PCMX WITH EMOLLIENT- chloroxylenol solution
Aplicare Products, LLC.

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.

PCMX with Emollient

3.3% w/w Chloroxylenol

Uses

For cleansing hands and forearms prior to surgery or patient care

Antiseptic

For external use only

Do not use in the eyes

Stop use and ask a doctor if

- skin shows symptoms of irritation, sensitivity, redness, pain or swelling
- discontinue use if irritation and redness develop. If condition persists more than 72 hours consult a doctor.

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

Directions

- Clean under nails with nail pick. Nails should be maintained with a 1 mm free edge
- Wet hands and forearms
- Apply 5 ml or palmful to hands and forearms
- Scrub thoroughly for 3 minutes with a wet brush
- Pay particular attention to the nails, cuticles and interdigital spaces
- Rinse thoroughly with running water
- Repeat

Other information

- Store at room temperature
- Avoid excessive heat above 40 deg C (104 deg F)
- Protect from freezing
- Latex Free

Citric Acid, PEG-120 Methyl Glucose Dioleate, Phenoxyethanol, Propylene Glycol, Purified Water, Sodium C12-15 Pareth-15 Sulfonate, Sodium Hydroxide, Sodium Lauroyl Sarcosinate, Sodium Laryl Sulfate, Soy Acid, Styrene/VP Copolymer

Questions?

Call: **1-800-523-0502** (Mon. - Fri. 8 AM-5 PM CST)

Scrub Care 3.3% Chloroxylenol

Emmollient CLeansing Solution

PCMX WITH EMOLLIENT

chloroxylenol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52380-9957
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	3.3 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM C12-15 PARETH-15 SULFONATE (UNII: 353VA59XH8)	
SOY ACID (UNII: HBA528N3PW)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
WATER (UNII: 059QF0KO0R)	
SODIUM LAUROYL SARCOSINATE (UNII: 632GS99618)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52380-9957-3	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/20/2018	
2	NDC:52380-9957-4	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/20/2018	
3	NDC:52380-9957-9	208198 mL in 1 DRUM; Type 0: Not a Combination Product	02/20/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2004	

Labeler - Aplicare Products, LLC. (081054904)

Establishment

Name	Address	ID/FEI	Business Operations
Aplicare Products, LLC.		081068252	manufacture(52380-9957)

Revised: 2/2018 Aplicare Products, LLC.