WINTER APPLE- hand sanitizer gel
Kroger, 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.

Winter Apple Hand Sanitizer

30142-803-16

Active ingredient

Ethyl alcohol 62%

Purpose

Antiseptic

Uses

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

Warnings

For external use only: hands

Flammable. Keep away from fire or 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

- irritation or redness develops
- condition 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.

Directions

- wet hand 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 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glycerin, carbomer, fragrance, red 40, red 33

EFFECTIVE AT ELIMINATING 99.99% OF MANY COMMON HARMFUL GERMS AND BACTERIA IN AS LITTLE AS 15 SECONDS*

DISTRIBUTED BY THE KROGER CO., CINCINNATI, OHIO 45202

QUALITY GUARANTEE

800-632-6900 www.kroger.com

Principal Panel Display

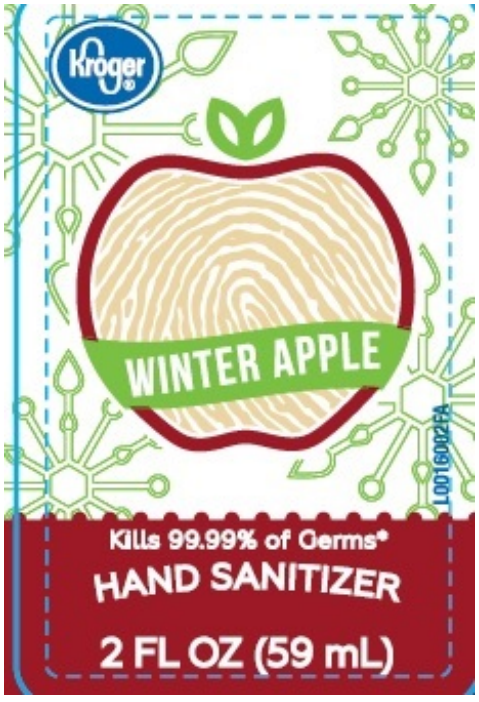
Kroger

WINTER APPLE

Kills 99.99% of Germs

HAND SANITIZER

2 FL OZ (59 mL)



WINTER APPLE

hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:30142-803
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:30142-803-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/24/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/24/2016	

Labeler - Kroger, CO., (006999528)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(30142-803)

Establishment

Name	Address	ID/FEI	Business Operations

Vi-Jon, LLC		088520668	manufacture(30142-803)

Revised: 12/2022 Kroger, CO.,