

ZAPRO HAND SANITIZER- hand sanitizer gel
GUANGDONG MINGLIU TECHNOLOGY DAIL-USE CHEMICAL 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.

75209-011
ZAPRO HAND SANITIZER

Active Ingredient(s)

Ethyl Alcohol 94%

Purpose

Antiseptic

Use

Use
for hand washing to decrease bacteria on he skin only when water is not ava able

Warnings

Flammable—Keep away from fire and flames
For external use only

Do not use

Do not get into eyes
If contact occurs, rinse thoroughly with water

When using this product

When using this product
Do not get into eyes
If contact occurs, rinse thoroughly with water

Stop use and ask a doctor if
Irritation and redness develop

Keep out of reach of children.If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and ask a doctor if
Irritation and redness develop

Keep out of reach of children.If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Press pump twice to dever two s quits of product on to the pam of your hand
Rub hands together and allow to dry without wiping

Other information


May discolor fabrics or surfaces
Store below 110°F(43°C)

Inactive ingredients

water, cellulosico-polymer, zinc pyrithione

Package Label - Principal Display Panel

75209-011-01 250ml



ZaPro™
HAND SANITIZER

NEW ADVANCED FORMULA

PROFESSIONAL STRENGTH

- Disinfects even through residual bio-fluid contamination, such as blood or saliva
- Residual Effect - Continues to work after application.
- Doesn't dry out skin/No sticky residue

NET QTY: 8.5 FL. OZ. (250 ML)

NON-TACKY
Designed to assist in easily donning latex and rubber gloves before procedures

Drug Facts	Purpose
Active Ingredients Ethyl Alcohol 94%	Antiseptic
Use for hand washing to decrease bacteria on he skin, only when water is not available	
Warnings ■ Flammable — Keep away from fire and flames ■ For external use only	
When using this product ■ Do not get into eyes ■ If contact occurs, rinse thoroughly with water	
Stop use and ask a doctor if ■ Irritation and redness develop	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions ■ Press pump twice to deliver two squirts of product onto the palm of your hand ■ Rub hands together and allow to dry without wiping	
Other Information ■ Store below 110° F (43° C) ■ May discolor fabrics or surfaces	
Inactive Ingredients water, cellulosico-polymer, zinc pyrithione	
Questions? 877-877-9778 M-F 7AM-5PM MST	


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West Jordan, UT 84084 USA
PAK-LA0080.v10.20

REORDER #006-00282

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75209-011-02 2000ml



ZaPro™
HAND SANITIZER

NEW ADVANCED FORMULA

PROFESSIONAL STRENGTH

- Disinfects even through residual bio-fluid contamination, such as blood or saliva
- Residual Effect - Continues to work after application.
- Doesn't dry out skin/No sticky residue

NET QTY: 67.6 FL. OZ. (2L)

NON-TACKY
Designed to assist in easily donning latex and rubber gloves before procedures

Drug Facts	Purpose
Active Ingredients Ethyl Alcohol 94%	Antiseptic
Use for hand washing to decrease bacteria on he skin, only when water is not available	
Warnings ■ Flammable — Keep away from fire and flames ■ For external use only	
When using this product ■ Do not get into eyes ■ If contact occurs, rinse thoroughly with water	
Stop use and ask a doctor if ■ Irritation and redness develop	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions ■ Press pump twice to deliver two squirts of product onto the palm of your hand ■ Rub hands together and allow to dry without wiping	
Other Information ■ Store below 110° F (43° C) ■ May discolor fabrics or surfaces	
Inactive Ingredients water, cellulosico-polymer, zinc pyrithione	
Questions? 877-877-9778 M-F 7AM-5PM MST	

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ZAPRO HAND SANITIZER

hand sanitizer gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:75209-011

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	94 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5)	
WATER (UNII: 059QF0KO0R)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75209-011-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/22/2020	
2	NDC:75209-011-02	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/22/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/22/2020	

Labeler - GUANGDONG MINGLIU TECHNOLOGY DAIL-USE CHEMICAL CO., LTD. (546631081)

Establishment

Name	Address	ID/FEI	Business Operations
GUANGDONG MINGLIU TECHNOLOGY DAIL-USE CHEMICAL CO., LTD.		546631081	manufacture(75209-011)

Revised: 12/2020

GUANGDONG MINGLIU TECHNOLOGY DAIL-USE CHEMICAL CO., LTD.