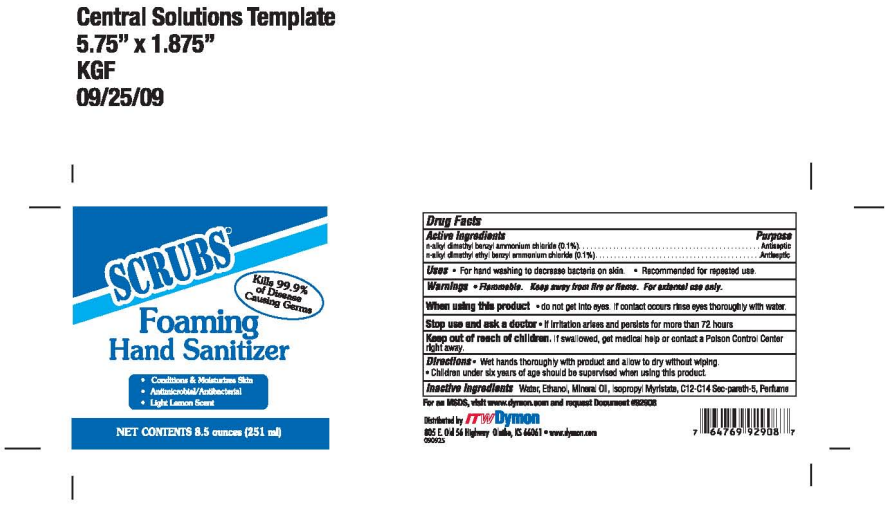
SCRUBS FOAMING HAND SANITIZER SCRUBS- hand sanitizer foam solution
ITW Dymon

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.



Active Ingredients *Purpose*

n-alkyl dimethyl benzyl ammonium chloride (0.1%).....Antiseptic

n-alkyl dimethyl ethyl benzyl ammonium chloride (0.1%).....Antiseptic

Stop use and ask a doctor

- if irritation arises and persists for more than 72 hours

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

When using this product

- do not get into eyes. If contact occurs rinse eyes thoroughly with water.

Uses

- For hand washing to decrease bacteria on skin.
- Recommended for repeated use.

Directions
- Wet hands thoroughly with product and allow to dry without wiping.
- Children under six years of age should be supervised when using this product.

Warnings
Flammable, keep away from heat and flame.
For external use only.

Inactive Ingredients
Water, ethanol, Mineral Oil, Isoproply Myristate, C12-C14 Sec-pareth-5

SCRUBS FOAMING HAND SANITIZER SCRUBS

hand sanitizer foam solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51239-1929
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.1 g in 970 mL
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 970 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Mineral Oil (UNII: T5L8T28FGP)	0.7 g in 970 mL
Isopropyl Myristate (UNII: 0RE8K4LNJS)	0.3 g in 970 mL
Alcohol (UNII: 3K9958V90M)	20 g in 970 mL
Water (UNII: 059QF0KO0R)	78.02 g in 970 mL

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor	LEMON	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:51239-1929-7	252 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	10/01/2009	

Labeler - ITW Dymon (103307604)

Registrant - ITW Dymon (103307604)

Establishment

Name	Address	ID/FEI	Business Operations
ITW Dymon		103307604	manufacture

Revised: 10/2009 ITW Dymon