CLEAN CARE PLUS HAND SANITIZER- alcohol gel
AT Bio Pharm 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

ethanol

Water, Glycerin, butylene glycol, carbomer, triethanolamine, rosmarinus officinalis (rosemary) leaf extract, paeonia suffruticosa bark extract, aloe barbadensis leaf extract, calendula officianalis flower extract

Antiseptic

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

hand sanitizer to help reduce bacteria on the skin

KEEP OUT OF REACH OF THE CHILDREN

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.

Dispense appropriate amount on your palm and thoroughly spread on both hands and rub into the skin until dry

For external use only.

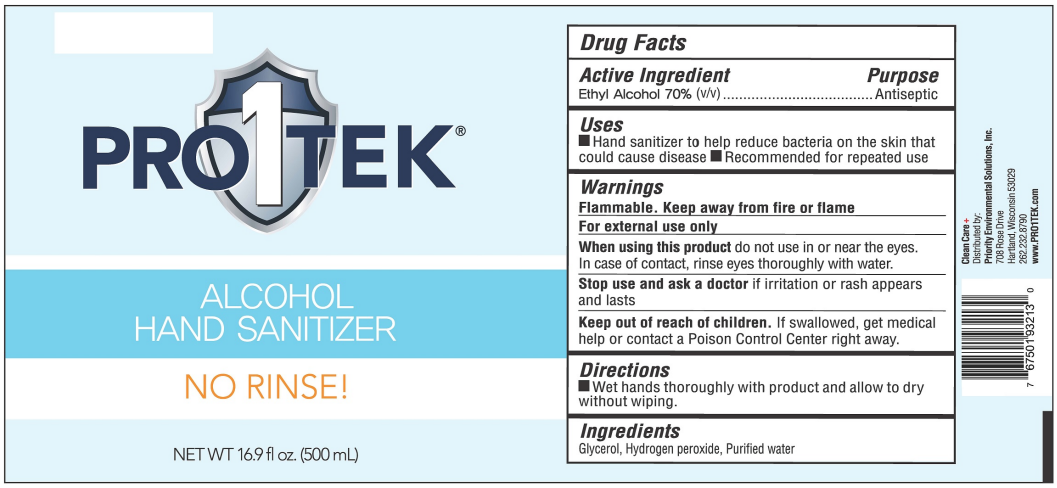
Flammable, keep away from fire or flame.

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.

for external use only



CLEAN CARE PLUS HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73932-0003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73932-0003-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/21/2020	

Labeler - AT Bio Pharm Co Ltd (695742996)

Registrant - AT Bio Pharm Co Ltd (695742996)

Establishment

Name	Address	ID/FEI	Business Operations
AT Bio Pharm Co Ltd		695742996	manufacture(73932-0003)

Revised: 4/2020 AT Bio Pharm Co Ltd