

INSTANT HAND SANITIZER WITH ALOE- ethyl alcohol gel
SAFEWAY INC

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.

HAND SANITIZER WITH ALOE

ACTIVE INGREDIENT

ETHYL ALCOHOL 65% (ANTISEPTIC)

USES

TO HELP REDUCE BACTERIA ON THE SKIN

WARNINGS

- FOR EXTERNAL USE ONLY.
- FLAMMABLE. KEEP AWAY FROM FIRE OR FLAME.

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE WITH WATER.

STOP USE AND ASK A DOCTOR IF

SKIN IRRITATION OR REDNESS DEVELOPS.

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

OTHER INFORMATION

STORE AT A TEMPERATURE BELOW 110⁰F (43⁰C).

PACKAGE FRONT AND BACK LABELS

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Drug Facts

Active ingredient	Purpose
Ethyl Alcohol 65%	Antiseptic

Uses ■ to help reduce bacteria on the skin.

Warnings ■ For external use only. ■ Flammable, keep away from fire or flame.

When using this product ■ avoid contact with eyes. If contact occurs, rinse with water.

Stop using and ask a doctor if ■ irritation or redness develops and lasts.

Keep out of reach of children. ■ In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions ■ Wet hands thoroughly and rub together until dry.

Other information Store below 110°F (43°C).

Questions/Comments? 1-888-723-3929

Inactive ingredients: Water (Aqua), Isopropyl Alcohol, Glycerin, Carbomer, Aminomethyl Propanol, Fragrance (Parfum), Propylene Glycol, Isopropyl Myristate, Aloe Barbadosensis Leaf Juice, Tocopheryl Acetate, Blue 1 (CI 42090), Yellow 5 (CI 19140).

*This product not manufactured by Johnson & Johnson Consumer Products Company, distributors of Purell® Hand Sanitizer.

DISTRIBUTED BY SAFEWAY INC.
P.O. BOX 99, PLEASANTON, CA 94566-0099
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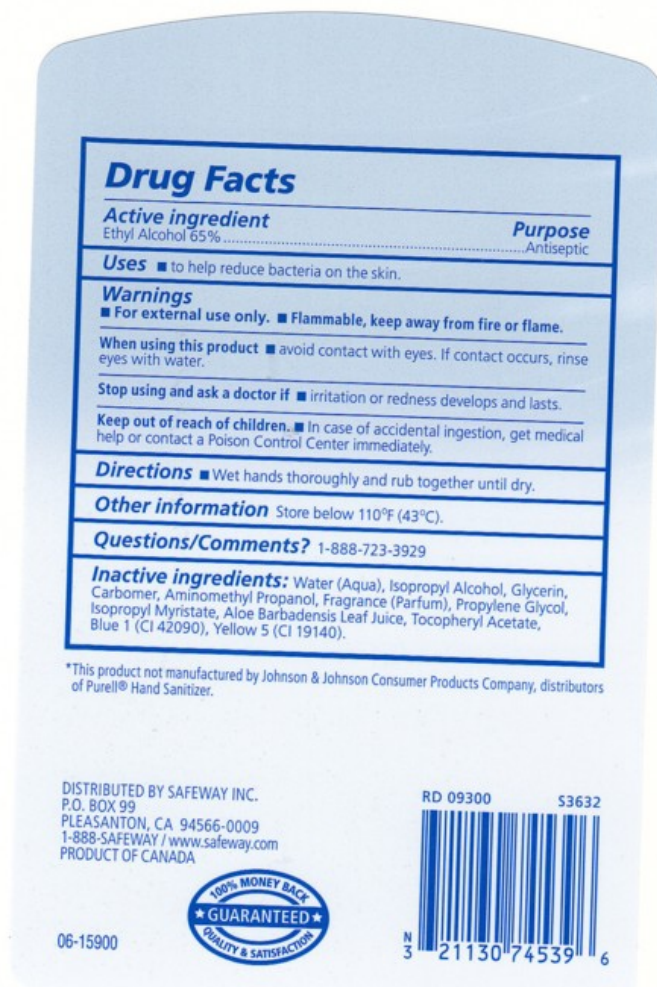
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32 FL OZ (946 mL)



INSTANT HAND SANITIZER WITH ALOE

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-254
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65.0000 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.0001 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:21130-254-02	59 mL in 1 BOTTLE, PLASTIC		
2	NDC:21130-254-32	946 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333	02/20/2010	

Labeler - SAFEWAY INC (009137209)

Revised: 2/2010

SAFEWAY INC