HAND SANITIZER PREVAIL- alcohol spray
Creation's Garden Natural Products, Inc

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Drug Facts

Ethyl Alcohol 62%.....Antiseptic

Uses

- To decrease bacteria on the skin that could cause disease.
- Recommended for repeated use.

Warnings

When using this product:

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water
- Avoid contact with broken skin
- For external use only-skin
- Flammable. Keep away from heat and flame.
- Stop use and ask doctor if skin irritation develops

Keep out of reach of children

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately

Directions

- Wet skin area with product
- Rub hands together until dry
- supervise children with the use of this product

Other information

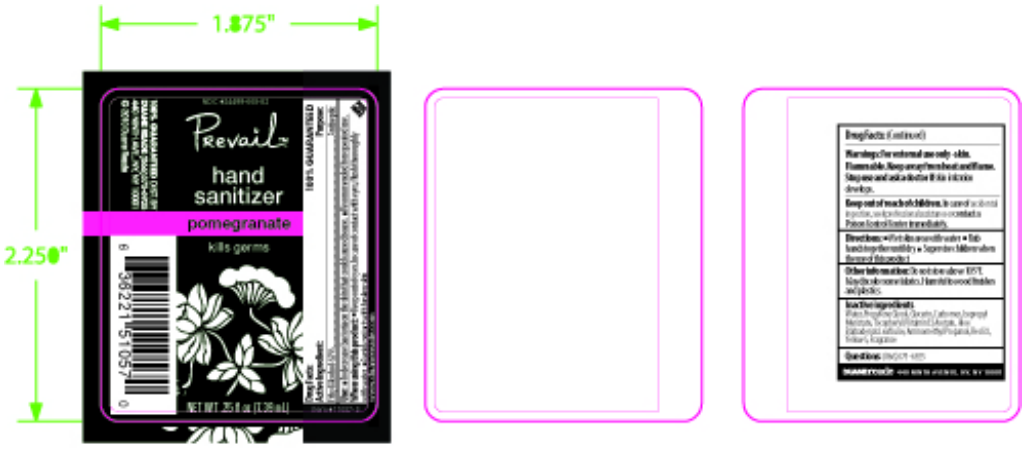
Do not store above 105°F. May discolor some fabrics. Harmful to wood finishes and plastics.

Inactive ingredients

Water, Propylene Glycol, Glycerin, Carbomer, Isopropyl Myristate, Tocopheryl (Vitamin E) Acetate, Aloe Barbadensis Leaf Juice, Aminomethyl Propanol, Yellow 5, Fragrance

Questions

(866) 375-6925



HAND SANITIZER PREVAIL

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24488-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	4.58 mL in 7.39 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24488-005-52	7.39 mL in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/12/2010	

Labeler - Creation's Garden Natural Products, Inc (839471216)

Registrant - Creation's Garden Natural Products, Inc (839471216)

Establishment

Name	Address	ID/FEI	Business Operations
Creation's Garden Natural Products, Inc		839471216	manufacture

Revised: 4/2010 Creation's Garden Natural Products, Inc