

**LYNX LX0864 PEACHY-FOAM ANTISEPTIC HANDSOAP- chloroxylenol soap
ABC Compounding Co., Inc.**

Lynx peachy foam 6864 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

Chloroxylenol 0.3%

Drug Facts Box OTC-Purpose Section

Antiseptic

Drug Facts Box OTC-Indications & Usage Section

for hand-washing to decrease bacteria on the skin

Drug Facts Box OTC-Warnings Section

For external use only

Drug Facts Box OTC-When Using Section

do not get into eyes

if contact occurs, rinse eyes thoroughly with water

Drug Facts Box OTC-Stop Use Section

irritation and redness develop

Drug Facts Box OTC-Keep Out of Reach of Children Section

if swallowed, get medical help or contact a Poison Control Center right away

Drug Facts Box OTC-Dosage & Administration Section

- wet hands and forearms
- apply 5 milliliters (teaspoonful) or palmful to hands and forearms
- scrub thoroughly for 1 minute and rinse

Drug Facts Box OTC-Inactive Ingredient Section

water, decyl glucoside, sodium laureth sulfate, cocamide MIPA, propylene glycol, sodium chloride, DMDM hydantoin, methylchloroisothiazolinone, methylisothiazolinone, fragrance, FD and C yellow no.5, food red 10

Lynx LX0864 peachy foam 6864

Lynx LX0864
peachy-foam

1000 ML
 (33.8 FL. OZ.)

**antiseptic
 hand soap**

CAUTION: KEEP OUT OF REACH OF CHILDREN

An effective antibacterial hand soap for use in food service facilities, hotels, athletic facilities, correctional facilities, business/office buildings, dressing rooms, factories, institutional facilities, locker rooms, nurseries, shower and bath areas, public facilities, public restrooms, schools, daycare centers, and campgrounds. Formulated with skin conditioners for extra mildness.

Drug Facts	
Active Ingredient	Purpose
Chloroxylenol 0.3%	Antiseptic
Uses for hand-washing to decrease bacteria on the skin	
Warnings	
For external use only	
When using this product	
<ul style="list-style-type: none"> ■ do not get into eyes ■ if contact occurs, rinse eyes thoroughly with water 	
Stop use and ask a doctor if	
<ul style="list-style-type: none"> ■ irritation and redness develop 	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away	
Directions	
<ul style="list-style-type: none"> ■ wet hands and forearms ■ apply 5 milliliters (teaspoonful) or palmful to hands and forearms ■ scrub thoroughly for 1 minute and rinse 	
Inactive Ingredients water, decyl glucoside, sodium laureth sulfate, cocamide MIPA, propylene glycol, DMDM hydantoin, fragrance, methylchloroisothiazolinone, methylisothiazolinone, aloe barbadensis, FD&C yellow no. 5, food red 10	

This product is designed exclusively for industrial and institutional use by trained personnel. This product is sold 'as is' and the manufacturer makes no warranty, expressed or implied, of merchantability, and/or fitness for a particular purpose or otherwise.

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Manufactured by: ABC COMPOUNDING CO., INC.
 P.O. Box 16247 • Atlanta, GA 30321



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Lynx peachy foam 6864

LYNX LX0864 PEACHY-FOAM ANTISEPTIC HANDSOAP

chloroxylenol soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62257-686
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
COCO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ACID RED 1 (UNII: 3365R6427R)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62257-686-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	04/27/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	04/26/2022	

Labeler - ABC Compounding Co., Inc. (003284353)**Registrant** - ABC Compounding Co., Inc. (003284353)**Establishment**

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(62257-686)