WATERLESS HAND SANITIZER- alcohol gel
Onpoinr, 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.

Drug Facts

Active Ingredient

Ethyl Alcohol 65%

Purpose

Antiseptic

Use

•to decrease bacteria on the skin that could cause disease
•recommended for reported 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 •may discolor some fabrics
•harmful to wood finishes and plastics

Inactive ingredients

aloe barbadensis leaf juice, benzophenone-4, blue 1, carbomer, fragrance, glycerin,

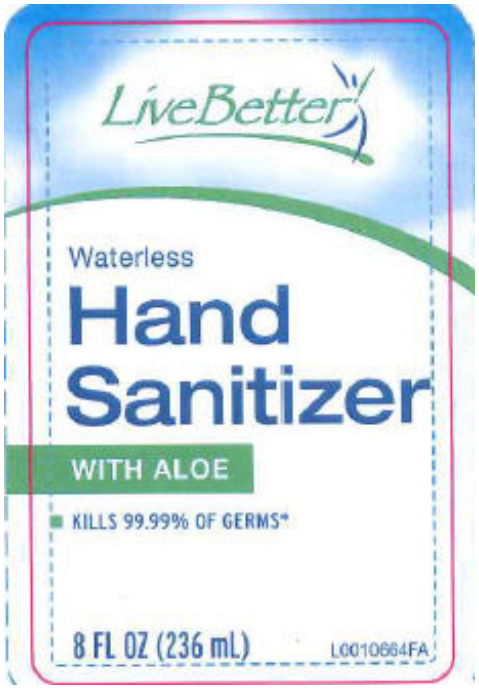
isopropyl myristate, propylene glycol, tocopheryl acetate, water, yellow 5

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.

Distributed by: Onpoint, Inc
2 Paragon Drive,
Montvale, NJ 07645

Live Better
Waterless Hand Sanitizer With Aloe
•Kills 99.99% of Germs
8 FL OZ (236 mL)



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WATERLESS HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51143-203
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: ZY81Z83H0X)	
SULISOBENZONE (UNII: 1W6L629B4K)	

FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
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:51143-203-34	.236 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/01/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2011	

Labeler - Onpoinr, Inc (001367366)

Registrant - Vi-Jon (088520668)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(51143-203)

Revised: 11/2018 Onpoinr, Inc