DR. DADDYS HAND SANITIZER- alcohol gel TB Healthcare Co., Ltd.

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

Alcohol

Water, Glycerin, Triethanolamine, Carbomer, Green Tea Extract, Aloe Extract, Sage Extract, Matricaria(Chamomilla) Extract

Antimicrobial

KEEP OUT OF REACH OF THE CHILDREN

- Store between 1-30C
- Avoid freezing and excessive heat above 40C

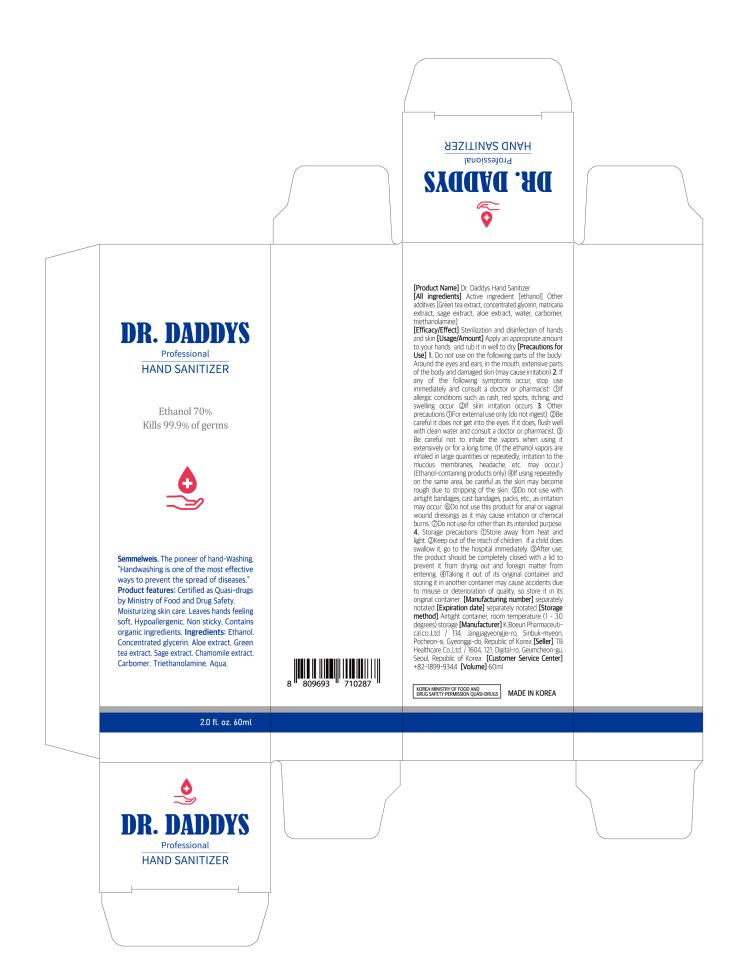
For external use only. Flammable, Keep away from fire or flames. Do not use

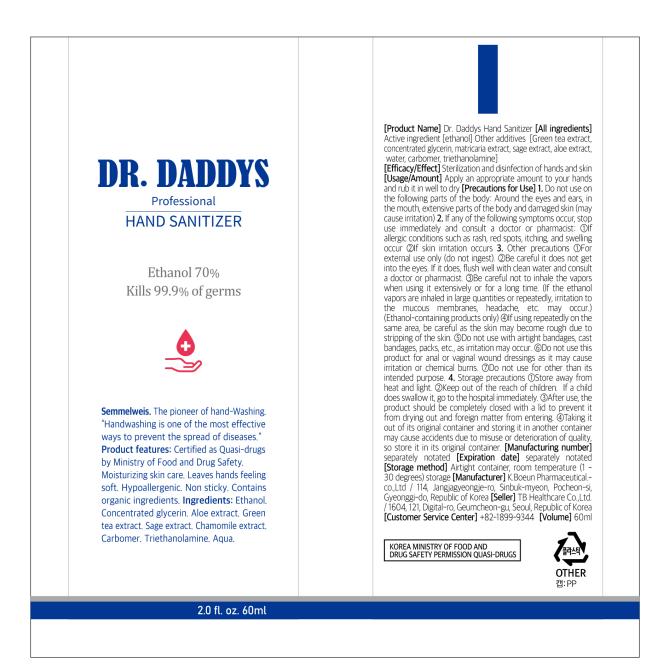
- in children less than 2 months of age
- on open skin wounds

when using this product keep out of eyes, ears, and mouth.

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.

- 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.





DR. DADDYS HAND SANITIZER

alcohol gel

 Product Information

 Product Type
 HUMAN OTC DRUG
 NDC:5884-0002

 Route of Administration
 TOPICAL
 V

 Active Ingredient/Active Mo:sure
 V
 V

 Ingretient Name
 Basis of Strength
 Strength

ALCOHOL	(UNII: 3K9958V90M)	(ALCOHOL - UNII:3K9958V90M)	

ALCOHOL

Inactive Ingredient	S				
Ingredient Name					
CARBOMER HOMOPOL	YMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
GLYCERIN (UNII: PDC6A	3C0OX)				
ALOE (UNII: V5VD430 YV	v9)				
TROLAMINE (UNII: 903)	K93S3TK)				
WATER (UNII: 059QF0KC	DOR)				
CHAMO MILE (UNII: FGL	3685T2X)				
GREEN TEA LEAF (UNII:	W2ZU1RY8B0)				
Packaging					
Packaging # Item Code	Package Description	Marketing Start Date	Marketing End Date		
# Item Code	Package Description 0 mL in 1 TUBE; Type 0: Not a Combination Product	Marketing Start Date	Marketing End Date		
# Item Code	.	_	Marketing End Date		
# Item Code	0 mL in 1 TUBE; Type 0: Not a Combination Product	_	Marketing End Date		
# Item Code 1 NDC:76884-0002-1 6	0 mL in 1 TUBE; Type 0: Not a Combination Product	_	Marketing End Date Marketing End Date		

Labeler - TB Healthcare Co., Ltd. (695035143)

Registrant - TB Healthcare Co., Ltd. (695035143)

Establishment

Name	Address	ID/FEI	Business Operations
TB Healthcare Co., Ltd.		695035143	label(76884-0002)

Establishment

Name	Address	ID/FEI	Business Operations
K.Boeun Pharmaceutical Co.,Ltd.		695674074	manufacture(76884-0002)

Revised: 5/2020

TB Healthcare Co., Ltd.