HAND SANITIZER- alcohol gel Brands International

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.

Germs Be Gone Hand sanitizers

Drug Facts

Active Ingredient

Ethyl Alcohol, 66% w/w (73%, v/v)

Purpose

Antiseptic

Uses

instant healthcare personnel hand antiseptic to reduce bacteria that potentially can cause disease

- instant hand antiseptic to decrease bacteria on the skin
- recommended for repeated use

Warnings

For external use only. Flammable, keep away from fire or flame.

When using this product keep out of eyes. If contact with eyes occurs, rinse promptly and thoroughly with water.

Stop use and ask a doctor if significant irritation or sensitization develops.

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

Directions

Apply to clean, dry hands. Apply sufficient amount to thoroughly wet all surfaces of hands and fingers. Rub onto hands until dry.

• Supervise children in the use of this product.

Other Information

Store below 43°C (110°F)

Inactive Ingredients

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, aminomethyl propanol, denatonium benzoate, disodium EDTA, glycerin, PEG-40 castor oil, propylene glycol, water, isopropyl myristate.

Please see label on the container.



Drug Facts / Informations médicament ACTIVE INGREDIENT: Ethyl alcohol 65% w/w PURPOSE: Antiseptic ACTIVE INGREDIENT: Ettyl alcohol 65% w/w PURPOSE: Antiseptic USE: to holp reduce bacteria on the skin WARNINGS: Flammable. Keep away from open flame and sources of heat. For external use only, Avoid contact with the eyes. If contact occurs, flush eyes with water. Discontinue use and consult a healthcare practitioner if imitation develops. Rarep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. DIRECTIONS: Place encough product in your paim to thoroughly cover your hands. Rub hands together brisky until dry, Children under 6 years of oge should be supervised when using this product. OTHER INFORMATION: Store below 110°F (43°C). May discolor certain tabrics or surfaces. INGRÉDIENT ACTIF: Alcool éthylique 65% w/w UTILITÉ: Antiseptique INGRÉDIENT ACTIF: Alcool éthylique 65% w/w UTILITÉ: Antiseptique USAGE: Pour aider à réduire les bactéries sur la peau. AVERTISSEMENTS: Inflammable. Tenir les produit à ficant du feu, des farmmes nues, des ampoules électriques oilumées et des autres sources de chaleur. Pour usage entenne seulement, éviter le contact avec les gruus et les muqueuses. Su une initiation apparaît, cesser d'utiliser produit et consulter un praticien de soins de samt 6, 4APDER HORS DE LA PORTÉE DES ENFANTS. En cas d'ingestion, communiquez immédiatement avec voire médicain ou le centre antipoison. MODE D'EMPLOI: Metter sufficienment de produit dans voite pauline pour MODE O'EMPLOI: Mettez suffisamment de produit dans votre paume pour bien couvrir vos mains. Frottez vivement vos mains ensemble jusqu'à ce qu'elle soient sèches. Les enfants de mains de 6 ans doivent être surveillés lorsqu'ils utilisent ce produit.

 Utilisent ce presum.

 AUTRES INFORMATIONS: À conserver en dessous de 43°C (110°F-) reux

 AUTRES INSues ou nurfaces.

 INACTIVE INGREDIENTS /INACTIFS: Water (Eau).

 bopropri Alcohol, Giyceri, cuthorre, Aninomethyl Poponol, Fragmano (Parlum). Proylene Giycol, isopropri Myrister, Alce Barbadensis Leuf Juice (Mare Vera). Toopher/ Acetato (Vasmin E).

 Lot# 20074006
 MFG : 15/03/2020
 EXP: 14/03/2023

 Manufactured by:
 NPM80021511

 Wearvarks (In UL3X S2, Canada www.brandsicorp.com
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HAND SANITIZER alcohol gel						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:50157-500		
Route of Administration	TOPICAL					
Active Ingredient/Active Moi	ety					
Ingredient Name			Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	66 mL in 100 mL		
Inactive Ingredients						
Ingredient Name					Strength	
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)						
WATER (UNII: 059QF0KO0R)						
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)						
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)						
GLYCERIN (UNII: PDC6A3C0OX)						
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)						
AMINOMETHYLPROPANOL (UNII: LU49E6626Q) PEG-40 CASTOR OIL (UNII: 4ERD2076EF)						
EDETATE DISODIUM (UNII: 7FLD91C						
DENATO NIUM BENZO ATE (UNII: 4YK						

Packaging						
# Item Code	Package Descriptio	n Marketing Start Date	Marketing End Date			
1 NDC:50157-500-01	236 mL in 1 BOTTLE; Type 0: Not a Co	mbination Product 03/19/2020				
Marketing Information						
Marketing Categ	ory Application Number or Mono	graph Citation Marketing Start Date	Marketing End Date			
OTC monograph not	inal part333E	03/19/2020				

Labeler - Brands International (243748238)

Revised: 3/2020

Brands International