KAY QSR- triclosan solution
Kay Chemical Co.

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

Triclosan 0.3%

Purpose

Antiseptic handwash

Uses

- For handwashing to decrease bacteria on the skin

Warnings

For external use only

- In eyes

- If in eyes, rinse promptly and throughly with water
- Discontinue use if irritation and redness develop

Stop use and ask a doctor if

- Skin irritation or redness occurs for more than 72 hours

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

Directions

- Wet hands and forearms
- Dispense a palmful of product to hands
- Scrub hands and forearms for 20 seconds
- Rinse throughly and dry

Other Information

- for additional information, see Material Safety Data Sheet (MSDS)
- **Medical Emergency:** (877) 231-2615 or call collect 0 (952) 853-1713

Inactive ingredients water (aqua), potassium cocoate, alcohol, sodium laureth sulfate, hexylene glycol, boric acid, cocamidopropyl PG-dimonium chloride phosphate, tetrasodium EDTA, cocamine oxide, fragrance, PEG-75 lanolin, methylparaben, propylparaben, aloe barbadensis leaf juice, FDC Blue 1

Questions? Call **1-800-529-5458**

Principal display panel and representative label

NDC NO: 63146-118-06

KAY

QSR Anti-Bacterial

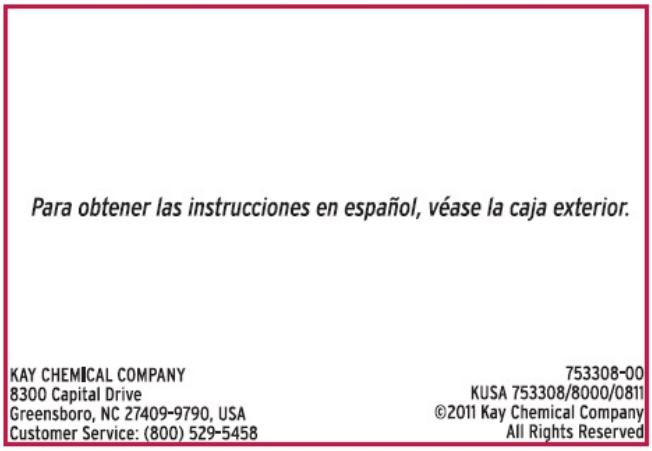
Foam Hand Soap

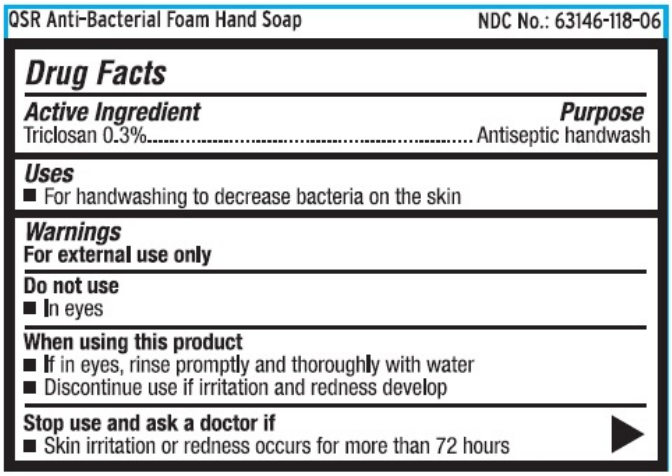
See inside back label for complete drug facts panel

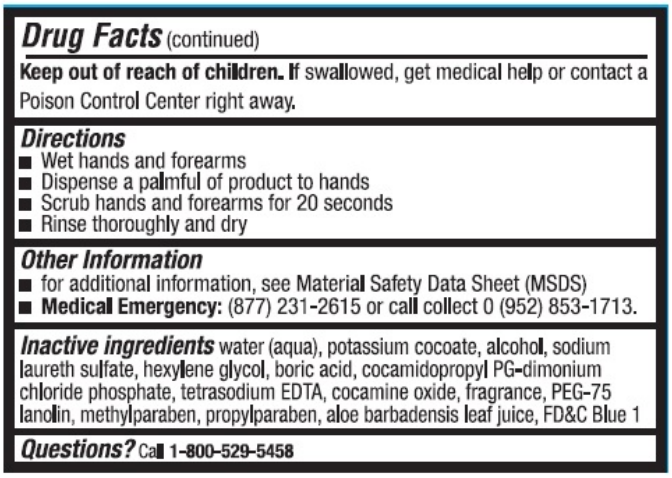
To obtain Spanish instructions, see outer carton.

Triclosan 0.3%

40.6 US fl oz (1200 ml)







KAY QSR

triclosan solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63146-118
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.3 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
POTASSIUM COCOATE (UNII: F8U72V8ZXP)	
ALCOHOL (UNII: 3K9958V90M)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
BORIC ACID (UNII: R57ZHV85D4)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
EDETATE SODIUM (UNII: MP1J8420LU)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63146-118-06	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/30/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/30/2011	

Labeler - Kay Chemical Co. (003237021)

Revised: 10/2019 Kay Chemical Co.