DRUMMOND AMERICAN TRADE WIND ANTIMICROBIAL- chloroxylenol liquid
Lawson Products, 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.

DRUMMOND AMERICAN TRADE WIND Antimicrobial Lotion Soap

Drug Facts

Active Ingredient

Chloroxylenol (1% v/v)

Purpose

Antiseptic Hand Sanitizer

Uses

- Helps reduce bacteria that potentially can cause disease
- Helps prevent cross contamination by hand contact

Warnings

- **For external use only**
- **Do not use near eyes**
- **Keep out of reach of children.**
- In case of eye contact flush with water for 15 minutes
- If irritation persists, get medical attention
- In case of accidental ingestion, seek medical attention or contact a poison control center immediately.

Directions

- Wet hands and forearms
- Apply appropriate amount to palm and hand
- Scrub hands and forearms for 60 seconds
- Rinse well
- Wipe dry with towel, repeat if necessary.

Other Information

- Assists with OSHA bloodborne pathogen standard compliance

Inactive Ingredients

Water, Cocoate, Oleate, Cocamidopropyl betaine, Hydroxyethyl Cellulose, Polyquaternium 7, Na_4EDTA, 2-Propanol, Peg-75 Lanolin, Aloe Vera Gel, Fragrance, Triclosan, DMDM Hydantoin.

Principal Display Panel – Bottle Label

DRUMMOND AMERICAN

TRADE WIND

Antimicrobial Lotion Soap

KEEP OUT OF REACH OF CHILDREN

SEE DRUG FACTS PANEL FOR ADDITIONAL INFORMATION.

Drug Facts

Active Ingredient: Chloroxylenol (1.0% w/w)

Purpose: Antiseptic Hand Cleanser

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Warning

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For Industrial And Institutional Use Only.

505-002-024-804

DL3070

DL3070

Sold exclusively by

DRUMMOND AMERICAN

Drummond American Corporation
600 Corporate Woods Pkwy.
Vernon Hills, IL 60061-3165 (847) 913-9313

DRUMMOND AMERICAN

TRADE WIND

Antimicrobial Lotion Soap

KEEP OUT OF REACH OF CHILDREN

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NET CONTENTS 1 U.S. GAL. (3.78 L)

DRUMMOND AMERICAN TRADE WIND ANTIMICROBIAL

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62428-505
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
chloroxylenol (UNII: 0F32U78V2Q) (chloroxylenol - UNII:0F32U78V2Q)	chloroxylenol	10.22 g in 1000 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
potassium oleate (UNII: 74WHF607EU)	
cocamidopropyl betaine (UNII: 5OCF3O11KX)	

hydroxyethyl cellulose (2000 CPS at 1%) (UNII: S38J6RZN16)	
isopropyl alcohol (UNII: ND2M416302)	
edetate sodium (UNII: MP1J8420LU)	
aloe vera leaf (UNII: ZY81Z83H0X)	
DMDM hydantoin (UNII: BYR0546TOW)	
triclosan (UNII: 4NM5039Y5X)	
potassium cocoate (UNII: F8U72V8ZXP)	
polyquaternium-7 (70/30 acrylamide/DADMAC; 1600 kd) (UNII: 0L414VCS5Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62428-505-55	3785 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333	10/28/1998	

Labeler - Lawson Products, Inc. (005438890)

Establishment

Name	Address	ID/FEI	Business Operations
Canberra Corporation		068080621	MANUFACTURE

Revised: 8/2010 Lawson Products, Inc.