CAREONE LEMON MERINGUE HAND SANITIZER- ethyl alcohol liquid American Sales Company

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.

Drug Facts

Active ingredient

Ethyl Alcohol 65%

Purpose

Antiseptic

Uses

• helps reduce bacteria on the skin

Warnings

For external use only

• flammable, keep away from fire or flame

When using this product

• avoid contact with eyes. If contact occurs, rinse with water.

Stop using this product and ask doctor if

• irritation or redness develops and lasts.

Keep out of reach of children.

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

• wet hands thoroughly and rub together until dry

Other information

store at a temperature below 110°F (43°C)

Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Propylene Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Carbomer, Aminomethyl Propanol, Fragrance (Parfum), Yellow 5 (CI 19140), Yellow 10 (CI 47005).

Label Copy





CAREONE LEMON MERINGUE HAND SANITIZER									
ethyl alcohol liquid									
Product Information									
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:41520-056					
Route of Administration	TOPICAL								
Active Ingredient/Active Moi	ety								
Ingredient Name			Basis of Strength		Strength				
ALCOHOL (UNII: 3K9958V90M) (ALC		ALCOHOL		650 mg in 1 mL					
Inactive Ingredients									
	Ingredient Name				Strength				
WATER (UNII: 059QF0KO0R)									
ISOPROPYL ALCOHOL (UNII: ND2M	416302)								
PROPYLENE GLYCOL (UNII: 6DC9Q	167V3)								
GLYCERIN (UNII: PDC6A3C0OX)									
ISOPROPYL MYRISTATE (UNII: 0 RE8									
ALPHATO COPHEROL ACETATE, I									
CARBOMER 934 (UNII: Z135WT9208)									

AMINO METHYLPROPANOL (UNII: LU49E6626Q)

гг	Nec VELLOW NO						
	0&C YELLOW NO		,				
D	&C YELLOW NO.	10 (UI	NII: 358 W5USQ3G)				
Packaging							
#	Item Code		Package Description	Marketing Start Date	Marketing End Date		
	NDC:41520-056- 08	236 m Produ	L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination ct	09/25/2017			
2	NDC:41520-056- 02	59 mI Produ	in 1 BOTTLE, PLASTIC; Type 0: Not a Combination ct	09/25/2017			
N	Iarketing In	forn	nation				
Marketing Category		ory	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
-	OTC monograph not final						

Labeler - American Sales Company (809183973)

Registrant - Apollo Health and Beauty Care Inc. (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(41520-056)

Revised: 9/2017

American Sales Company