

HAND SANITIZER- alcohol gel
NTML Group 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.

INMO 70% Alcohol Hand Gel (30ml)

Hand Sanitizer with NDC 00000-000 is a a human over the counter drug product labeled by NTML Group Ltd. The generic name of Hand Sanitizer is alcohol. The product's dosage form is gel and is administered via topical form.

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

For handwashing to decrease bacteria on the skin

Recommended for repeated use

Warnings

For external use only

Flammable, keep away from fire and flame

Do not drink

If taken internally will produce serious gastric disturbances

Do not use

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

When using this product avoid the eyes and mucous membrane

In the case of eyes or mucous membranes contact, rinse area thoroughly with water

Stop use and ask a doctor if condition worsens, redness or irritation develops

If condition persists for more than 3 days

Keep out of reach of children. If swallowed contact a doctor or Poison Control Center immediately.

Directions

Rub dime sized amount between hands until dry

Supervise children in the use of this product

In the case of eye contact, rinse eyes thoroughly with water

Other information

Store below 105 F

May discolour some fabrics

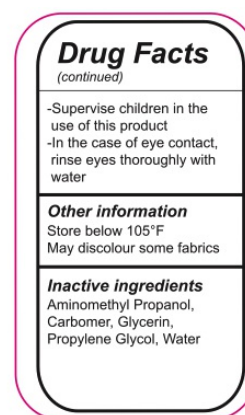
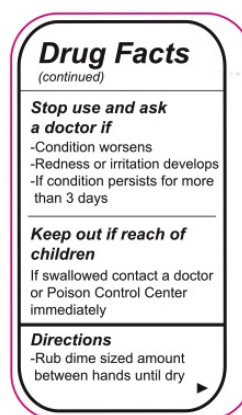
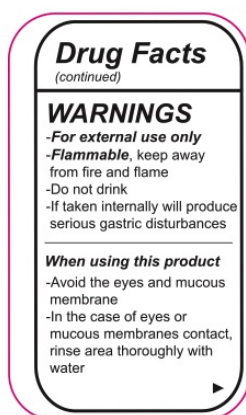
Inactive ingredients

Aminomethyl Propanol, Carbomer, Glycerin, Propylene Glycol, Water.

Package Label - Principal Display Panel

70% Alcohol Gel

30 mL (1 fl oz)



HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77366-001
Route of Administration	TOPICAL		

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
WATER (UNII: 059QF0K00R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77366-001-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - NTML Group Ltd (664159445)

Revised: 4/2020

NTML Group Ltd