

**TRICLOSAN- triclosan liquid**  
**GOJO Industries, Inc.**

*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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**Active ingredient**

Triclosan 0.3%

**Purpose**

Antimicrobial

**Use**

- Handwash to help decrease bacteria on the skin
- Recommended for repeated use

**Warnings**

For external use only

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts

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

**Directions**

- Wet hands
- Apply a small amount of product and work into a lather
- Rinse well and dry hands completely

**Inactive ingredients**

Water (Aqua), Coconut Acid, Sodium Sulfate, Oleic Acid, Ethanolamine, Aloe Barbadensis Leaf Juice, Cocamide MEA, Coco-Betaine, Hydrolyzed Vegetable Protein, Propylene Glycol, Retinyl Palmitate, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Hydroxypropyl Methylcellulose, Methylchloroisothiazolinone, Methylisothiazolinone

Brought to you by GOJO  
Llega a usted gracias a GOJO

**PROVON**  
BRAND

Medicated  
**Lotion Soap**  
with Triclosan

**Jabón Loción**  
Medicado  
con Triclosano



Distributed by / Distribuido por:  
**GOJO Industries, Inc.**, Akron, OH 44309  
800-321-9647 • 330-255-6000 www.GOJO.com  
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NDC 21749-415-10

**Drug Facts**

**Active ingredient**      **Purpose**  
Triclosan 0.3% ..... Antimicrobial

**Uses** • Handwash to help decrease bacteria on the skin • Recommended for repeated use

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**Datos Farmacológicos (cont'd)**

**Ingrediente activo**      **Propósito**  
Triclosano 0,3% ..... Antimicrobiano

**Usos** • Lavado de manos empleado para disminuir la cantidad de bacterias en la piel • Recomendado para uso reiterado

**Advertencias**

**Sólo para uso externo**

**Al utilizar este producto,** evitar el contacto con los ojos o con la zona alrededor de los ojos. En caso de contacto, enjuagar completamente los ojos con agua.

**Dejar de usar el producto y consultar a un médico** si aparece y persiste una irritación o erupción cutánea

**Mantener fuera del alcance de los niños.** En caso de ingestión, de inmediato acudir a un médico o ponerse en contacto con un centro para el control de tóxicos.

**Modo de Uso** • Mojarse las manos • Aplicar una pequeña cantidad del producto y frotar las manos hasta producir una espuma abundante • Enjuagar bien • Secarse las manos completamente

**Ingredientes inactivos**

Agua, Acido de coco, Sulfato de sodio, Acido oleico, Etanolamina, Zumo de la hoja de Aloe Barbadensis, MEA, cocamida, Coco-betaina, Proteina vegetal hidrolizada, Propilenglicol, Palmitato de retinilo, EDTA tetrasódico, Acetato de tocoferilo, Aceite de maíz (Zea Mays), Hidroxipropil metilcelulosa, Metilcloroisotiazolinona, Metilisotiazolinona

1 L (33.8 US/ÉU FL OZ)

2158

U.S. Pat. 6,619,512  
U.S. Pat. 6,877,642 • U.S. Pat. D431,404  
U.S. Pat. 6,216,916 • U.S. Pat. D432,547

**TRICLOSAN**

triclosan liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:21749-415
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	0.003 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCONUT ACID (UNII: 40U37V505D)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
OLEIC ACID (UNII: 2UM9U37CP)	

MONOETHANOLAMINE (UNII: 5KV86114PT)
ALOE VERA LEAF (UNII: ZY81Z83H0X)
COCO MONOETHANOLAMIDE (UNII: C80684146D)
COCO-BETAINE (UNII: 03DH2IZ3FY)
Propylene Glycol (UNII: 6DC9Q167V3)
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)
EDETATE SODIUM (UNII: MP1J8420LU)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
CORN OIL (UNII: 8470G57WFM)
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)
Methylchloroisothiazolinone (UNII: DEL7T5QRPN)
Methylisothiazolinone (UNII: 229D0E1QFA)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-415-10	1000 mL in 1 BAG; Type 0: Not a Combination Product	03/10/2010	
2	NDC:21749-415-20	2000 mL in 1 BOX; Type 0: Not a Combination Product	03/10/2010	
3	NDC:21749-415-16	473 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/10/2010	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/10/2010	12/01/2021

**Labeler** - GOJO Industries, Inc. (004162038)

**Registrant** - GOJO Industries, Inc. (004162038)

### Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-415)