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.

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 andredness 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

SANNYTIZE HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-317
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients

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

Revised: 3/2021 Dynarex Corporation