local Anesthetics.

Do not use - on broken or inflamed skin -if your partner is pregnant

Ask a doctor before use if you have, or ever had, liver or kidney problems

Ask a doctor or pharmacist before use if you are already taking prescribed drugs

when using this product

- do not get into eyes

- do not inhale

- do not exceed 24 sprays in 24 hours

Dosage and Administration

-apply 3 or more sprays, not to exceed 10, to the head and shaft of the penis before intercourse, or as directed by a doctor

-wash product after intercourse

-correct quantity and time of application will be determined by individual requirements and you should always use the minimum effective quantity

Product Label

TIGER DELAYSPRAY

lidocaine spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59240-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	10 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59240-006-06	6 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/03/2017	
2	NDC:59240-006-08	8 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/03/2017	
3	NDC:59240-006-12	12 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/03/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part346	01/03/2017	

Labeler - Magverz INC (078712269)

Registrant - Magverz INC (078712269)

Revised: 1/2017 Magverz INC