RELIEF HAND SANITIZING WIPES- ethanol cloth
Daehan P&H 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.

Relief HAND SANITIZING WIPES | 1 ct. in 1 packet < 20 ct. in 1 box | 79002-162-20

Active Ingredient(s)

Ethanol 62% Purpose: Antiseptic

Purpose

Antiseptic

Uses

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

Warnings

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

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

Directions

■ Remove the wipes one by one, and use on the skin gently.
■ There is a risk of contamination if the wet wipe is put back into the packaging with unclean hands.
■ Do not throw away wipes in a flush toilet, as there is a risk of clogging it.
■ Supervise children under 6 years of age when using.
■ Do not allow infants and toddlers swallow or put in the mouth.
■ Avoid direct inhalation of vapors during application. (Headaches and irritation to mucous membranes may occur when directly inhaled.)
■ Keep product in its original container as storing the product in anything other than the original container may result in accidents or cause the integrity of the product to diminish.

Other information

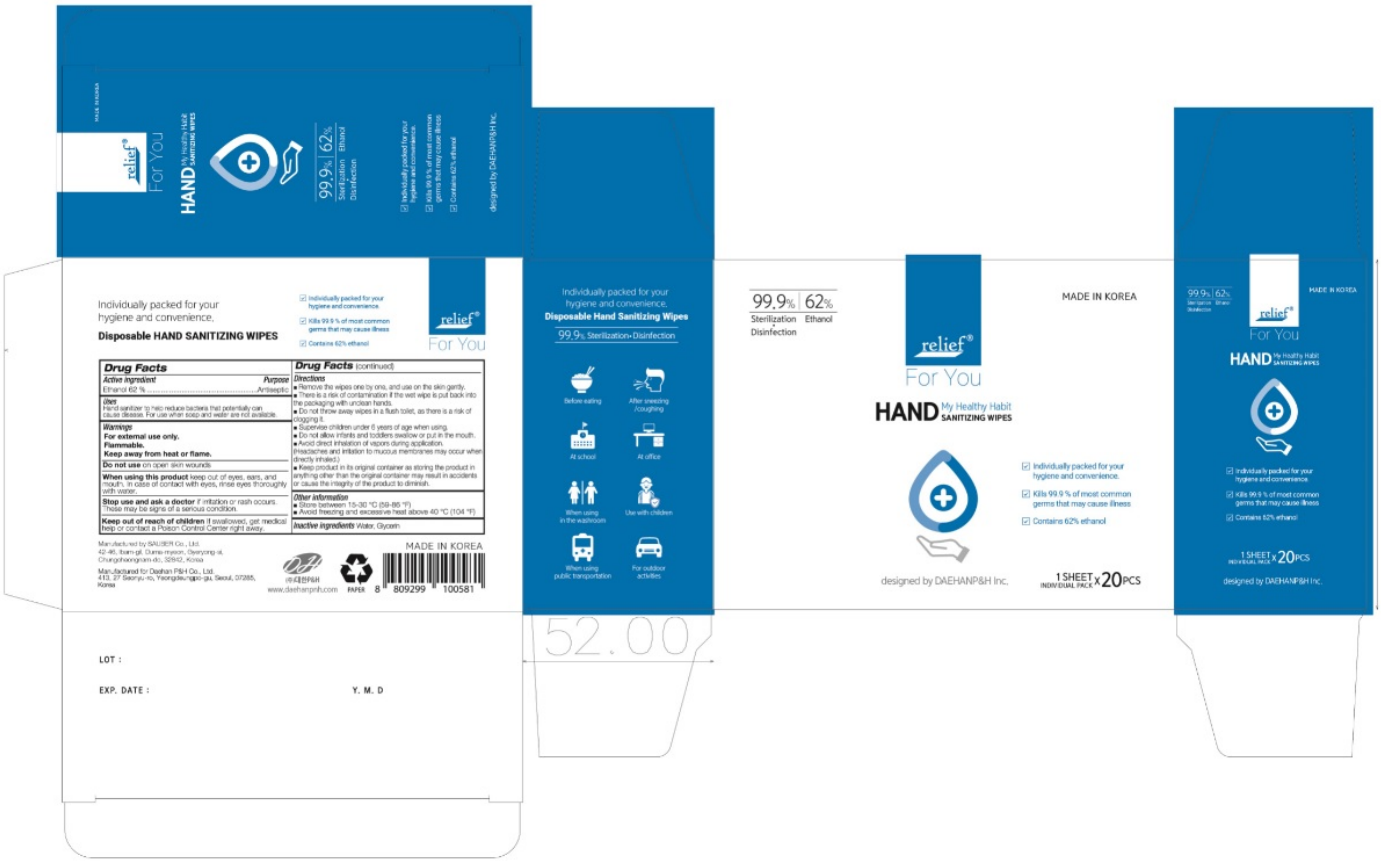
■ Store between 15-30°C (59-86°F)

■ Avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients

Water, Glycerin

Package Label - Principal Display Panel



RELIEF HAND SANITIZING WIPES

ethanol cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79002-162
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79002-162-20	20 in 1 BOX	06/22/2020	
1	NDC:79002-162-01	1 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/22/2020	

Labeler - Daehan P&H Co., Ltd. (557818402)

Establishment

Name	Address	ID/FEI	Business Operations
Sauber Co.,Ltd.		690395844	manufacture(79002-162)

Revised: 6/2020 Daehan P&H Co., Ltd.