PHARMACY PRESCRIPTION 1.8OZ HAND SANITIZER - SWEET PEACH- alcohol gel
Ningbo Liyuan Daily Chemical Products Co.,Ltd.

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.

Active Ingredient

Ethyl Alcohol 62.0%

Purpose

Antimicrobial

Uses

- Helps reduce bacteria on the skin that could cause disease.

Recommended for repeated use.

Warnings

For external use only.

Flammable. Keep away from fire or flame.

When using this product

do not use in or near the eyes. In case of eye contact, flush eyes thoroughly with water.

Stop use and ask a doctor if:

irritation and redness develops and lasts more than 72 hours.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions-

Place enough product in palm to thoroughly cover your hands. Rub hands together briskly until dry.

Children under 6 years of age should be supervised when using this product.

Other information: Store below 110 Fahrenheit degree (43 degrees Celsius)

May discolor certain fabrics or surfaces.

Inactive Ingredients:

Water, Aloe Barbadensis Gel, Carbomer, Fragrance, Glycerin, Triethanolamine, Tocopheryl Acetate, (Sweet Peach FD&C Red No. 40, FD&C Yellow No. 5)

Drug Facts

PHARMACY PRESCRIPTION 1.8OZ HAND SANITIZER - SWEET PEACH

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76176-072
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76176-072-01	53 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/01/2018	

Labeler - Ningbo Liyuan Daily Chemical Products Co.,Ltd. (530766098)

Registrant - Ningbo Liyuan Daily Chemical Products Co.,Ltd. (530766098)

Establishment

Name	Address	ID/FEI	Business Operations
Ningbo Liyuan Daily Chemical Products Co.,Ltd.		530766098	manufacture(76176-072)

Revised: 1/2018 Ningbo Liyuan Daily Chemical Products Co.,Ltd.