SANNYTIZE HAND SANITIZER- hand sanitizer gel
Dynarex Corporation

1427 Hand Sanitizer, 1 oz. 67777-010-01, 1428 Hand Sanitizer, 2 oz. 67777-010-02
1429 Hand Sanitizer, 4 oz. 67777-010-03, 1431 Hand Sanitizer, 8 oz. 67777-010-04
1432 Hand Sanitizer, 16 oz. 67777-010-05, 1433 Hand Sanitizer, .9g 67777-010-06
1433UB-10 Hand Sanitizer, 0.9g 67777-010-20

Active Ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic Handwash

Use

- For hand washing to decrease bacteria on the skin.

Warnings

For External Use Only

Flammable. Keep away from fire or flame

Do not use

In 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 develop.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

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

Other Information

- Do not store above 110°F (43ºC)
- May discolor some fabrics
- Recommended for repeated use

Inactive Ingredients

Aloe Vera Leaf Extract, Carbopol Ultrez 10, Fragrance, Glycerin, Propylene Glycol, Purified Water,Triethanolamine

Questions?

1-888-396-2739 Monday - Friday, 9AM - 5PM EST

Label

1428 Hand Sanitizer

Label

1427 Hand Sanitizer

Label

1427 Hand Sanitizer

Label

1429 Hand Sanitizer

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1431 Hand Sanitizer

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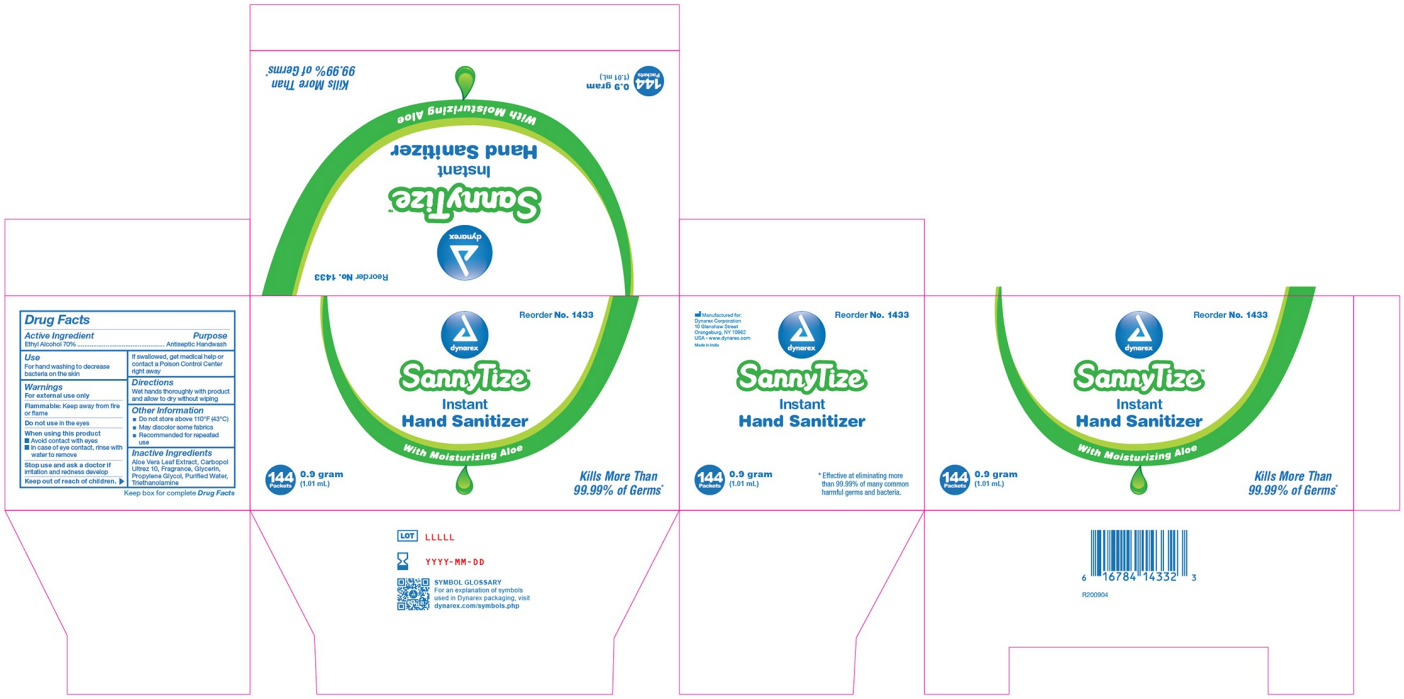
1432 Hand Sanitizer

