FOOT HANGOVER- lidocaine hcl liquid Ari Brands, 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.

FOOT HANGOVER

Drug Facts

Active ingredients (in each bottle)

Lidocaine HCL 4%

Purpose

External analgesic

Uses

For the temporary relief of pain and itching due to minor burns, sunburn, minor cuts, abrasions, insect bites, and minor skin irritations.

Warnings

For external use only. Avoid contact with eyes.

Flammable: Do not use while smoking or near heat or flame.

Do not use in large quantities, particularly over raw surfaces or blistered areas, on large areas of the body or on a cut, irritated or swollen skin, or for more than one week without consulting a physician.

Stop use and ask a doctor if: conditions worsen, symptoms persist for more than 7 days or clear up and occur again within a few days.

If pregnant or breast feeding, ask a health professional before use.

Keep out of reach of children and pets. If swallowed, get medical help or contact the Poison Control Center right away.

Directions	
adults and children 2 years and older	apply externally to the affected area up to 3 to 4 times a day
children under 2 years	ask a doctor

Other Information

Contents under pressure. Do not store at temperatures above 120° F

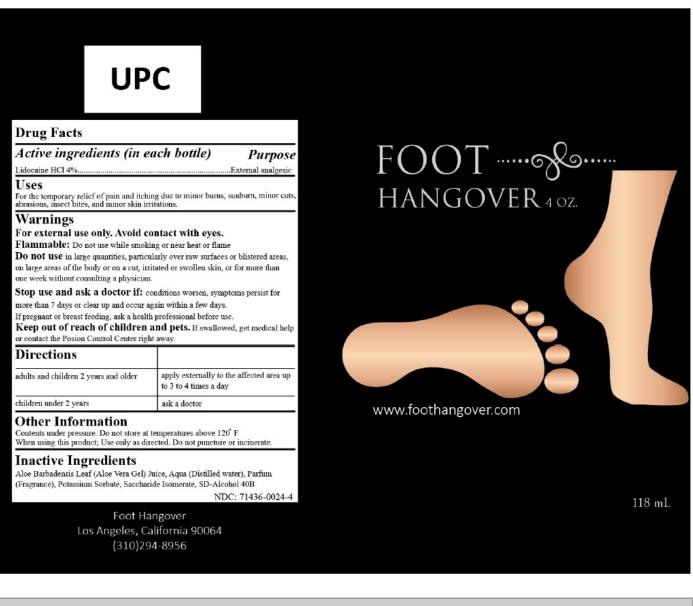
When using this product; Use only as directed. Do not puncture or incinerate.

Inactive Ingredients

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Distilled water), Parfum (Fragrance), Potassium Sorbate, Saccharide Isomerate, SD-Alcohol 40B

Foot Hangover Los Angeles, California 90064 (310)294-8956

Packaging



lidocaine hcl liquid Product Information Product Type HUMAN OTC DRUG Route of Administration TOPICAL Active Ingredient/Active Moisson Ingredient Name Basis of Strength	FOOT HANGOVER					
Product TypeHUMAN OTC DRUGItem Code (Source)NDC:71436-0024Route of AdministrationTOPICAL	lidocaine hcl liquid					
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Active Ingredient/Active Moiety	Product Type	HUMAN OTC DRUG	Ite m C	Code (Source)	NDC:71436-0	024
	Route of Administration	TOPICAL				
Ingredient Name Basis of Strength Strength	Active Ingredient/Active Moiety					
	Ingredient Name		Basis of Strength		Strength	

Ina	active Ingredie	nts				
Ingredient Name					Strength	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)						
WA	TER (UNII: 059QF0	KO0R)				
PO	TASSIUM SORBAT	FE (UNII: 1VP	U26JZZ4)			
SAC	CCHARIDE ISOME	RATE (UNII:	W8K377W98I)			
AL	C OHOL (UNII: 3K9	958V90M)				
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Labeler -	Ari Brands, Llc (080658382)
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Revised: 1/2018

Ari Brands, Llc