DR.DAY S HAND CLEAN GEL- alcohol gel Labis co Corp.

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.

Labisco Corp. - Dr.Day S Hand Clean Gel

Alcohol

water, carbomer, triethanolamine, glycerin, propylene glycol, aloe vera leaf, green tea leaf, jojoba oil

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

keep out of reach of the children

Place enough product on hands to cover all surfaces. Rub hands together until dry.

Supervise children under 6 years of age when using this product to avoid swallowing.

For external use only. Flammable. Keep away from heat or flame.

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

for external use only



DR.DAY S HAND CLEAN GEL

alcohol gel

Product Information

Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:75535-0003
	Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	350 mL in 500 mL		

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
CARBO MER HO MO PO LYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
GREEN TEA LEAF (UNII: W2ZU1RY8B0)				
JOJOBA OIL (UNII: 724GKU717M)				
TROLAMINE (UNII: 9O3K93S3TK)				
GLYCERIN (UNII: PDC6A3C0OX)				

ı	Packaging					
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
ı	1	NDC:75535-0003-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/04/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	04/10/2020		

Labeler - Labisco Corp. (695712140)

Registrant - Labisco Corp. (695712140)

Establishment				
Name	Address	ID/FEI	Business Operations	
Labisco Corp.		695712140	manufacture(75535-0003), label(75535-0003), pack(75535-0003)	

Revised: 5/2020 Labisco Corp.