Label



1433 Hand Sanitizer

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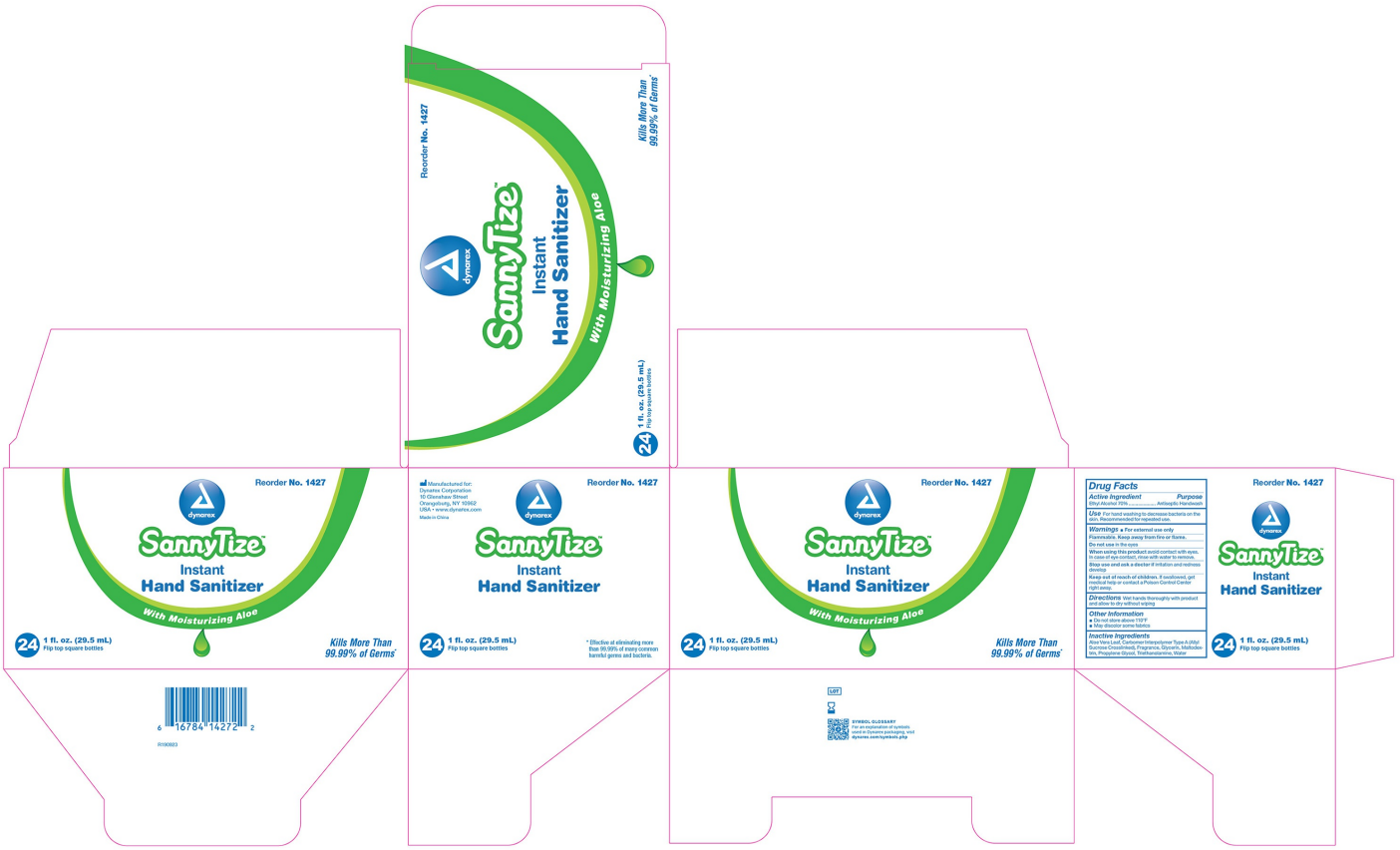

