HAND SANITIZER SUN BLOSSOM WITH MOISTURE BEADS- ethyl alcohol gel APOLLO HEALTH AND BEAUTY CARE

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 MOISTURE BEADS SUN BLOSSOM

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.

QUESTIONS/COMMENTS?:

1-866-428-7327

OTHER INFORMATION

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

Package Front and Back Labels

• 2OZ Front and Back Labels: eob2.jpg



• 9OZ Front and Back Labels: eob9.jpg



8.75 FL OZ (259 mL)



HAND SANITIZER SUN BLOSSOM WITH MOISTURE BEADS

ethyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG **Route of Administration**

Item Code (Source)

NDC:63148-261

TOPICAL

Active Ingredient/A	ctive Moiety			
Ingredient Name			Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	$62.0000\ mL$ in 100 mL
Packaging				
# Item Code	Package Description	Marketing Start Date		Marketing End Date
NDC:63148-261-02	59 mL in 1 BOTTLE, PLASTIC			
NDC:63148-261-09	259 mL in 1 BOTTLE, PUMP			
Marketing Inform	mation			
Marketing Category	Application Number or Monograph Citation		Marketing Start Da	te Marketing End Date
OTC monograph not final	part333		02/15/2010	

Labeler - APOLLO HEALTH AND BEAUTY CARE (201901209)

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