

## **WATERLESS HAND SANITIZER- alcohol gel**

**Kroger**

*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.*

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### **Hand Sanitizer**

**203**

#### **Active Ingredient**

Ethyl Alcohol 65%

#### **Purpose**

Antiseptic

#### **Use**

- to decrease bacteria on the skin that could cause disease
- recommended 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.
- do not inhale or ingest
- avoid contact with broken skin

#### **Stop use and ask a doctor if**

skin irritation 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 105°F •may discolor some fabrics
- harmful to wood finishes and plastics

#### **Inactive ingredients**

aloe barbadensis leaf juice, benzophenone-4, blue 1, carbomer, fragrance, glycerin, isopropyl myristate, propylene glycol, tocopheryl acetate, water, yellow 5

Not manufactured or distributed by GOJO Industries, owner of the registered trademark Purell

Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds.  
SDS-MO-15036 SDA WI-2059 SDA WI-2486 DSP-MO-28 DSP-MO-34

Distributed by the Kroger Co.  
Cincinnati, Ohio 45202

203.000/203AB

25% MORE than the 8 fl oz size

Kroger

Waterless Hand Sanitizer

Kills 99.99% of Germs

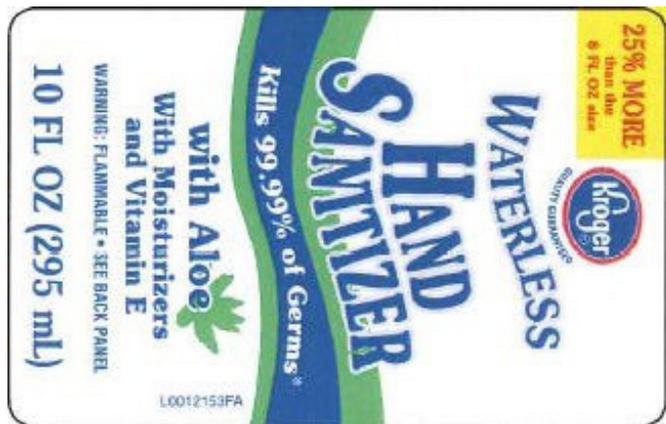
with Aloe

With Moisturizers

and vitamin E

WARNING: FLAMMABLE•SEE BACK PANEL

10 FL OZ (295 mL)



## WATERLESS HAND SANITIZER

alcohol gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30 142-203
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	.65 mL in 100 L

### Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
SULISOBENZONE (UNII: 1W6L629B4K)	

FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-203-34	.236 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/12/1997	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/12/1997	

**Labeler -** Kroger (006999528)

**Registrant -** Vi-Jon (088520668)

**Establishment**

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(30142-203)