KAY ALCOHOL FOAM HAND SANITIZER - alcohol solution
Kay Chemical Company

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, 62% w/w (equivalent to 70% v/v)

Purpose

Antiseptic handwash

Uses

- for hand washing to decrease bacteria on the skin

Warnings

- **For external use only**
- **FLAMMABLE, keep away from fire or flame, heat, sparks and sources of static discharge.**

Do not use

- in eyes

When using this product

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

Stop use and ask doctor if skin irritation and redness persists from more than 72 hours

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

Directions

- wash hands to remove soil
- spread to cover hands thoroughly, rub to dry

Other information

- for additional information, see Safety Data Sheet (SDS)
- for emergency medical information in USA, call (877) 231-2615 or call collect - 0

(952) 853-1713

Inactive ingredients water (aqua), PEG-10 dimethicone, ethylhexylglycerin, farnesol, bisabolol, tert-butyl alcohol, denatonium benzoate

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

Principal display panel and representative label

NDC 63146-313-09 KAY

Alcohol Foam

Hand Sanitizer

Active ingredient: Ethyl alcohol 62% w/w (equivalent to 70% v/v)

Net contents: 40.6 US fl oz (1200 mL)

SDS-NC-872, SDS-WI-15014, SDS-NJ-20007

Distributed by

Kay Chemical Company

8300 Capital Drive - Greensboro NC 27409-9790 USA

Customer Service: (800) 529-5458

Made in USA

KAY ALCOHOL FOAM HAND SANITIZER

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63146-313
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
FARNESOL (UNII: EB41QIU6JL)	
LEVOMENOL (UNII: 24WE03BX2T)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63146-313-09	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/02/2020	

Labeler - Kay Chemical Company (003237021)

Revised: 11/2022 Kay Chemical Company