FOAMING HAND SANITIZER WITH ALOE- benzalkonium chloride 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 BOX (BACK LABEL)

ACTIVE INGREDIENT

BENZALKONIUM CHLORIDE 0.1% (ANTISEPTIC)

USES AND DIRECTIONS

- **USES:** TO HELP REDUCE BACTERIA ON THE SKIN THAT COULD CAUSE DISEASE. RECOMMENDED FOR REPEATED USE.
- **DIRECTIONS:** PUMP ENOUGH PRODUCT TO YOUR PALM TO THOROUGHLY COVER YOUR HANDS, RUB TOGETHER UNTIL DRY.

WARNINGS

- FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT

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

STOP USE AND ASK A DOCTOR IF

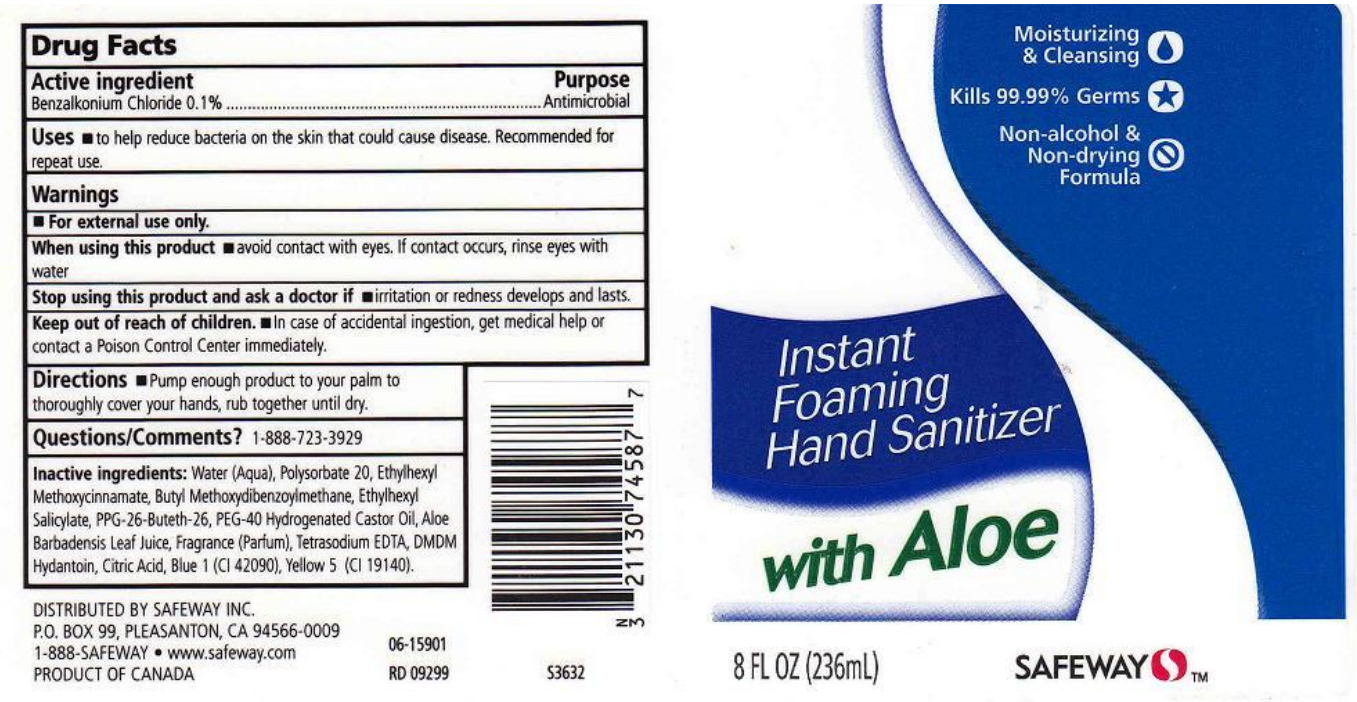
- SKIN IRRITATION OR REDNESS DEVELOPS AND LASTS.

KEEP OUT OF REACH OF CHILDREN

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

PACKAGE FRONT AND BACK LABELS

- 8OZ FRONT AND BACK LABELS: safeway8.jpg



QUESTIONS/COMMENTS?

1-888-723-3929

FOAMING HAND SANITIZER WITH ALOE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21130-240
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1000 mL in 100 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:21130-240-08	236 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	07/14/2010	

Labeler - SAFEWAY INC (009137209)

Revised: 7/2010 SAFEWAY INC