

# **PEARLIZED ANTIBACTERIAL SKIN CLEANSER- chloroxylenol liquid Fastenal**

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## **Pearlized Antibacterial Skin Cleanser**

### **Active Ingredient**

Chloroxylenol 0.40%

### **Uses**

- Antibacterial hand cleaner.
- Use in daycare, hospitals, nursing homes, physicians offices, dental offices and clinics

### **Warnings**

- **For external use only.**
- Avoid contact with eyes.
- Children under the age of 6 should be supervised by an adult when using this product.
- Discontinue use if irritation or redness develops.
- If irritation persists for more than 72 hours, consult a physician.
- **KEEP OUT OF REACH OF CHILDREN.**
- If swallowed, get medical help or contact a Poison Control Center right away.

### **Directions**

- **Read the entire label before using this product.**
- Dispense 2 pumps of product onto palm of hand and scrub thoroughly over all surfaces of both hands.
- Rinse with clean water.

### **Inactive Ingredients**

Water, Sodium Laureth Sulfate, Cocamidopropyl betadine, Phenoxyethanol, Coco MIPA, Glycol Stearate, DMDM Hydantoin, Propylene Glycol, Sodium Chloride, Fragrance, Glycerin, Ethanol, D&C Green #5, FD&C Yellow #5.

### **Purpose**

Antibacterial

Winning Hands Antibacterial

KEEP OUT OF REACH OF CHILDREN

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Drug Facts	
<b>Active Ingredient</b> Chloroxyleneol 0.4%	<b>Purpose</b> Antibacterials
<b>Uses</b>	
<ul style="list-style-type: none"> <li>Antibacterial hand cleanser.</li> <li>Use in daycare, hospitals, nursing homes, physician offices, dental offices and clinics.</li> </ul>	
<b>Warnings</b>	
For external use only.	
When using this product, avoid contact with eyes. If contact occurs, rinse thoroughly with water.	
Stop use and ask a doctor if irritation or rash occurs.	
<b>KEEP OUT OF THE REACH OF CHILDREN.</b>	
If swallowed, get medical help or contact a Poison Control Center right away.	
<b>Directions</b>	
<ul style="list-style-type: none"> <li>Read the entire label before using this product.</li> <li>Place enough product on your palm and scrub thoroughly over all surfaces of both hands.</li> <li>Rinse with clean water.</li> </ul>	
<b>Inactive Ingredients</b>	
Water, Sodium Lauryl Sulfate, Cocamidopropyl Betaine, Sodium Lauryl Sulfate, Ethylene Glycol Phenyl Ether, Cocamidopropylamine Oxide, Sodium Chloride, Propylene Glycol, Sodium Benzoate, Glycol Distearate, Fragrance, Laramide MPA, Sodium Hydroxide, Disodium Cocamido MPA-Sulfosuccinate, D&C Green 5, FD&C Yellow 5.	
<b>Questions? 807-454-5374</b>	

Datos del Producto	
<b>Ingrediente Activo</b> Cloroxileno 0.4%	<b>Propósito</b> Antibacterianos
<b>Usos</b>	
<ul style="list-style-type: none"> <li>Limpador de manos antibacteriano.</li> <li>Para usar en guarderías, hospitales, hogares para ancianos, consultorios médicos y odontológicos y clínicas.</li> </ul>	
<b>Advertencias</b>	
Solo para uso externo.	
Al utilizar este producto, evite el contacto con los ojos.	
En caso de contacto, enjuáguese los ojos con agua.	
Deje de usar y consulte a un médico si se produce irritación o sarpullido.	
<b>MANTENER FUERA DEL ALCANCE DE LOS NIÑOS.</b>	
En caso de ingestión, obtenga asistencia médica o diríjase a un centro de toxicología de inmediato.	
<b>Instrucciones</b>	
<ul style="list-style-type: none"> <li>Lea toda la etiqueta antes de usar este producto.</li> <li>Poner suficiente cantidad del producto en la palma y frótese bien en todas las superficies de ambas manos.</li> <li>Enjuague con agua limpia.</li> </ul>	
<b>Ingredientes Inactivos</b>	
Agua, sulfato de lauril de sodio, cocamidopropil betaina, sulfato de lauril de sodio, éter fenílico de etilenglicol, óxido de cocamidopropilamina, cloruro de sodio, propilenglicol, benzoato de sodio, distearato de glicol, fragancia, laramida MPA, hidróxido de sodio, sulfosuccinato disódico de cocamido MPA, D&C Green 5, FD&C Yellow 5.	
<b>Preguntas? 807-454-5374</b>	