1427 Hand Sanitizer

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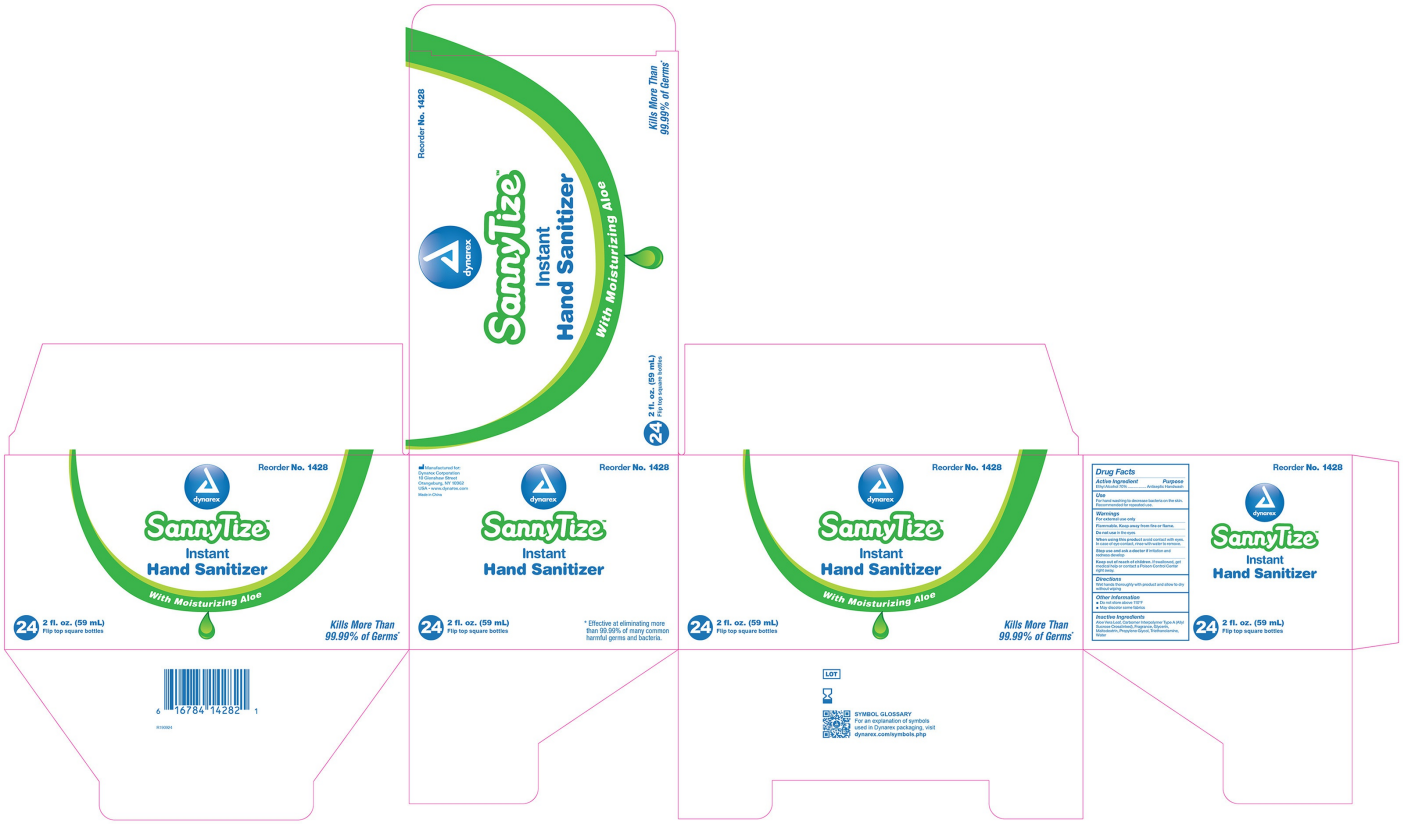

