HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E- ethyl alcohol gel HEB

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.

BACK LABEL - DRUG FACTS BOX

ACTIVE INGREDIENT

ETHYL ALCOHOL 62% (ANTISEPTIC)

USES AND DIRECTIONS

- USES: TO HELP REDUCE BACTERIA ON THE SKIN.
- **DIRECTIONS:** PUT A SMALL AMOUNT INTO YOUR PALM AND BRISKLY RUB HANDS TOGETHER UNTIL DRY.

WARNINGS

 FOR EXTERNAL USE ONLY.IN CASE OF ACCIDENTAL INGESTION, SEEK MEDICAL ASSISTANCE OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

WHEN USING THIS PRODUCT

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

STOP USE AND ASK A DOCTOR IF

SKIN IRRITATION OR REDNESS DEVELOPS.

KEEP OUT OF REACH OF CHILDREN

FOR CHILDREN UNDER 6, USE ONLY WITH ADULT SUPERVISION. NOT RECOMMENDED FOR USE ON INFANTS.

OTHER INFORMATION

FLAMMABLE. KEEP AWAY FROM HEAT AND FLAME.

PACKAGE FRONT AND BACK LABELS

• 8OZ FRONT AND BACK LABELS: HCF8OZ.jpg



HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E

ethyl alcohol gel

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-225

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)

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Packaging

l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:37808-225-08	237 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	07/05/2010	
OIC monograph not final	part333	0 //0 5/20 10	

Revised: 7/2010 HEB