DR. PALM HAND SANITIZER- alcohol denat gel
SKINNYDIP LIMITED

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.

Dr. Palm

Active Ingredient

Alcohol 75%

Purpose

Antiseptic

Uses

- Hand sanitizer to help reduce bacteria on the skin that could cause disease.
- For use when soap and water are not available

warnings

For external use only. Flammable. Keep away from heat or flame.

- **Stop use and ask a doctor** if irritation or rash occurs. These may be signs of a serious condition.

Do not use

- in children less than 2 months of age.
- on open skin wounds.

When using this product

- keep out of eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Directions

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

Keep out of reach of children.

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

Inactive Ingredients.

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Extract, Aminomethyl Propanol,Butylene Glycol, Glycerin, Water (Aqua).

Other information.

- Store between 15-30°C (59-86°F)
- Avoid freezing and excessive heat above 40°C(104°F)

Product Image

DR. PALM HAND SANITIZER

alcohol denat gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81236-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81236-001-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
2	NDC:81236-001-02	35 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
3	NDC:81236-001-03	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
4	NDC:81236-001-04	75 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
5	NDC:81236-001-05	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
6	NDC:81236-001-06	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
7	NDC:81236-001-07	465 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
8	NDC:81236-001-08	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
9	NDC:81236-001-09	975 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2020	
10	NDC:81236-001-10	27 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/08/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/20/2020	

Labeler - SKINNYDIP LIMITED (217094578)

Revised: 4/2021 SKINNYDIP LIMITED