

ZEP APPLAUD AB- chloroxylenol liquid

Zep 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.

66949-385 Applaud AB

Chloroxylenol 0.3%

Antiseptic

Uses

Hand washing to decrease bacteria on skin.

For external use only.

- **For external use only.**

Do not use

Do not use in the eyes; if in eyes, rinse promptly and thoroughly with water.

When using this product

- Do not swallow.
- If swallowed, do not induce vomiting and if individual is conscious, give large quantities of water to drink and consult a physician immediately.

Stop use ask a doctor if skin irritation or redness persists for more than 72 hours.

Keep out of reach of children except under adult supervision.

Directions

- Wet hands with water.
- Place hands under dispenser.
- Apply liquid soap.
- Massage soap into hands and wrists, emphasizing back of hands, knuckles, and cuticles.
- Rinse thoroughly.

Other information

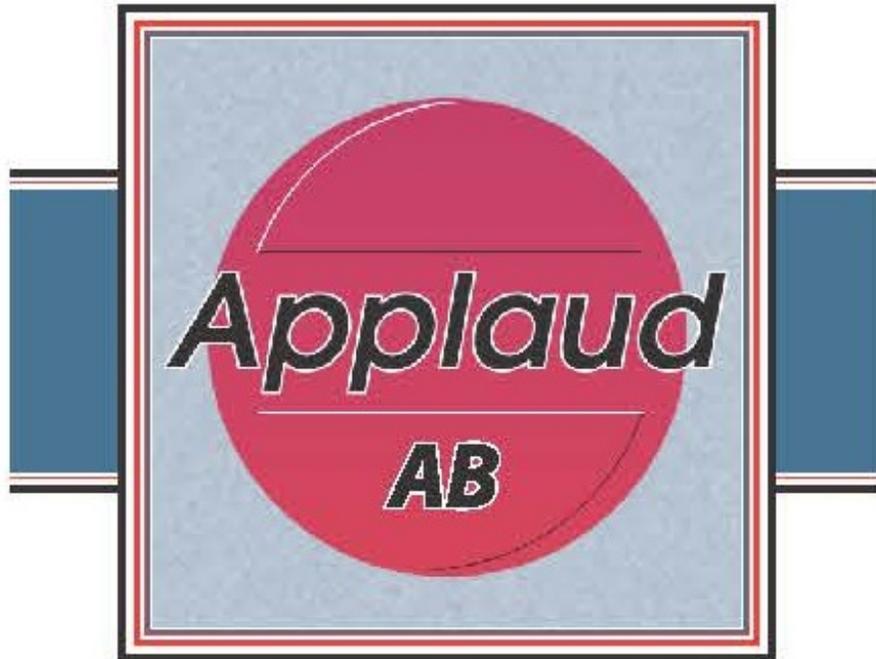
- Store at room temperature.
- Do not freeze.
- Dispose in accordance with all applicable federal, state and local regulations.

Inactive ingredients

Water, Potassium Tallate, Cocamide DIPA, Lauric Acid, Sodium Chloride, Tetrasodium EDTA, Acrylates/PEG-10 maleate/Styrene Copolymer, Isopropyl Alcohol, Fragrance, BHT, Red 4, Red 33

Questions or comments?

Call 1-877-BUY-ZEP (1-877-428-9937)



Lotion Hand Soap
With Anti-Bacterial Agent
Contains PCMX

ZEP APPLAUD AB

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66949-385
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)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
COCO DIISOPROPANOLAMIDE (UNII: S485AM948Q)	
POTASSIUM LAURATE (UNII: V4361R8N4Z)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHACRYLATE/METHOXY PEG-10 MALEATE/STYRENE COPOLYMER (UNII: 39DK5WQ2PR)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-385-01	11400 mL in 1 CONTAINER; Type 0: Not a Combination Product	03/10/2017	
2	NDC:66949-385-24	15140 mL in 1 CONTAINER; Type 0: Not a Combination Product	03/10/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/10/2017	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-385)

Revised: 12/2019

Zep Inc.