

**EXCELON ANTIMICROBIAL FOAM- chloroxylenol liquid**  
**GOJO Industries, Inc.**

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**EXCELON Antimicrobial Foam Soap**

**Active ingredient**

Chloroxylenol 0.3%

**Purpose**

Antimicrobial

**Uses**

- 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), Alcohol, Ammonium Laureth Sulfate, Ammonium Lauryl Sulfate, Propylene Glycol, Ammonium Xylenesulfonate, Cocamide MEA, Glycerin, Isopropyl Alcohol, Lactic Acid, Retinyl Palmitate, Simmondsia Chinensis (Jojoba) Seed Oil, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Ammonium Sulfate, Fragrance (Parfum), Methylchlorisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 6 (CI 15985)

E6162-04



Made in USA for:  
 Hecho en los E.E.U.U. por:  
**Excelon, Inc.**  
**Akron, OH 44309**  
**1-800-321-9647**  
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**ANTIMICROBIAL FOAM SOAP**  
**ESPUMA DE JABÓN**  
**ANTIMICROBIANA**

1.25L (42 FL OZ)

6162-640-EXL-F

**Drug Facts**

<b>Active Ingredient</b>	<b>Purpose</b>
Chloroxylenol 0.3%.....	Antimicrobial

**Uses**

- 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 product and thoroughly cover hands with lather
- Rinse well and dry hands completely

**Drug Facts (continued)**

**Directions**

- Wet hands
- Apply product and thoroughly cover hands with lather
- Rinse well and dry hands completely

**Inactive Ingredients** Water (Aqua), Alcohol, Ammonium Laureth Sulfate, Ammonium Lauryl Sulfate, Propylene Glycol, Ammonium Xylenesulfonate, Cocamide MEA, Glycerin, Isopropyl Alcohol, Lactic Acid, Retinyl Palmitate, Simmondsia Chinensis (Jojoba) Seed Oil, Tetrasodium EDTA, Tocopheryl Acetate, Zea Mays (Corn) Oil, Ammonium Sulfate, Fragrance (Parfum), Methylchloroisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 6 (CI 15985)

5362-644-ES-E

**EXCELON ANTIMICROBIAL FOAM**

chloroxylenol liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:21749-580
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
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<b>CHLOROXYLENOL</b> (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.003 mg in 1 mL
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### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>AMMONIUM LAURETH-2 SULFATE</b> (UNII: 698O4Z48G6)	
<b>AMMONIUM LAURYL SULFATE</b> (UNII: Q7AO2R1M0B)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>AMMONIUM XYLENESULFONATE</b> (UNII: 4FZY6L6XCM)	
<b>COCO MONOETHANOLAMIDE</b> (UNII: C80684146D)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)	
<b>LACTIC ACID</b> (UNII: 33X04XA5AT)	
<b>VITAMIN A PALMITATE</b> (UNII: 1D1K0N0VVC)	
<b>JOJOBA OIL</b> (UNII: 724GKU717M)	
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)	
<b>.ALPHA.-TOCOPHEROL ACETATE, D-</b> (UNII: A7E6112E4N)	
<b>CORN OIL</b> (UNII: 8470G57WFM)	
<b>AMMONIUM SULFATE</b> (UNII: SU46BAM238)	
<b>METHYLCHLOROISOTHIAZOLINONE</b> (UNII: DEL7T5QRPN)	
<b>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)	
<b>FD&amp;C RED NO. 4</b> (UNII: X3W0AM1JLX)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-580-89	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	09/09/2004	
2	NDC:21749-580-90	1250 mL in 1 PACKAGE; Type 0: Not a Combination Product	09/09/2004	
3	NDC:21749-580-43	1500 mL in 1 PACKAGE; Type 0: Not a Combination Product	09/09/2004	11/22/2024
4	NDC:21749-580-67	2000 mL in 1 PACKAGE; Type 0: Not a Combination Product	09/09/2004	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/09/2004	

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

## Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-580)

## Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		088312414	label(21749-580) , pack(21749-580)

Revised: 11/2024

GOJO Industries, Inc.