

**PRIME SOURCE E2 FOAM SO AP- chloroxylenol liquid
BUNZL**

PRIME SOURC® E2 Foam Soap

Active ingredient

Chloroxylenol 0.5%

Purpose

Antimicrobial

Use

- 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.

Inactive ingredients

Water (Aqua), Alcohol, Lauric Acid, Ethanolamine, Dipropylene Glycol, Lactic Acid, Poloxamer 124, Isopropyl Alcohol, Sodium Metabisulfite, Sodium Sulfite, Tetrasodium EDTA, Sodium Sulfate, Methylparaben, Propylparaben



Drug Facts	
Active ingredient Chloroxylenol 0.5%	Purpose Antimicrobial
Uses	
<ul style="list-style-type: none"> 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.	

Datos Farmacológicos	
Ingrediente activo Cloroxileno 0.5%	Propósito Antimicrobiano
Usos	
<ul style="list-style-type: none"> Lavado de manos empleado para disminuir la cantidad de bacterias en la piel Recomendado para uso reiterado 	
Advertencias	
Sólo para uso externo	
Al utilizar este producto, evitar el contacto con los ojos o con la zona alrededor de los ojos. En caso de contacto, enjuagar completamente los ojos con agua.	
Dejar de usar el producto y consultar a un médico si aparece y persiste una irritación o erupción cutánea	
Mantener fuera del alcance de los niños. En caso de ingestión, de inmediato acudir a un médico o ponerse en contacto con un centro para el control de tóxicos.	

Drug Facts (continued)	
Directions	
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Inactive ingredients	
Water (Aqua), Alcohol, Lauric Acid, Ethanolamine, Dipropylene Glycol, Lactic Acid, Poloxamer 124, Isopropyl Alcohol, Sodium Metabisulfite, Sodium Sulfite, Tetrasodium EDTA, Sodium Sulfate, Methylparaben, Propylparaben	
Questions or comments?	
Call 1-800-321-9647 Monday through Friday 8:00 AM to 5:00 PM	

Modo de uso	
<ul style="list-style-type: none"> Moje las manos Aplique el producto hasta cubrir completamente sus manos con la espuma Enjuague bien y seque las manos completamente 	
Ingredientes inactivos	
Agua, Alcohol etílico, Acido láurico, Etanolamina, Dipropilenglicol, Acido láctico, Poloxámero 124, Alcohol isopropílico, Metabisulfito de sodio, Sulfito de sodio, EDTA tetrasódico, Sulfato de sodio, Metilparabeno, Propilparabén	
¿Preguntas o comentarios?	
Llame al 1-800-321-9647 de lunes a viernes de 8:00 AM a 5:00 PM	

PRIME SOURCE E2 FOAM SOAP

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66294-523
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
WATER (UNII: 059QF0K00R)	
ALCOHOL (UNII: 3K9958V90M)	
LAURIC ACID (UNII: 1160N9NU9U)	
MONOETHANOLAMINE (UNII: 5KV86114PT)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
POLOXAMER 124 (UNII: 1S66E28KXA)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
SODIUM SULFITE (UNII: VTK01UQK3G)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66294-523-42	1250 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/15/2015	
2	NDC:66294-523-67	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/30/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/15/2015	

Labeler - BUNZL (799540588)

Revised: 12/2024

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