SKIN CRAVE NON-ALCOHOL HAND SANITIZER- antibacterial non-alcohol hand sanitizer spray
Tropical Enterprises International, 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.

Antibacterial Non-Alcohol Hand Sanitizer Benzalkonium Spray

ACTIVE INGREDIENT

Benzalkonium Chloride 0.13%

DOSAGE AND ADMINISTRATION

Directions: Pump or spray a small amount of foam into palm of hand. Rub thoroughly over all surfaces of both hands. Rub hands together briskly until dry.

INACTIVE INGREDIENT

Water, 2-Phenoxyethanol

INDICATIONS AND USAGE

Uses: For hand sanitizing to decrease bacteria on the skin. Recommended for repeat use.

KEEP OUT OF REACH OF CHILDREN

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

OTC PURPOSE

Purpose... Antimicrobial

WARNINGS

WARNINGS For external use only

SKIN CRAVE NON-ALCOHOL HAND SANITIZER

antibacterial non-alcohol hand sanitizer spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58418-786
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	13 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

2-(2-(2-(2-PHENOXYETHOXY)ETHOXY)ETHOXY)ETHANOL (UNII: Y050HYR4XA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58418-786-00	10 mL in 1 PACKET; Type 0: Not a Combination Product	08/01/2012	
2	NDC:58418-786-05	15 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/01/2012	
3	NDC:58418-786-10	10 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/01/2012	
4	NDC:58418-786-01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/01/2012	
5	NDC:58418-786-02	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/01/2012	
6	NDC:58418-786-04	120 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/01/2012	
7	NDC:58418-786-08	240 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/01/2012	
8	NDC:58418-786-12	360 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2012	
9	NDC:58418-786-16	480 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2012	
10	NDC:58418-786-64	1920 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2012	
11	NDC:58418-786-28	3840 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2012	
12	NDC:58418-786-17	50.275 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2012	

Labeler - Tropical Enterprises International, Inc. (091986179)

Registrant - Tropical Enterprises International, Inc. (091986179)

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