

GOLDEN MEDICATED- medicated lotion soap solution
Kutol Products Company

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.

Kutol F060

Chloroxylenol 0.3% w/w.....Antibacterial Handwashing

Water, Tall Oil Acid, Potassium Hydroxide, Sodium Lauryl Sulfate, Coconut Acid, Cocamide DIPA, Phenoxyethanol, Tetrasodium EDTA, Sodium Sulfate, Sodium Laureth Sulfate, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate (Vitamin E), Iodopropynyl Butylcarbamate, Fragrance, Red 33, Yellow 5.

Handwash to help decrease bacteria on the skin.

For external use only.

Do not use in the eyes. In case of contact, immediately flush 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.

- Apply a small amount, covering hands with product for 30 seconds. Add water, lather and rinse.
- Children under 6 years of age should be supervised when using this product.

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GOLDEN		Contains NO Triclosan!
Medicated Lotion Soap		
Drug Facts		
Active Ingredient	Purpose	
Chloroxylenol 0.3% w/w	Antibacterial Handwashing	
Uses ■ Handwash to help decrease bacteria on the skin.		
Warnings		
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Directions ■ Apply a small amount, covering hands with product for 30 seconds. Add water, lather, rinse.		
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Inactive Ingredients Water, Tall Oil Acid, Potassium Hydroxide, Sodium Lauryl Sulfate, Coconut Acid, Cocamide DIPA, Phenoxyethanol, Tetrasodium EDTA, Sodium Sulfate, Sodium Laureth Sulfate, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate (Vitamin E), Iodopropynyl Butylcarbamate, Fragrance, Red 33, Yellow 5.		
KUTOL PRODUCTS CO. Hand Hygiene Specialists Cincinnati, Ohio www.kutol.com		Net Contents: 3.785 L (one gallon) Manufactured in the USA in a LEED® Silver certified facility



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SOFT & SILKY

Contains NO Triclosan!

Golden Medicated Lotion Soap
1000mL (33.8 Fl. Oz.) Boxless Bag

Drug Facts

Active Ingredient	Purpose
Chloroxylenol 0.3% w/w.....	Antibacterial Handwashing

Uses

■ Handwash to help decrease bacteria on the skin.

Warnings

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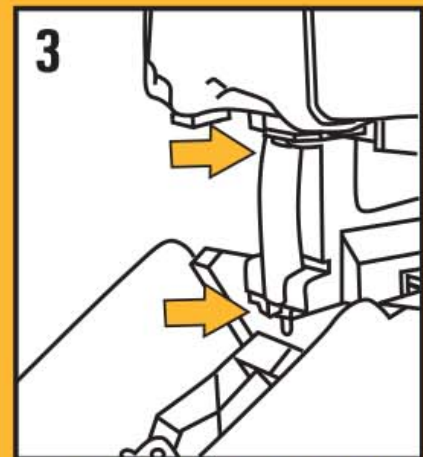
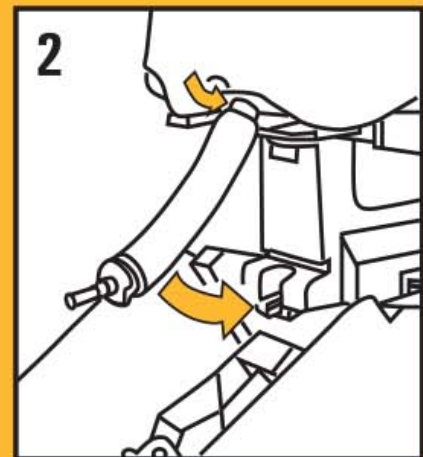
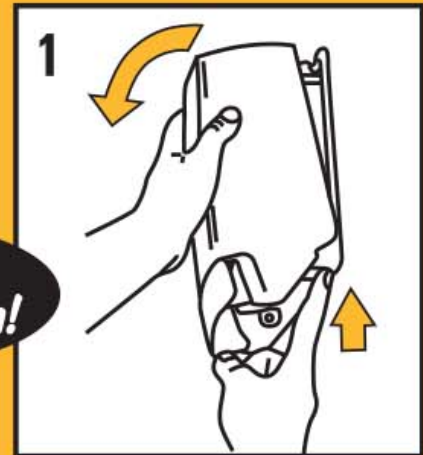
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Inactive Ingredients

Water, Tall Oil Acid, Potassium Hydroxide,



Close cover carefully

KUTOL

Hand Hygiene Specialists

Cincinnati, OH
1-800-543-4641

Sodium Lauryl Sulfate, Coconut Acid,
Cocamide DIPA, Phenoxyethanol, Tetrasodium
EDTA, Sodium Sulfate, Sodium Laureth Sulfate,
Aloe Barbadensis Leaf Juice, Tocopheryl
Acetate (Vitamin E), Iodopropynyl
Butylcarbamate, Fragrance, Red 33, Yellow 5.

www.kutol.com

Made in the USA in a
LEED® Silver certified facility

Net Contents 1000mL (33.8 Fl. Oz.)
Contenu net 1000mL
Contenido Neto 1000 mL

LA705R-0314

50865-060-27.jpg

GOLDEN MEDICATED

medicated lotion soap solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50865-060
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
COCONUT ACID (UNII: 40U37V505D)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
EDETATE SODIUM (UNII: MP1J8420LU)	
TALL OIL ACID (UNII: H9HR63474M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
COCO DIISOPROPANOLAMIDE (UNII: S485AM948Q)	

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End
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#	Item Code	Package Description	Date	Date
1	NDC:50865-060-09	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/11/2014	
2	NDC:50865-060-27	1000 mL in 1 BAG; Type 0: Not a Combination Product	03/11/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/11/2014	

Labeler - Kutol Products Company (004236139)

Registrant - Kutol Products Company (004236139)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products Company, Inc.		004236139	manufacture(50865-060)

Revised: 8/2023

Kutol Products Company