HAND SANITIZER- ethyl alcohol gel
Allure Labs 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.

DRUG FACTS

Active Ingredients:

Ethyl Alcohol - 65%

Purpose: Antimicrobial

Uses: To decrease the risk of skin infection in minor cuts, scrapes and burns.

Warning: For External use only - Hands

Flammable: Keep away from heat and Flame

When using this products:

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water.

Stop use and ask a doctor if skin irritation develops.

Keep out of reach of children:

- If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions:

- Wet hands throughly with product and allow to dry without wiping. For children under 6, use only under adult supervision. Not recommended for infants.

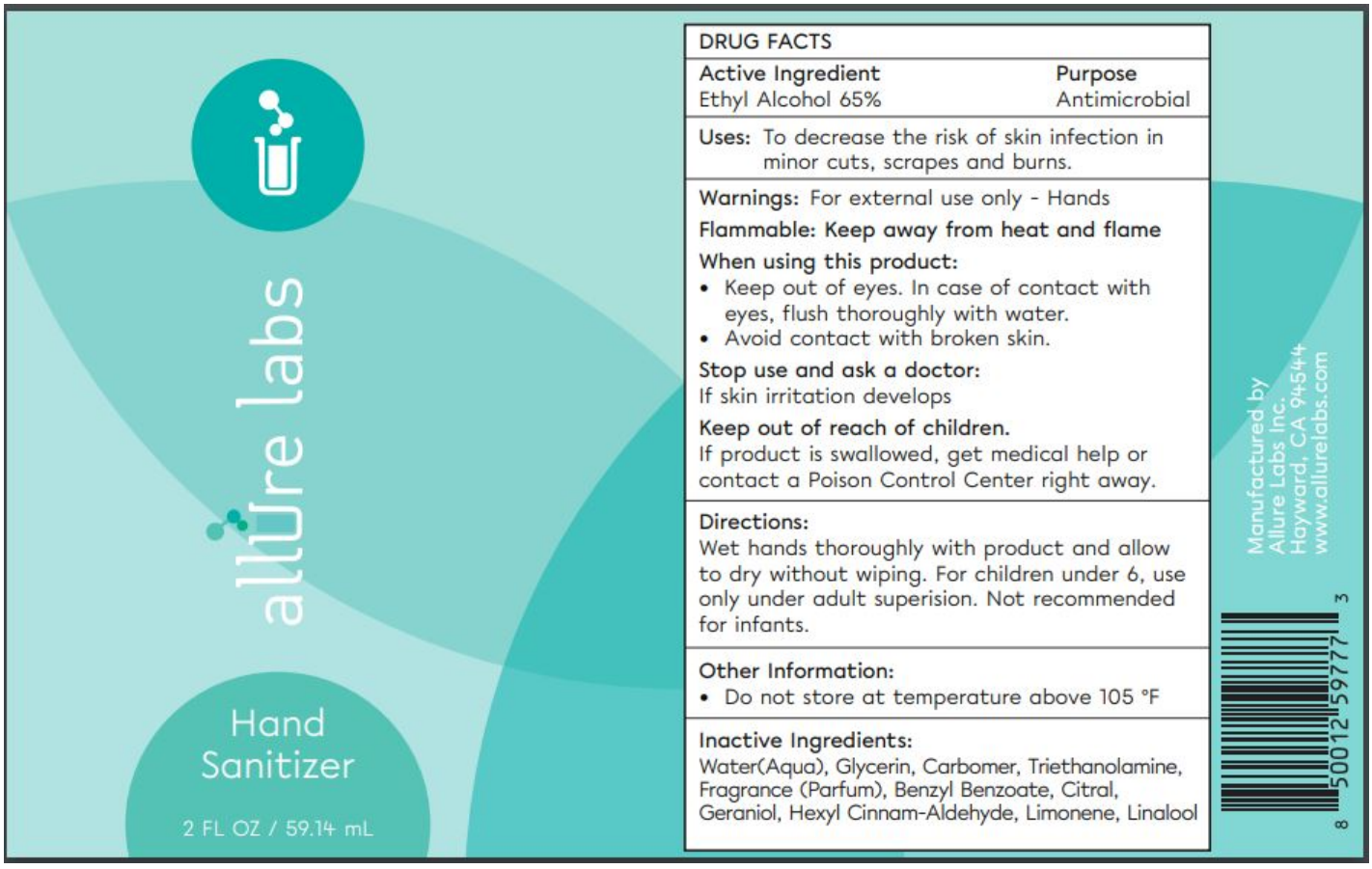
Inactive Ingredients: Water (Aqua), Glycerin, Carbomer, Triethanolamine, Fragrance (Parfume), Benzyl Benzoate, Citral, Geraniol, Hexyl Cinnam Aldehyde, Limonene, Lonalool.

