DELUXE ALL-PURPOSE - triclosan liquid
Johnson Labs, 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.

Active Ingredient

Drug Facts Active Ingredients Triclosan 0.2 %

Purpose

Purpose Handwash

Keep Out of Reach of Children

Keep out of reach of children - If swallowed get medical help or contact a Poison Control Center immediately.

Uses

Antimicrobial handwash.

Warnings

Warnings - For external use only.

When using this product - Keep out of eyes. In case of contact with eyes, flush thoroughly with water.

Directions

Directions - Dispense a small amount of soap into hands or on a washcloth. Work up a lather and rinse with water.

Inactive Ingredients

Inactive Ingredients - As-40, Cd-6, Color, Dantogard, Propylene Glycol, Water.

Package Label

Johnson Labs DELUXE ALL-PURPOSE LIQUID HAND SOAP

This state-of-the-art hand soap was developed to be used in industries, schools, offices or wherever people need a high quality hand soap. This product can also be used as a hair and body shampoo and clothes wash.

1 Gallon (3.78L) made in U.S.A. Johnson Labs, Inc. Troy, AL 36081 334-566-9152 800-473-9152

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DRUG FACTS

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Troy, AL 36081
334-566-9152
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DELUXE ALL-PURPOSE
triclosan liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59854-301
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	7.57 mL in 3.78 L

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59854-301-01	3.78 L in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2000	

Labeler - Johnson Labs, Inc. (805806742)

Registrant - Johnson Labs, Inc. (805806742)

Establishment

Name	Address	ID/FEI	Business Operations
Johnson Labs, Inc.		805806742	manufacture

Revised: 4/2012 Johnson Labs, Inc.