Pearlized Antibacterial Skin Cleanser  
Limpador perlizado antibacteriano para la piel



NDC: 72024-841-04

Part # 923228098
1 GAL. (3.78 L)

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## PEARLIZED ANTIBACTERIAL SKIN CLEANSER

chloroxyleneol liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72024-841
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	4 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
.ALPHA.-AMYL CINNAMALDEHYDE (UNII: WC51CA3418)	
.BETA.-CITRONELLOL, (+/-)- (UNII: 565OK72VNF)	
3-(3,4-METHYLENEDIOXYPHENYL)-2-METHYLPROPANAL (UNII: L65EG8H6PA)	
4-ACETOXY-3-PENTYLTETRAHYDROPYRAN (UNII: 30E3255185)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
2-ISOBUTYL-4-METHYLTETRAHYDROPYRAN-4-OL (UNII: VK5ZHH2T3F)	
BENZYL SALICYLATE (UNII: WAO5MKN9TU)	
2-TERT-BUTYLCYCLOHEXYL ACETATE (UNII: 364FV60913)	
HEXYL SALICYLATE (UNII: 8F78EY72YL)	
.GAMMA.-UNDECALACTONE (UNII: QB1T0AG2YL)	
SODIUM CHLORIDE (UNII: 451W471Q8X)	
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
LAURIC ISOPROPANOLAMIDE (UNII: 82DUX3RRVU)	
SODIUM CARBONATE (UNII: 45P3261C7T)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	

<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)
<b>CALCIUM SILICATE</b> (UNII: S4255P4G5M)
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)
<b>LINALOOL, (+/-)-</b> (UNII: D81QY6I88E)
<b>TRIBASIC CALCIUM PHOSPHATE</b> (UNII: 91D9GV0Z28)
<b>SODIUM ALUMINIUM SILICATE</b> (UNII: 058TS43PSM)
<b>WATER</b> (UNII: 059QF0KO0R)
<b>SODIUM LAURETH SULFATE</b> (UNII: BPV390UAP0)
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)
<b>COCAMIDOPROPYLAMINE OXIDE</b> (UNII: M4SL82J7HK)
<b>BENZYL ACETATE</b> (UNII: 0ECG3V79ZJ)
<b>.ALPHA.-HEXYLCINNAMALDEHYDE</b> (UNII: 7X6O37OK2I)
<b>1-(2,3,8,8-TETRAMETHYL-1,2,3,4,5,6,7,8-OCTAHYDRONAPHTHALEN-2-YL)ETHANONE</b> (UNII: 1GD7ODM28Y)
<b>DIHYDROMYRCENOL</b> (UNII: 46L1B02ND9)
<b>HEXAHYDRO-4,7-METHANOINDEN-6-YL PROPIONATE</b> (UNII: S3ASM14UAS)
<b>VERDYL ACETATE</b> (UNII: 5232EN3X2F)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72024-841-04	3780 mL in 1 JUG; Type 0: Not a Combination Product	11/12/2012	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/12/2012	

**Labeler** - Fastenal (042653634)

**Registrant** - Betco corporation, Ltd. (005050158)

### Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd.		005050158	label(72024-841) , manufacture(72024-841)