HAND SANITIZER- alcohol gel 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.

Advanced Hand Sanitizer 370

Active Ingredient

Ethyl alcohol 70%

purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommend for repeated use

Warnings

For external use only-hands

Flammable

Keep away from heat and flame

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

• skin irritation or redness develops

Keep out of reach of children

If swallowed get medical help or contact a Poison Control Center right away

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 1050 F
- may discolor some fabrics

• harmful to wood finishes and plastics

Inactive ingredients

water, glyceryl caprylate/caprate, glycerin, isopropyl myristate, tocopheryl acetate, acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4

Questions?

1-888-723-3929

*Effective at eliminating more than 99.99% of many common harmful germs and bacteria in as little as 15 seconds

**This product is not manufactured or distributed by GOJO Industries, Inc. distributor of Purell Advanced Hand Sanitizer Refreshing Gel

SDS-MO-15036 SDA-WI-2486 DSP-MO-28 DSP-MO-34

DISTRIBUTED BY SAFEWAY INC.

P.O. BOX 99

PLEASANTON, CA 94566-0009

1-888-SAFEWAY www.safeway.com

OUR PROMISE QUALITY & SATISFACTION 100% GUARANTEED OR YOUR MONEY BACK. 370.000/370AB

PRINCIPAL DISPLAY PANEL

SAFEWAY CARE

ADVANCED HAND SANITIZER

Kills more than 99.99% of germs*

Compare to Purell Advanced Hand Sanitizer**

Refreshing Gel

Quality guaranteed

32 FL OZ (946 mL)



HAND SANITIZER

alcohol gel

Prod	nct l	Info	rma	tion
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:21130-370

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

indute ingredients				
Ingredient Name	Strength			
water (UNII: 059QF0KO0R)				
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)				
GLYCERIN (UNII: PDC6 A3C0 OX)				

ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)		
.ALPHATO COPHERO L ACETATE (UNII: 9E8X80D2L0)		
CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809 Y72KV36)		
SULISOBENZONE (UNII: 1W6L629B4K)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-370- 16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/06/2014	
2	NDC:21130-370- 34	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/06/2014	
3	NDC:21130-370- 45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/06/2014	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/06/2014	

Labeler - Safeway Inc. (009137209)

Registrant - Vi-Jon (790752542)

Establishment				
Name	Address	ID/FEI	Business Operations	
Vi-Jon		088520668	manufacture(21130-370)	

Revised: 5/2020 Safeway Inc.