FRESH DECOR CITRUS POMEGRANATE- ethyl alcohol liquid
SAFEWAY 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.

DRUG FACTS

ACTIVE INGREDIENT

ETHYL ALCOHOL 65%

PURPOSE

ANTISEPTIC

USES

TO DECREASE BACTERIA ON THE SKIN.

WARNINGS

FOR EXTERNAL USE ONLY. FLAMMABLE, KEEP AWAY FROM SOURCES OF HEAT OR FIRE.

STOP USING AND ASK A DOCTOR

IF IRRITATION OR REDNESS DEVELOPS AND LASTS.

WHEN USING THIS PRODUCT

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

KEEP OUT OF REACH OF CHILDREN.

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

DIRECTIONS

WET HANDS THOROUGHLY AND RUB TOGETHER UNTIL DRY.

OTHER INFORMATION

DO NOT STORE ABOVE 110F (43C)

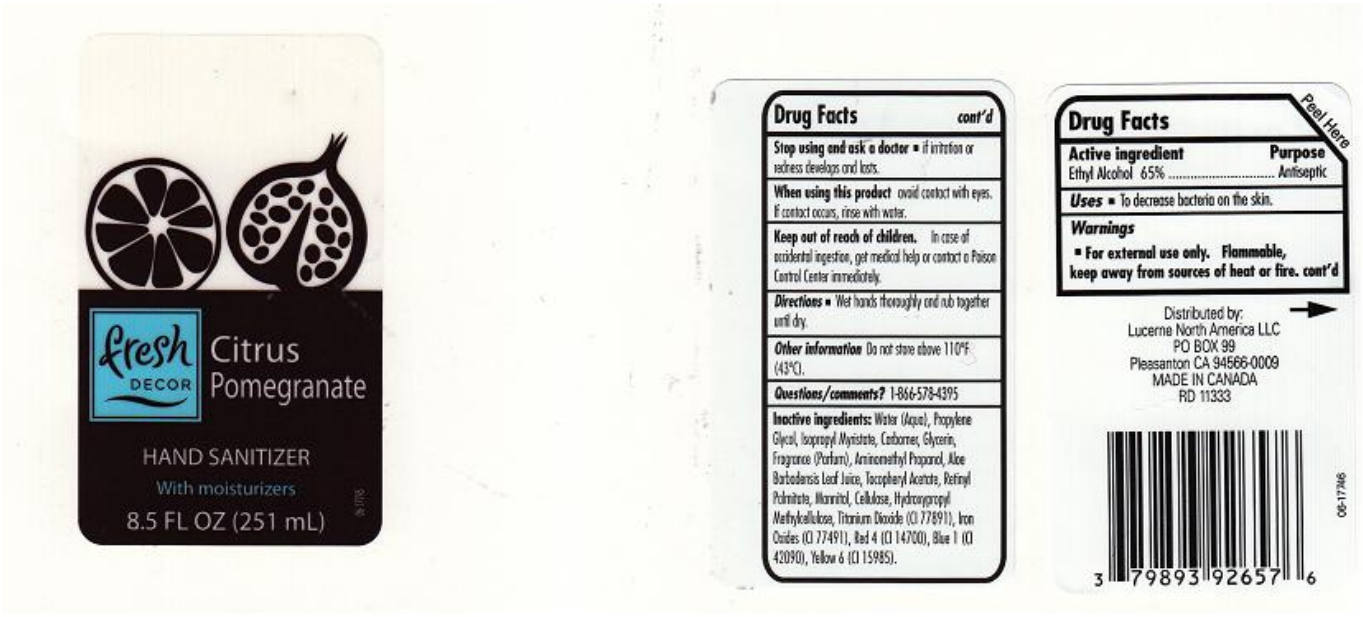
QUESTIONS/COMMENTS?

1-866-578-4395

INACTIVE INGREDIENTS:

WATER (AQUA), PROPYLENE GLYCOL, ISOPROPYL MYRISTATE, CARBOMER, GLYCERIN, FRAGRANCE (PARFUM), AMINOMETHYL PROPANOL, ALOE BARBADENSIS LEAF JUICE, TOCOPHERYL ACETATE, RETINYL PALMITATE, MANNITOL, CELLULOSE, HYDROXYPROPYL METHYLCELLULOSE, TITANIUM DIOXIDE (CI 77891), IRON OXIDES (CI 77491), RED 4 (CI 14700), BLUE 1 (CI 42090), YELLOW 6 (CI 15985).

LABEL COPY



FRESH DECOR CITRUS POMEGRANATE

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21130-463
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65.0 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
CARBOMER 934 (UNII: Z135WT9208)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	

VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
MANNITOL (UNII: 3OWL53L36A)	
ALPHA CELLULOSE (UNII: I355QGZ19A)	
HYPROMELLOSE 2208 (15000 MPA.S) (UNII: Z78RG6M2N2)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-463-09	251 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/03/2012	

Labeler - SAFEWAY INC. (009137209)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture

Revised: 12/2011 SAFEWAY INC.