HANDY ANDYS- hand sanitizer gel gel
LXR Biotech LLC

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.

LXR-Handy Andy Sanitizer Gel

Active Ingredient(s)

Alcohol 80% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

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

Warnings

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

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.
Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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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.

Other information

Store between 59-86F (15-30C)
Avoid freezing and excessive heat above 104F (40C)
May discolor certain fabrics or surfaces

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP, acrylate, copolymer

Package Label - Principal Display Panel

57 mL NDC: 78299-142-11

HANDY ANDYS
hand sanitizer gel gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78299-142
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0KO0R)	
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78299-142-12	57 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/12/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/12/2020	

Labeler - LXR Biotech LLC (117520926)

Establishment

Name	Address	ID/FEI	Business Operations
LXR Biotech LLC		968357405	manufacture(78299-142)

Revised: 1/2022 LXR Biotech LLC