1428 Hand Sanitizer

Label

1429 Hand Sanitizer

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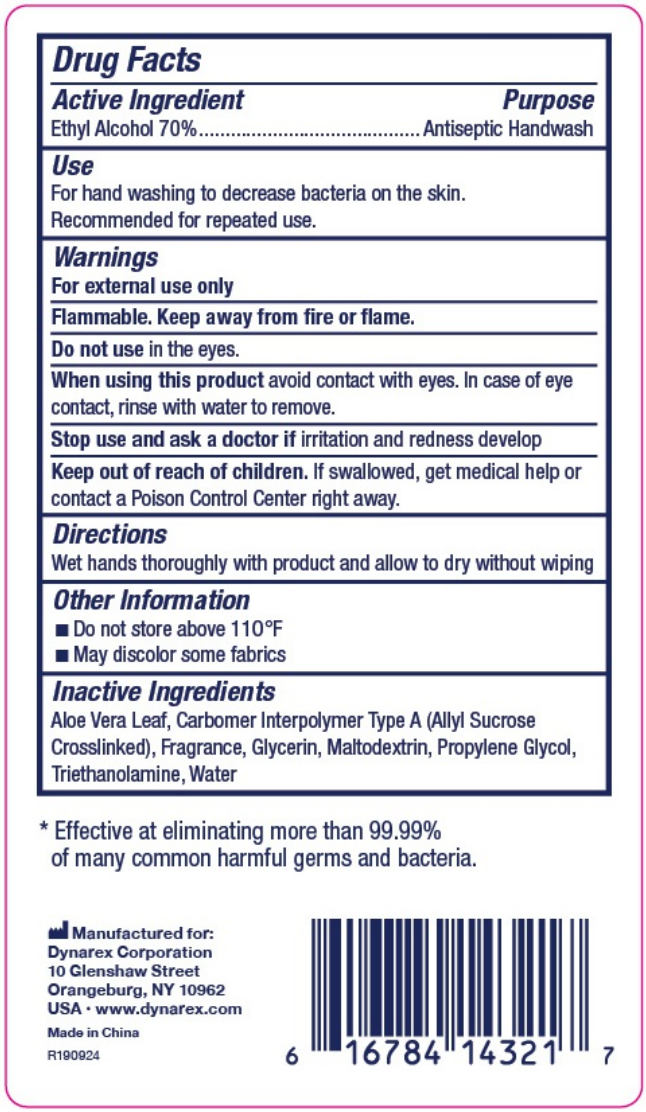
1431 Hand Sabitizer

