**TATTOO GOO- chloroxylenol liquid**
**The Magni Group 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: Chloroxylenol 0.5%**
**Purpose: Antibacterial**


NDC: 43689-0056-1

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**Use:**
**Helps eliminate germs on skin**

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**Warnings:**
**For external use only**

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**When using this product:**
**-avoid contact with eyes**

**-In case of eye contact, flush with water**

NDC: 43689-0056-1

**Stop use and ask a doctor if**
**irritation or redness develops.**

NDC: 43689-0056-1

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

NDC: 43689-0056-1

**Directions**
**-Three to four times daily for tattoos, one to two times daily for piercings.**
**-Squeeze into palm of hand.**
**-Work up lather in hands.**
**-Gently rub lather over tattoo or piercing.**
**-Cold water rinse, pat, then air dry.**

NDC: 43689-0056-1

**Inactive Ingredients:**
**Water (Aqua), Sodium PEG-7 Olive Oil Carboxylate, Propanediol, Olive Oil PEG-7 Esters, Sodium Laureth Sulfate, Polyquaternium 10, Cocoamide MEA, Cocamidopropyl Betaine, Diazolidinyl Urea, Sodium Benzoate, Potassium Sorbate, Citric Acid.**

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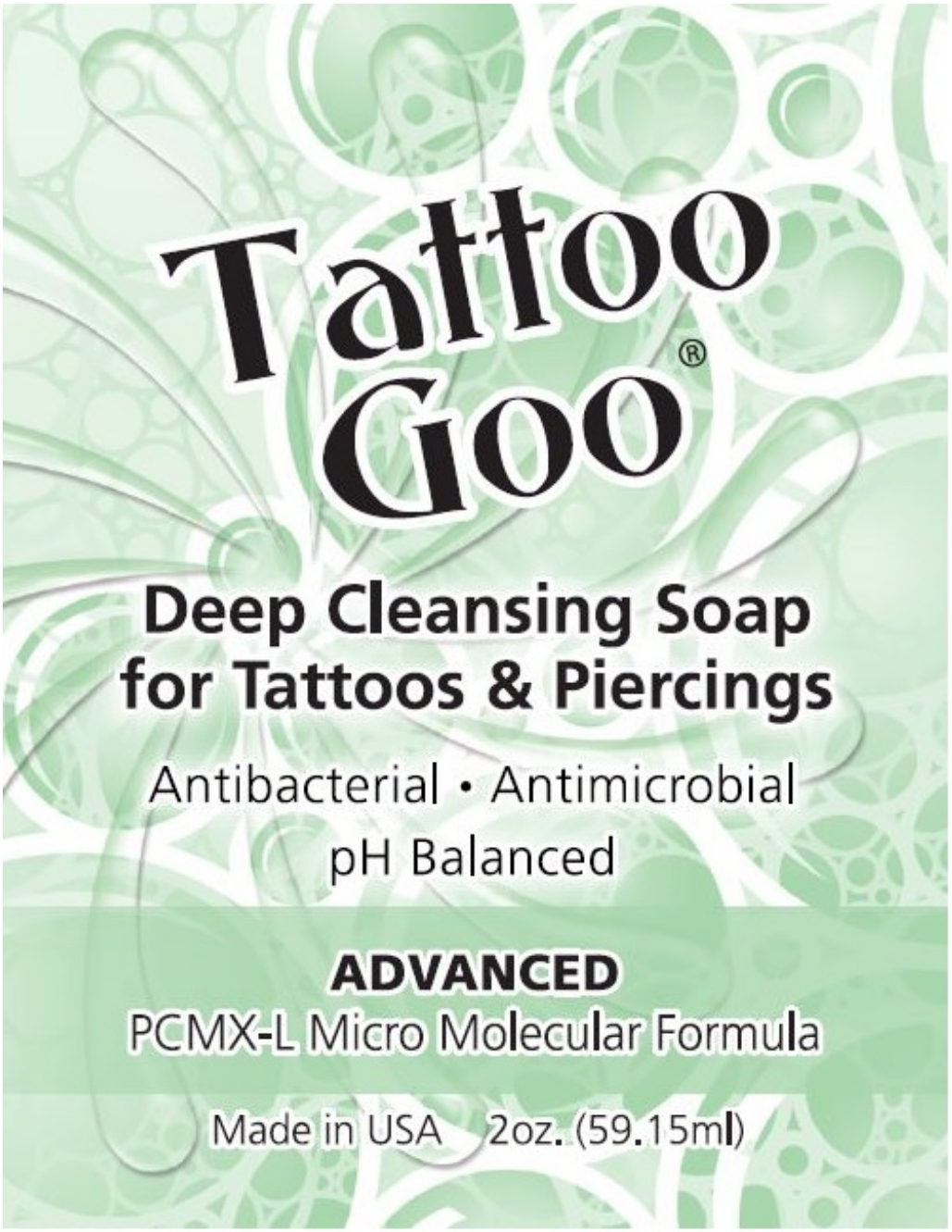
**Principal Display Panel**

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# TATTOO GOO

chloroxylenol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:43689-0056 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **CHLOROXYLENOL** (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q) | CHLOROXYLENOL | 0.5 mg in 100 mL |

| Inactive Ingredients | |
|---|---|
| Ingredient Name | Strength |
| **SODIUM LAURETH SULFATE** (UNII: BPV390UAP0) | |
| **ANHYDROUS CITRIC ACID** (UNII: XF417D3PSL) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **POLYQUATERNIUM-10 (1000 MPA.S AT 2%)** (UNII: GMR4PEN8PK) | |
| **COCO MONOETHANOLAMIDE** (UNII: C80684146D) | |
| **COCAMIDOPROPYL BETAINE** (UNII: 5OCF3O11KX) | |
| **POTASSIUM SORBATE** (UNII: 1VPU26JZZ4) | |
| **DIAZOLIDINYL UREA** (UNII: H5RIZ3MPW4) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **OLIVE OIL POLYGLYCERYL-6 ESTERS** (UNII: 4KDO9AFM9I) | |
| **PROPANEDIOL** (UNII: 5965N8W85T) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WJQ0SDW1A) | |

| Packaging | | | | |
|---|---|---|---|---|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| **1** | NDC:43689-0056-1 | 59.15 mL in 1 TUBE; Type 0: Not a Combination Product | 08/25/2020 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333E | 08/25/2020 | |

## Labeler - The Magni Group Inc (113501902)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| United Laboratories Manufacturing LLC | | 807878116 | manufacture(43689-0056) |

Revised: 8/2020 The Magni Group Inc