CLEAN CARE HAND SANITIZER- alcohol gel
URBANBCL 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

ethyl alcohol

The convenient Sense Fresh Advanced Instant Hand Sanitizer with Aloe is an antimicrobial liquid that contains 70 percent ethyl alcohol to help reduce the number of germs on hands, and three moisturizers to help keep skin hydrated. No rinsing with water or drying with towels is needed. Alcohol-based hand sanitizers can quickly reduce the number of germs on hands within 30 seconds. The hand sanitizer meets U.S. Food and Drug Administration (FDA) healthcare personnel hand-washing requirements. It’s perfect for on-the-go uses when traveling outside of the home, and fits neatly in purses, backpacks or briefcases.

KEEP OUT OF REACH OF THE CHILDREN

Just spray enough product in your palm to cover hands and rub hands together briskly until dry.

Children under 6 years of age should be supervised when using Sense Fresh.

Hand sanitizer to help reduce bacteria on the skin that could cause disease. Recommended for repeated use.

■ Flammable. Keep away from fire or flame.

■ For external use only.

■ Do not use in eyes.

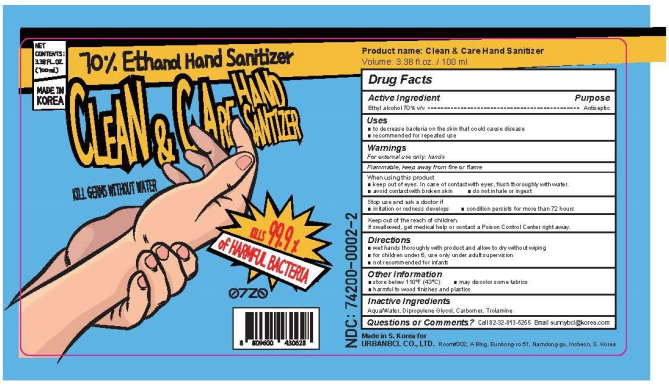
■ lf swallowed, get medical help promptly.

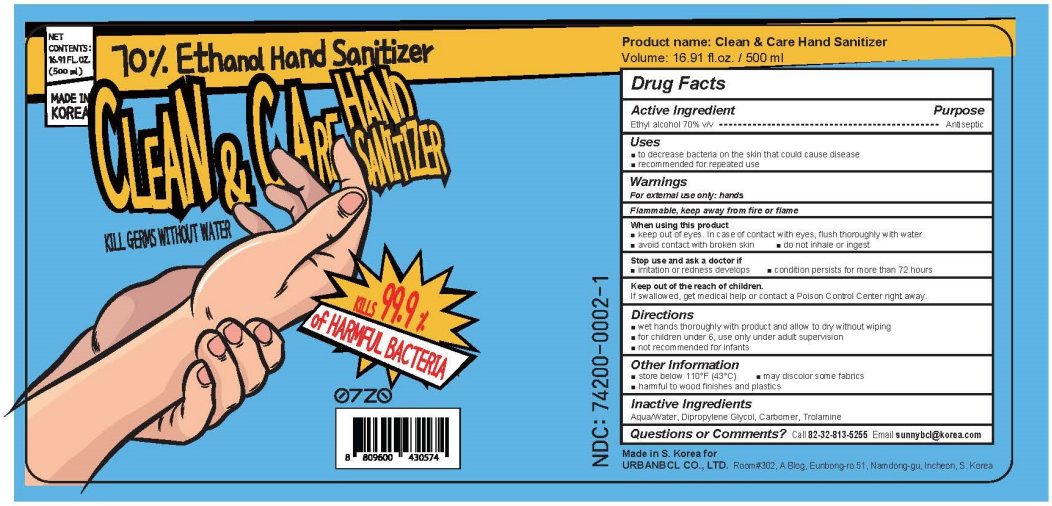
■ Stop use, ask doctor lf irritation occurs.

■ Keep out of reach of children.

for external use only

water, trolamine, dipropylene glycol, carbomer





CLEAN CARE HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74200-0012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
CARBOMER 940 (UNII: 4Q93RCW27E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74200-0012-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	
2	NDC:74200-0012-2	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

Labeler - URBANBCL CO LTD (694794065)

Registrant - URBANBCL CO LTD (694794065)

Establishment

Name	Address	ID/FEI	Business Operations
URBANBCL CO LTD		694794065	manufacture(74200-0012)

Revised: 4/2020 URBANBCL CO LTD