

LYNX ANTIBACTERIAL - chloroxylenol soap
ABC Compounding Co., 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.

Lynx Antibacterial Hand Soap 6544 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 forearm
- apply 5 milliliters (teaspoonful) or palmful to hands and forearm
- 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, citric acid, DMDM hydantoin, fragrance, FD and C yellow no. 5, food red 10

Lynx Antibacterial Hand Soap 6544 1 gallon

1606

Lynx

antibacterial hand soap

■ ANTISEPTIC HAND WASH

CAUTION: KEEP OUT OF REACH OF CHILDREN

Si usted no entiende la etiqueta, busque a alguien para que se la explique a usted en detalle.
(If you do not understand this label, find someone to explain it to you in detail.)

NET CONTENTS: 1 GALLON (3.8 L)



This product is an effective antibacterial hand soap for use in food processing plants and restaurants by personnel prior to handling food and/or food processing equipment.

This formulation effectively reduces the bacterial flora of the skin. When tested via the Time Kill Test, it demonstrates 99% or greater kill against *Staphylococcus aureus*, *E. Coli*, and *Pseudomonas aeruginosa*. Formulated with skin conditioners for extra mildness.

Drug Facts

Active Ingredient	Purpose
Chloroxylenol 0.3%	Antiseptic

Uses
For handwashing to decrease bacteria on the skin.

Warnings

For external use only

When using this product

- do not get into eyes
- if contact occurs, rinse eyes thoroughly with water

- Stop use and ask a doctor if:
 - irritation and redness develop

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

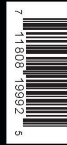
Directions
<ul style="list-style-type: none"> wash hands and forearms apply 5 milliliters (teaspoonful) or equivalent to hands and forearms scrub thoroughly for 1 minute and rinse

Inactive Ingredients
water, deyl glucose, sodium lauryl sulfate, cocamide MIPA, propylene glycol, DMDM hydantoin, sodium chloride, citric acid, fragrance, 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, express or implied, of merchantability, fitness for a particular purpose or otherwise.

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

Lynx is a registered trademark
of ABC Compounding Co., Inc.
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L1166.121018



Lynx Antibacterial Hand Soap Label

LYNX ANTIBACTERIAL

chloroxylenol soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62257-160
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)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CO CO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62257-160-06	1 in 1 BOX	07/14/2006	
1		800 mL in 1 BAG; Type 0: Not a Combination Product		

2	NDC:62257-160-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
3	NDC:62257-160-24	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
4	NDC:62257-160-01	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/14/2006	
5	NDC:62257-160-03	350 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/14/2006	
6	NDC:62257-160-05	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
7	NDC:62257-160-07	700 mL in 1 BAG; Type 0: Not a Combination Product	07/14/2006	
8	NDC:62257-160-09	2000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/14/2006	
9	NDC:62257-160-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/14/2006	
10	NDC:62257-160-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
11	NDC:62257-160-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	07/14/2006	
12	NDC:62257-160-13	800 mL in 1 BAG; Type 0: Not a Combination Product	07/14/2006	
13	NDC:62257-160-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
14	NDC:62257-160-15	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
15	NDC:62257-160-28	149 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
16	NDC:62257-160-27	800 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/14/2006	
17	NDC:62257-160-55	208200 mL in 1 DRUM; Type 0: Not a Combination Product	07/14/2006	
18	NDC:62257-160-08	1 in 1 BOX	07/14/2006	
18		1000 mL in 1 BAG; Type 0: Not a Combination Product		
19	NDC:62257-160-16	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
20	NDC:62257-160-18	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/14/2006	
21	NDC:62257-160-19	18900 mL in 1 CONTAINER; Type 0: Not a Combination Product	07/14/2006	
22	NDC:62257-160-20	75600 mL in 1 DRUM; Type 0: Not a Combination Product	07/14/2006	
23	NDC:62257-160-35	132500 mL in 1 DRUM; Type 0: Not a Combination Product	07/14/2006	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/14/2006	

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(62257-160)

Revised: 12/2018

ABC Compounding Co., Inc.