

**HAND SANITIZING MANDARIN- alcohol gel**  
**GANZ U.S.A., 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.*

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**time & again™**  
**HAND SANITIZING GEL**  
**MANDARIN**

***Drug Facts***

**Active ingredient**

Alcohol 62%

**Purpose**

Antiseptic

**Uses**

To decrease bacteria on hands.

**Warnings**

**For external use only.**

**Flammable, keep away from fire or flame.**

**When using this product**

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- Avoid contact with broken skin.
- Do not inhale or ingest.

**Stop use and ask a doctor** if skin irritation develops.

**Keep out of reach of children.** In case of accidental ingestion, seek professional assistance or contact Poison Control Center immediately.

**Directions**

Wet hands thoroughly with product and allow to dry without wiping.

**Other information**

- Do not store above 105°F.
- May discolor some fabrics.
- Harmful to wood finishes and plastics.
- You may report a serious adverse reaction to this product to: Ganz U.S.A., LLC #043, 60 Industrial Parkway, Cheektowaga, New York, 14227-9903

**Inactive ingredients**

Water, Fragrance, Glycerin, Triethanolamine, PEG-7 Glyceryl Cocoate, Hydrolyzed Jojoba Esters, Chamomilla Recutita (Matricaria) Flower Extract, Salvia Officinalis (Sage) Leaf Extract, Urtica Dioica (Nettle) Extract, Rosmarinus Officinalis (Rosemary) Leaf Extract, Aloe Barbadensis Leaf Extract, Aloe Barbadensis Leaf Juice, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Yellow 6.

**PRINCIPAL DISPLAY PANEL - 232 mL Bottle Label**

*time & again*™

**HAND SANITIZING GEL**

Formulated with Glycerin,  
Aloe and Jojoba to Moisturize.

Kills harmful bacteria and germs.

**MANDARIN**

232 mL 7.8 fl. OZ.

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TA397328 Mandarin

GANZ  
Cheektowaga, N.Y.  
14227-9903  
www.timeandagain.com  
MADE IN CANADA

6 61371 49537 8

**HAND SANITIZING MANDARIN**

alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:75862-002
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mg in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
GLYCERYL COCOATE (UNII: WVK1CT5994)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
JOJOBA OIL (UNII: 724GKU717M)	
CHAMAEMELUM NOBILE FLOWER (UNII: O2T154T6OG)	
SAGE (UNII: 065C5D077J)	
URTICA DIOICA LEAF (UNII: X6M0DRN46Q)	
ROSEMARY (UNII: IJ67X351P9)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75862-002-01	232 mL in 1 BOTTLE, PUMP		
2	NDC:75862-002-02	59 mL in 1 BOTTLE		
3	NDC:75862-002-03	15 mL in 1 BOTTLE		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	01/01/2011	

**Labeler** - GANZ U.S.A., LLC (798785242)

Revised: 12/2010

GANZ U.S.A., LLC