ANTACGEL- hand sanitizer gel with peppermint and lavender, 73% alcohol gel MEDIPHARCO PHARMACEUTICAL JOINT STOCK 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.

Alcohol 73%

Antiseptic

Hand sanitizer gel helps to sanitize hands, eliminate unpleasant odor and leave hands feeling soft and fresh.

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

* in children less than 2 months of age

* on open skin wounds

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

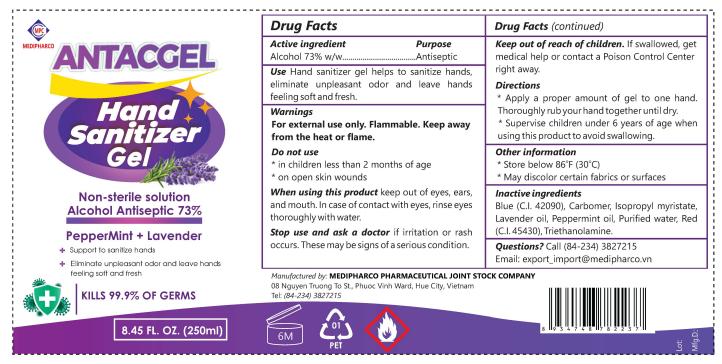
if irritation or rash occurs. These may be signs of a serious condition.

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

- * Apply a proper amount of gel to one hand. Thoroughly rub your hand together until dry.
- * Supervise children under 6 years of age when using this product to avoid swallowing.
- * Store below 86°F (30°C)
- * May discolor certain fabrics or surfaces

Blue (C.I. 42090), Carbomer, Isopropyl myristate, Lavender oil, Peppermint oil, Purified water, Red (C.I. 45430), Triethanolamine.

Apply a proper amount of gel to one hand.





ANTACGEL

hand sanitizer gel with peppermint and lavender, 73% alcohol gel

Product Type	HUMAN OTC DRUG	Item Code (Source) NDC		NDC:79	2:79633-202	
Route of Administration	TOPICAL					
Active Ingredient/Active Mo	iety					
Ingr	edient Name		Basis of Stren	gth	Strength	
ALCOHOL (UNII: 3K9958V90M) (AL	COHOL - UNII:3K9958V90M)		ALCOHOL		73 g in 100 g	
Inactive Ingredients						
	Ingredient Name				Strength	
	Ingreatent Name					
LAVENDER OIL (UNII: ZBP1YXW0H8	•					
LAVENDER OIL (UNII: ZBP1YXW0H8 TROLAMINE (UNII: 903K93S3TK)	•					
LAVENDER OIL (UNII: ZBP1YXW0H8 TROLAMINE (UNII: 903K93S3TK) ERYTHROSINE SODIUM ANHYDRO)					

IS								
C								
P								
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)								
Packaging								
#	Item Code		Package Description	Marketing Start Date	Marketing End Date			
1	NDC:79633-202-25	217	g in 1 BOTTLE; Type 0: Not a Combination Product	07/22/2020				
2	NDC:79633-202-45	434	g in 1 BOTTLE; Type 0: Not a Combination Product	07/22/2020				
Marketing Information								
	Marketing Catego	ry	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
0	TC monograph not fin	nal	part333A	07/22/2020				

Labeler - MEDIPHARCO PHARMACEUTICAL JOINT STOCK COMPANY (555273697)

Establishment

Name	Address	ID/FEI	Business Operations
MEDIPHARCO PHARMACEUTICAL JOINT STOCK COMPANY		555273697	manufacture(79633-202)
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Revised: 7/2020

MEDIPHARCO PHARMACEUTICAL JOINT STOCK COMPANY