### PURELL- alcohol gel Navajo Manufacturing Company Inc.

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Purell

### **Drug Facts**

### Active ingredient

Ethyl Alcohol 70% v/v

### **Purpose**

Antimicrobial

#### Uses

Hand sanitized to help reduce bacteria on the skin

### Warnings

Flammable. Keep away from fire or flame.

For external use only.

# When using this product

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

# Stop use and ask a doctor if

irritation or rash appears and lasts

# Keep out of reach of children.

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

#### **Directions**

- Put enough product in your palm to cover hands and rub hands together briskly until dry
- Children under 6 years of age should be supervised when using this product

#### Other information

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces

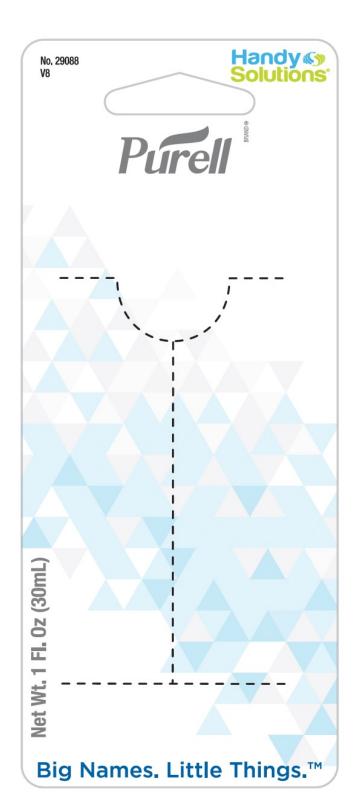
# Inactive ingredients

Water (Aqua), Isopropyl Alcohol, Caprylyl Glycol, Glycerin, Isopropyl Myristate, Tocopheryl Acetate, Acrylates/C10-30, Alkly Acrylate, Crosspolymer, Aminomethyl Propanol, Fragrance (Parfum)

**Questions?** 

1-888-4-PURELL www.PURELL.com

Purell, 1oz



No. 29088 V8





Kills more than 99.99% of most common germs that may cause illness

### **Drug Facts**

#### lises

Hand sanitizer to help reduce bacteria on the skin

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#### Questions? 1-888-4-PURELL www.PURELL.com

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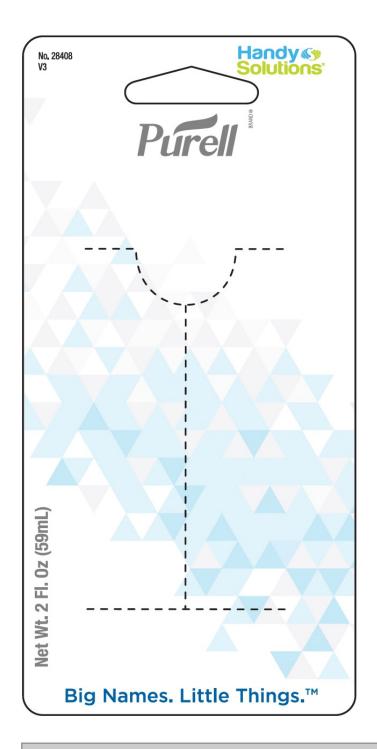
Please Recycle



www.navajoinc.com



Purell, 2oz



No. 28408



Kills more than 99.99% of most common germs that may cause illness

#### **Drug Facts**

Active ingredient Ethyl Alcohol 70% v/v ..... Purpose Antimicrobial

•Hand sanitizer to help reduce bacteria on the skin

### Warnings

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#### Questions? 1-888-4-PURELL www.PURELL.com

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#### **PURELL**

alcohol gel

### **Product Information**

**Route of Administration** 

**Product Type** HUMAN OTC DRUG Item Code (Source) NDC:67751-188(NDC:21749-704)

**TOPICAL** 

# Active Ingredient/Active Majety

ı	Active Ingredient/Active Molety				
	Ingredient Name	<b>Basis of Strength</b>	Strength		
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL		

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ISOPROPYL ALCOHOL (UNII: ND2M416302)		
CAPRYLYL GLYCOL (UNII: 00YIU5438U)		
GLYCERIN (UNII: PDC6A3C0OX)		
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)		

F	Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:67751-188- 01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2017		
2	NDC:67751-188- 02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/04/2017	09/01/2021	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	12/04/2017		

# Labeler - Navajo Manufacturing Company Inc. (091917799)

# Registrant - Navajo Manufacturing Company Inc. (091917799)

Establishment				
Name	Address	ID/FEI	Business Operations	
Navajo Manufacturing Company Inc.		136941411	relabel(67751-188), repack(67751-188)	

Revised: 4/2024 Navajo Manufacturing Company Inc.