CALEXOTICS FUCK SAUCE ANAL- numbing lubricant gel Private Label Productions LLC

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.

Blackwork Numbing Lubricant 236 mL

Active Ingredient(s)

Lidocaine HCL 3.5%

Purpose

External Analgesic

Use

For temporary relief of pain, soreness or burning in the perianal area.

Warnings

■ For external use only. Do not use this product with a mechanical device.
■ Certain persons can develop allergic reactions to ingredients in this product. If the symptom being treated does not subside or if redness, irritation, swelling, pain or other symptoms develop or increase, discontinue use and consult a doctor.
■ Do not exceed the daily recommended dosage.
■ In case of any bleeding, discontinue use and consult a doctor.
■ Keep this and all drugs out of reach of children. In case of accidential overdose, seek medical help or call a Poison Control Center right away.
■ As with any drug, consult a health care professional if you are pregnant or nursing. Ask a health care professional before using this product.

Directions

 \blacksquare Apply to any area of the body where you desire extra moisture. Secure cap. \blacksquare Apply externally to the desired area up to 6 times daily.

Keep out of reach of children.

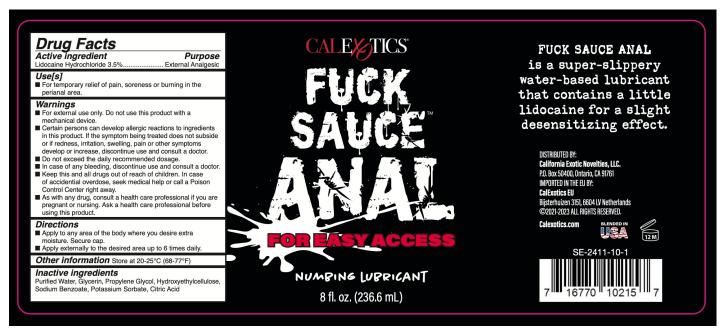
Other information

Store at 20-25°C (equals 68-77°F)

Inactive ingredients

Purified Water, Glycerin, Propylene Glycol, Hydroxyethylcellulose, Sodium Benzoate,

Package Label - Principal Display Panel



236 mL: NDC: 77632-125-01

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numbing lubricant gel

Product Information	Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77632-125			
Route of Administration	TOPICAL					

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	3.5 mg in 100 mL			

Inactive Ingredients		
Strength		

l	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:77632-125-	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/31/2023		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part348	05/31/2023			

Labeler - Private Label Productions LLC (046278265)

Revised: 7/2023 Private Label Productions LLC