

CRANBERRY CHAMPAGNE HAND SANITIZER- cranberry champagne hand sanitizer liquid
Zhejiang Meimi Technology Co., Ltd.

Zhejiang Meimi Technology Co., Ltd.
Cranberry Champagne Hand Sanitizer 45ml
41366-084-01

Active Ingredient(s)

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Use

Hand sanitizer to help reduce bacteria on the skin that may cause disease.

Warnings

- for external use only.
- Flammable.
- Keep away from heat and flame.

Stop use and ask a doctor if irritation or redness develops.

Keep out of reach of children.
if swallowed, get medical help or contact a Poison Control Center.

Directions

Spray enough product in your palm to thoroughly cover your hands. Rub hands together briskly until dry. Recommended for repeated use. Children under 6 years of age should be supervised when using this product.

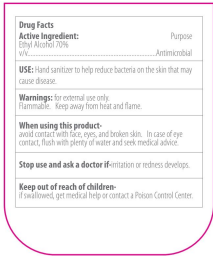
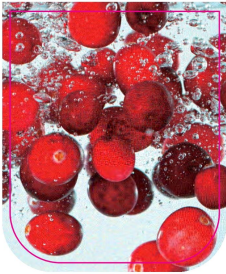
Inactive ingredients

Water(Aqua), Fragrance(Parfum), Glycerin, Benzyl Alcohol, Linalyl Acetate, Limonene, Linalool, Citronellol, Vanillin, Red 40.

When using this product:
avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Package Label - Principal Display Panel

45 mL in 1 BOTTLE
NDC: 41366-084-01



CRANBERRY CHAMPAGNE HAND SANITIZER

cranberry champagne hand sanitizer liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41366-084	
Route of Administration	EXTRACORPOREAL			
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	
VANILLIN (UNII: CHI530446X)				
GLYCERIN (UNII: PDC6A3C0OX)				
WATER (UNII: 059QF0KO0R)				
LIMONENE, (+)- (UNII: GFD7C86Q1W)				
LINALOOL (UNII: D81QY6I88E)				
RED 40 (UNII: WZB9127XOA)				
BENZYL ALCOHOL (UNII: LKG8494WBH)				
.BETA.-CITRONELLOL, (+/-)- (UNII: 565OK72VNF)				
LINALYL ACETATE (UNII: 5K47SSQ51G)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41366-084-01	45 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/10/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/10/2025	

Labeler - Zhejiang Meimi Technology Co., Ltd. (413668440)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Meimi Technology Co., Ltd.		413668440	manufacture(41366-084)

Revised: 6/2025

Zhejiang Meimi Technology Co., Ltd.