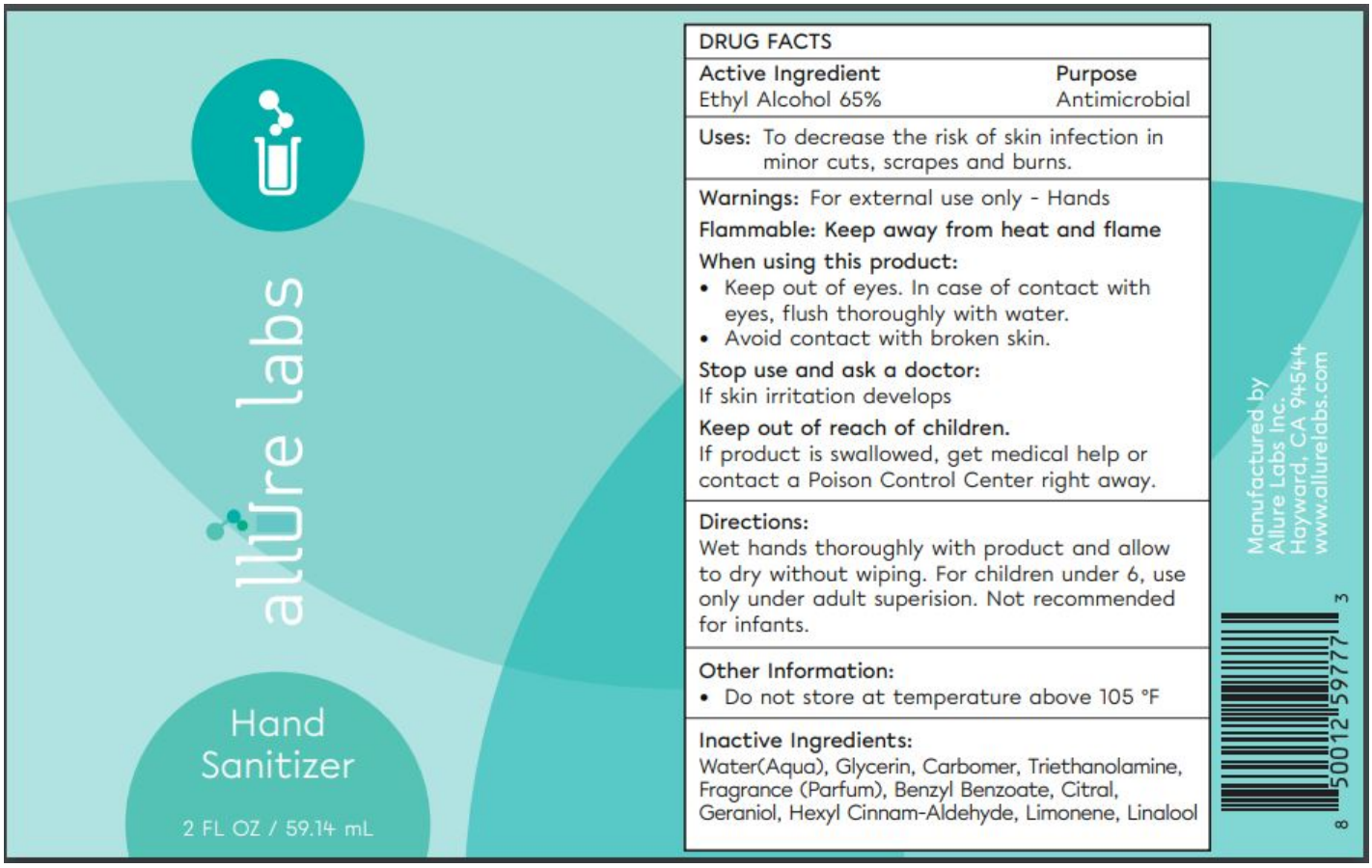
Manufactured By

Allure Labs, Inc.

Hayward, CA 94544

www.allurelabs.com





HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62742-4184
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	650 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
TROLAMINE (UNII: 9O3K93S3TK)	
BENZYL BENZOATE (UNII: N863NB338G)	
CITRAL (UNII: T7EU0O9VPP)	
GERANIOL (UNII: L837108USY)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62742-4184-1	59.14 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/17/2020	
2	NDC:62742-4184-2	355 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/17/2020	

Labeler - Allure Labs Inc (926831603)

Revised: 3/2020 Allure Labs Inc