Labels



1432 Hand Sanitizer

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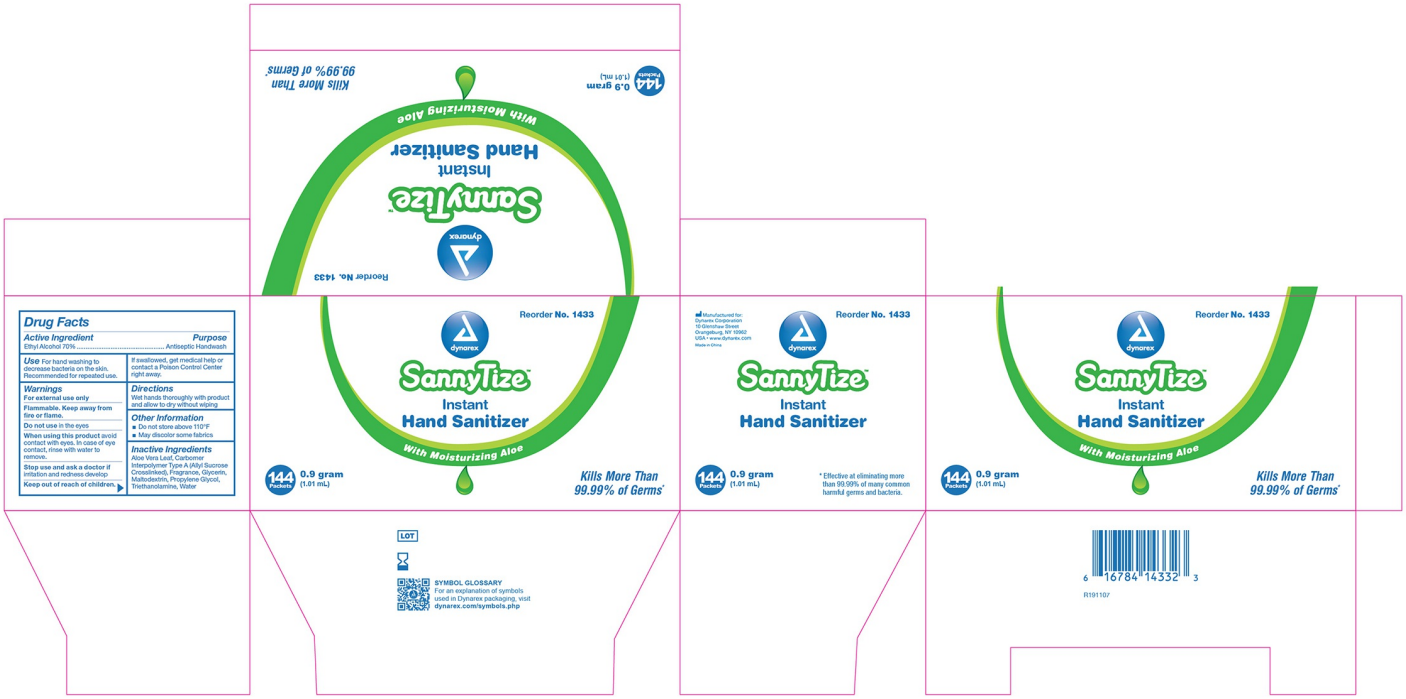

1433 Hand Sanitizer

Label 1433UB-10

1433UB-10

SANNYTIZE HAND SANITIZER

hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	

Product Characteristics

Color		**Score**	
Shape	FREEFORM	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-010-01	144 in 1 CASE	11/01/2019	
1	NDC:67777-010-11	24 in 1 BOX		
1	NDC:67777-010-10	29.5 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:67777-010-02	144 in 1 CASE	11/01/2019	
2	NDC:67777-010-13	24 in 1 BOX		
2	NDC:67777-010-12	59 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:67777-010-03	96 in 1 CASE	11/01/2019	
3	NDC:67777-010-15	24 in 1 BOX		
3	NDC:67777-010-14	118.3 mL in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:67777-010-04	48 in 1 CASE	11/01/2019	
4	NDC:67777-010-16	236.5 mL in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:67777-010-05	12 in 1 CASE	11/01/2019	
5	NDC:67777-010-17	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

6	NDC:67777-010-06	1728 in 1 CASE	11/01/2019	
6	NDC:67777-010-18	144 in 1 BOX		
6		1.01 mL in 1 PACKET; Type 0: Not a Combination Product		
7	NDC:67777-010-20	1000 in 1 CASE	11/01/2019	
7	NDC:67777-010-19	10 in 1 BOX		
7		1.01 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	11/01/2019	

Labeler - Dynarex Corporation (008124539)

Revised: 1/2024 Dynarex Corporation