

## **SANNYTIZE HAND SANITIZER- alcohol gel**

### **Dynarex Corporation**

*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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**1427 Dynarex SannyTize Hand Sanitizer NDC 67777-317-01**  
**1428 Dynarex SannyTize Hand Sanitizer NDC 67777-317-02**  
**1429 Dynarex SannyTize Hand Sanitizer NDC 67777-317-03**  
**1431 Dynarex SannyTize Hand Sanitizer NDC 67777-317-04**  
**1432 Dynarex SannyTize Hand Sanitizer NDC 67777-317-05**  
**1433 Dynarex SannyTize Hand Sanitizer NDC 67777-317-06**

### **ACTIVE INGREDIENT**

Active Ingredient	Purpose
Ethyl Alcohol 0.62%	Antiseptic Handwash

### **WARNINGS**

- **For external use only**
- **Flammable, keep away from fire and flame**
- **Do not use** in or near the eyes
- **When using this product** avoid contact with eyes. In case of eye contact, rinse with water to remove.
- **Stop use and ask a doctor if** irritation and redness develops

### **PURPOSE**

An antiseptic handwash.

### **INDICATIONS & USAGE**

Cleaning and antiseptic cleansing of hands and skin.

### **DOSAGE & ADMINISTRATION**

- Wet hands thoroughly with product and allow to dry without wiping

### **KEEP OUT OF REACH OF CHILDREN**

**KEEP OUT OF REACH OF CHILDREN**, if swallowed get medical help or contact a Poison Control Center right away.

### **INACTIVE INGREDIENTS**

Inactive ingredients: Water, Glycerin, Propylene glycol, Carbomer, Triethanolamine, Aloe barbadensis juice, Fragrance

**Label**

**dynarex**

**SannyTize™**

**Instant Hand Sanitizer**

*With Moisturizing Aloe*

**Kills More Than 99.99% of Germs\***

**Reorder No. 1432**

16 fl. oz. (473 mL)

<b>Drug Facts</b>	
<b>Active Ingredient</b>	<b>Purpose</b>
Ethyl Alcohol 70% .....	Antiseptic Handwash
<b>Use</b>	
For hand washing to decrease bacteria on the skin	
<b>Warnings</b>	
For external use only	
<b>Flammable:</b> Keep away from fire or flame	
<b>Do not use</b> in the eyes	
<b>When using this product</b>	
<ul style="list-style-type: none"> <li>■ Avoid contact with eyes</li> <li>■ In case of eye contact, rinse with water to remove</li> </ul>	
<b>Stop use and ask a doctor</b> if irritation and redness develop	
<b>Keep out of reach of children</b> If swallowed, get medical help or contact a Poison Control Center right away	
<b>Directions</b>	
Wet hands thoroughly with product and allow to dry without wiping	
<b>Other Information</b>	
<ul style="list-style-type: none"> <li>■ Do not store above 110°F (43°C)</li> <li>■ May discolor some fabrics</li> <li>■ Recommended for repeated use</li> </ul>	
<b>Inactive Ingredients</b>	
Aloe Vera Leaf Extract, Carbopol Ultrez 10, Fragrance, Glycerin, Propylene Glycol, Purified Water, Triethanolamine	
* Effective at eliminating more than 99.99% of many common harmful germs and bacteria.	
<b>Manufactured for:</b> Dynarex Corporation 10 Glenshaw Street Orangeburg, NY 10962 USA • www.dynarex.com Made in India R201125	
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<b>SANNYTIZE HAND SANITIZER</b>			
alcohol gel			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:67777-317
<b>Route of Administration</b>	TOPICAL		
<b>Active Ingredient/Active Moiety</b>			
Ingredient Name	Basis of Strength	Strength	

<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL
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### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>CARBOMER COPOLYMER TYPE A</b> (UNII: 71DD5V995L)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-317-01	29.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/25/2017	
2	NDC:67777-317-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/25/2017	
3	NDC:67777-317-03	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/25/2017	
4	NDC:67777-317-04	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/25/2017	
5	NDC:67777-317-05	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/25/2017	
6	NDC:67777-317-06	1.035 mL in 1 PACKET; Type 0: Not a Combination Product	04/25/2017	
7	NDC:67777-317-07	1.035 mL in 1 PACKET; Type 0: Not a Combination Product	04/25/2017	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/25/2017	

**Labeler** - Dynarex Corporation (008124539)

**Registrant** - Dynarex Corporation (008124539)