INSTANT HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E- ethyl alcohol gel RITE AID CORPORATION

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 MOISTURIZERS AND VITAMIN E

ACTIVE INGREDIENT

ETHYL ALCOHOL 62% (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

• RA2FRONT.jpg



Drug Facts

Active ingredient
Ethyl Alcohol 62%

Purpose Antimicrobial

Uses ■ to help reduce bacteria on the skin.

Warnings

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

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

Stop using and ask a doctor ■ if irritation or rash appears and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions ■ Wet hands thoroughly and rub together until dry. ■ Children under 6 should be supervised when using this product.

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

Inactive ingredients: Water (Aqua), Glycerin, Isopropyl Myristate, Propylene Glycol, Tocopheryl Acetate, Aminomethyl Propanol, Carbomer, Fragrance (Parfum).

*Not manufactured or distributed by Johnson and Johnson, the owner and distributor of Purell® Hand Sanitizer.



DISTRIBUTED BY:
Rite Aid Corporation
30 Hunter Lane, Camp Hill, PA 17011

Product of Canada







INSTANT HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E

ethyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11822-2250

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:11822-2250-8	237 mL in 1 BOTTLE, PUMP				
2	NDC:11822-2250-2	59 mL in 1 BOTTLE, PLASTIC				

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

Labeler - RITE AID CORPORATION (014578892)

Revised: 2/2010 RITE AID CORPORATION