

WATSON HAND SANITIZER ALOE- alcohol gel

Tash Market Place

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.

watson Hand sanitizer aloe

Drug Facts

Active ingredient

Ethyl alcohol 62%

Purpose

Antimicrobial

Use

- Hand sanitizer to help reduce bacteria on the skin

Warnings

Flammable. Keep away from fire or flame.

For external use only

When using this product

rinse eyes thoroughly with water.

Stop use and ask a Doctor

if irritation and redness develop condition persists for more than 72 hours.

Keep out of reach of children.

Poison Control Center right away.

Directions

- Rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product.

Other information

- May discolor certain fabrics or surfaces

Inactive ingredients

Water, Carbomer, Glycerin, Aminomethyl Propanol, Propylene Glycol, Tocopheryl Acetate, Barbados Leaf Extract, Fragrance (Parfun), Blue 1.

Package Labeling:59ml

FRONT LABEL



Package Labeling:237ml

BACK LABEL



FRONT LABEL

WATSON

Hand Sanitizer

Kills 99.9% of Germs
Fast & Effective
With Added Moisturizer



8 FL OZ (237 mL)

BACK LABEL

Drug Facts

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Ethyl alcohol 62%.....	Antimicrobial

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Inactive ingredients

Water, Carbomer, Glycerin, Aminomethyl, Propanol, Propylene Glycol, Tocopheryl Acetate, Aloe Barbadensis Leaf Extract, Fragrance (Parfum), Blue 1.



DANGER



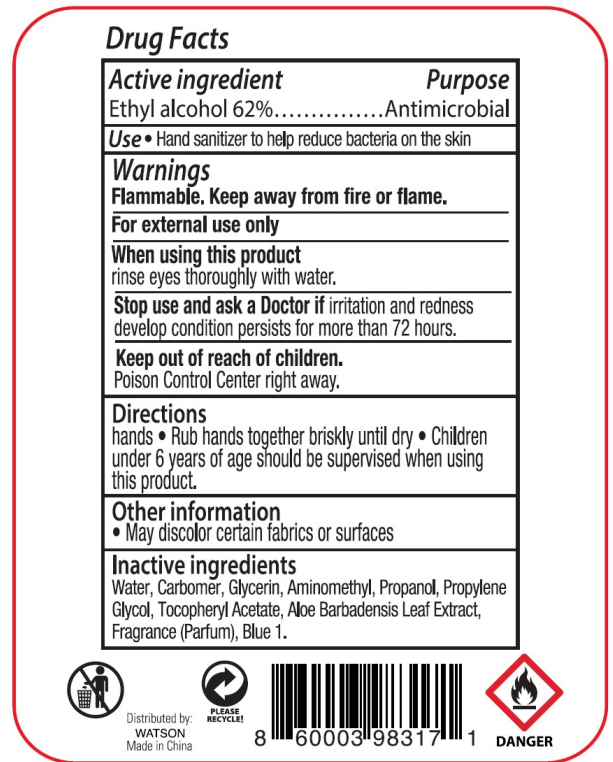
Distributed by:
WATSON
Made in China

Package Labeling:500ml

FRONT LABEL



BACK LABEL



Package Labeling:999ml

FRONT LABEL

WATSON

Hand Sanitizer

Kills 99.9% of Germs
Fast & Effective
With Added Moisturizer



33.8 FL OZ (999 mL)

BACK LABEL

Drug Facts

Active ingredient	Purpose
Ethyl alcohol 62%.....	Antimicrobial

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Distributed by:
WATSON
Made in China

WATSON HAND SANITIZER ALOE

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:90061-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	0.62 mL in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
GLYCERIN (UNII: PDC6A3C0OX)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90061-010-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
2	NDC:90061-010-02	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
3	NDC:90061-010-03	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
4	NDC:90061-010-04	999 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333E	08/18/2020	

Labeler - Tash Market Place (041805793)