THE AEGIS PRIME S HAND SANITIZER- alcohol liquid WOWBIOTECH

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.

WOWBIOTECH - THE AEGIS PRIME S HAND SANITIZER

Alcohol

water, carbomer, triethanolamine, glycerin, butylene glycol, aloe barbadensis extract, fragrance(lemon-001s)

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

keep out of reach of the children

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.

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

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. In case of contact with eyes, rinse 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.

for external use only

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Basis of Strength	HAND SAN Easy to use without w Bacteria removal and moistu Colorless and transpar	NITIZER vater rizing effect	Warnings Fo Do not apply arc When using this In case of conta Discontinue use Keep out of reac Do not drink. In O Directions put hands. *rub into Other informati *may discolor fa Inactive ingred Glycerin, Butyler	r external use only. Flammat bund eyes. Do not use in ears & product, avoid contact with eye ct, flush eyes with water. if irritation or redness develops ho f children. Children must be case of accidental ingestion, se amp as needed into your palms skin until dry. ion store in a cool dry place bel brics. ients Ethanol, Water, Carbome ne glycol, Aloe barbadensis ext	k mouth es. If condition supervised i lek professior and thorough low 30 °C (1~3 er, Triethanola ract, Fragram	persist, consult a doctor. n use of this product. nal assistance. hly spread on both 80°C)
Accive Ingredient/Active Moie Active Ingredient Name Active Ingredient Name Act				WOWBIOTEC 30, Gajeong-ro 67beon- Incheon, Republic of	H gil, Seo-gu, f Korea	99 ->
Product Type HUMAN OTC DRUG Item Code (Source) NDC:76:007 Route of Administration TOPICAL Verter Source) Verter Source) Active Ingredient/Active Moieter Basis of Strength Strength Actore Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 350 mL in 500 mL Inactive Ingredients Strength Strength Matter (UNII: 059QF0K00R) Strength Strength Route of Sygen Strength Strength Strength WATER (UNII: 059QF0K00R) Strength Strength GROLAMINE (UNII: 903K93S3TK) Strength Strength CARBOMER HOMOPOLYMER, UNSECIFIED TYPE (UNII: 0.45MM3/FC) Strength GLYCERIN (UNII: PDC6A3C00X) Strength						
Route of Administration TOPICAL Active Ingredient/Active Moietter Active Ingredient/Active Moietter Basis of Strength AlcoHol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL (UNII: 903K93S3TK) Strenge ALCOHOL (UNII: 903K93S3TK) CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC) GIYCERIN (UNII: PDC6A3C00X)	lcohol liquid	HAND SANITIZER				
Ingredient NameBasis of StrengthStrengthALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)ALCOHOL350 mL in 500 mlInactive IngredientsStrengthStrengthIngredientsFARER (UNII: 059QF0K00R)TROLAMINE (UNII: 903K93S3TK)CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)GLYCERIN (UNII: PDC6A3C00X)	lcohol liquid Product Information		Item Code ((Source)	NDC:76	6627-0007
Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 350 mL in 500 ml Inactive Ingredients Strength Inactive Ingredients Strength Ingredient Name Strength Ingredint Name Strength <th>lcohol liquid Product Information Product Type</th> <th>HUMAN OTC DRUG</th> <th>Item Code (</th> <th>(Source)</th> <th>NDC:76</th> <th>6627-0007</th>	lcohol liquid Product Information Product Type	HUMAN OTC DRUG	Item Code ((Source)	NDC:76	6627-0007
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 350 mL in 500 ml Inactive Ingredients Ingredient Name Streng WATER (UNII: 059QF0K00R) TROLAMINE (UNII: 903K93S3TK) CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC) GLYCERIN (UNII: PDC6A3C00X) (100 ml)	lcohol liquid Product Information Product Type Route of Administration	HUMAN OTC DRUG TOPICAL	Item Code ((Source)	NDC:76	6627-0007
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Ingredient NameStrengWATER (UNII: 059QF0K00R)TROLAMINE (UNII: 903K93S3TK)CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)GLYCERIN (UNII: PDC6A3C00X)	Alcohol liquid Product Information Product Type Route of Administration Active Ingredient/Active Mo Ingre	HUMAN OTC DRUG TOPICAL		Basis of Strengt	h	Strength
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TROLAMINE (UNII: 903K93S3TK) CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC) GLYCERIN (UNII: PDC6A3C00X)	lcohol liquid Product Information Product Type Route of Administration Active Ingredient/Active Me Ingre ALCOHOL (UNII: 3K9958V90M) (Al	HUMAN OTC DRUG TOPICAL		Basis of Strengt	h	Strength) mL in 500 mL
GLYCERIN (UNII: PDC6A3C0OX)	lcohol liquid Product Information Product Type Route of Administration Active Ingredient/Active Mo Ingre ALCOHOL (UNII: 3K9958V90M) (Al	HUMAN OTC DRUG TOPICAL		Basis of Strengt	h	Strength
	Icohol liquid Product Information Product Type Route of Administration Active Ingredient/Active Mo Ingre ALCOHOL (UNII: 3K9958V90M) (AI Inactive Ingredients WATER (UNII: 059QF0K00R)	HUMAN OTC DRUG TOPICAL		Basis of Strengt	h	Strength) mL in 500 mL
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	lcohol liquid Product Information Product Type Route of Administration Active Ingredient/Active Me Ingre ALCOHOL (UNII: 3K9958V90M) (Al Inactive Ingredients WATER (UNII: 059QF0K00R) TROLAMINE (UNII: 903K93S3TK)	HUMAN OTC DRUG TOPICAL Die ty edient Name LCOHOL - UNII:3K9958V90M)	F	Basis of Strengt	h	Strength) mL in 500 mL
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Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date					
1 NDC:76627- 0007-1	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/11/2020						
Marketing Information								
Marketing Categ	ory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date					
OTC monograph not	inal part333A	05/11/2020						

Labeler - WOWBIOTECH (695625834)

Registrant - WOWBIOTECH (695625834)

Establishment						
Name	Address	ID/FEI	Business Operations			
WOWBIOTECH		695625834	manufacture(76627-0007), label(76627-0007), pack(76627-0007)			

Revised: 5/2020

WOWBIOTECH