PALM PARTY- alcohol gel gel
Dongyang Loulee Cosmetics Co.,Ltd.

Active ingredient(s)

Alcohol 60%

Purpose(s)

Antisepti

Uses

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

- Flammable. Keep away from fire or flame.
- For external use only.

Do not use

In 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

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Inactive ingredients

WATER, FRAGRANCE, ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER, ARGININE, PHENOXYETHANOL

Questions or comments?

213-396-1046 / MalachiPerez291@gmail.com

Box Front Panel

PALM PARTY
alcohol gel gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87322-1614
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	60 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
FRAGRANCE 13576 (UNII: 5EM498GW35)	5 mL in 100 mL
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.4 mL in 100 mL
WATER (UNII: 059QF0KO0R)	34.2 mL in 100 mL
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	0.2 mL in 100 mL
ARGININE (UNII: 94ZLA3W45F)	0.2 mL in 100 mL

Product Characteristics

Color	white (Transparent gel)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87322-1614-1	30 mL in 1 CARTON; Type 0: Not a Combination Product	01/16/2026	
2	NDC:87322-1614-2	35 mL in 1 CARTON; Type 0: Not a Combination Product	01/16/2026	
3	NDC:87322-1614-3	42 mL in 1 CARTON; Type 0: Not a Combination Product	01/16/2026	
4	NDC:87322-1614-4	100 mL in 1 CARTON; Type 0: Not a Combination Product	01/16/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/16/2026	

Labeler - Dongyang Loulee Cosmetics Co.,Ltd. (529518089)

Registrant - Dongyang Loulee Cosmetics Co.,Ltd. (529518089)

Establishment

Name	Address	ID/FEI	Business Operations
Nanjing Ningjitong Pharmaceutical Biotechnology Co., Ltd		450168555	manufacture(87322-1614)

Establishment

Name	Address	ID/FEI	Business Operations
Dongyang Loulee Cosmetics Co., Ltd		529518089	label(87322-1614)

Revised: 1/2026

Dongyang Loulee Cosmetics Co.,Ltd.