YMLABS- aloe hand sanitizer gel Yusef Manufacturing Laboratories

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.

YMlabs Gel Hand Sanitizer Aloe

DIRECTIONS: Put small amount into palm of hand. Rub together until dry.

WARNINGS: For External Use Only. FLAMMABLE - Do not use near heat, fire, flame or while smoking. *Do not store where temperature exceeds 105°F. Do not get in eyes. In case of eye contact flush with water. *Do not Inhale or Ingest. If swallowed, get medical help or contact poison control immediately. *Stop use and ask a doctor if skin irritation develops. *KEEP OUT OF REACH OF CHILDREN.

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Drug Facts

Active ingredients Purpose

Ethyl Alcohol (62%)......Sanitizer

USES: Decreases Bacteria on the hands.

INACTIVE INGREDIENTS: Water, Carbomer, Aloe Barbadensis Leaf Gel, Treithanolamine.

Made In USA
Distributed by
<Vendor, City, ST>
Questions? 877.827.5425
NEVER Tested On Animals.

Drug FactsActive ingredients Purpose Ethyl Alcohol (62%)......Sanitizer

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YMLABS

aloe hand sanitizer gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10827-1022	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 g in 100 g		

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
CARBOMER HOMOPOLYMER TYPE A (UNII: F68VH75CJC)			
TROLAMINE (UNII: 903K93S3TK)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10827- 1022-1	30 g in 1 BOTTLE; Type 0: Not a Combination Product	12/30/2021	
2	NDC:10827- 1022-2	56 g in 1 BOTTLE; Type 0: Not a Combination Product	12/30/2021	

-	NDC:10827- 1022-3	59 g in 1 BOTTLE; Type 0: Not a Combination Product	12/30/2021	
4	NDC:10827- 1022-4	118 g in 1 BOTTLE; Type 0: Not a Combination Product	12/30/2021	
5	NDC:10827- 1022-5	228 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/30/2021	

Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
part333E	12/30/2021		
	Application Number or Monograph Citation	Application Number or Monograph Marketing Start Citation Date	

Labeler - Yusef Manufacturing Laboratories (144150674)

Registrant - Yusef Manufacturing Laboratories (144150674)

Establishment				
Name	Address	ID/FEI	Business Operations	
Yusef Manufacturing Laboratories		144150674	manufacture(10827-1022)	

Revised: 2/2022 Yusef Manufacturing Laboratories