POWERHOUSE ULTRA DISH ANTIBACTERIAL- chloroxylenol liquid
Delta Brands 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.

Drug Facts

Active ingredient

Chloroxylenol 0.30%

Purpose

Antibacterial

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only: hands 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 develops ■ condition persists for more than 72 hours

Keep out of reach of children

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

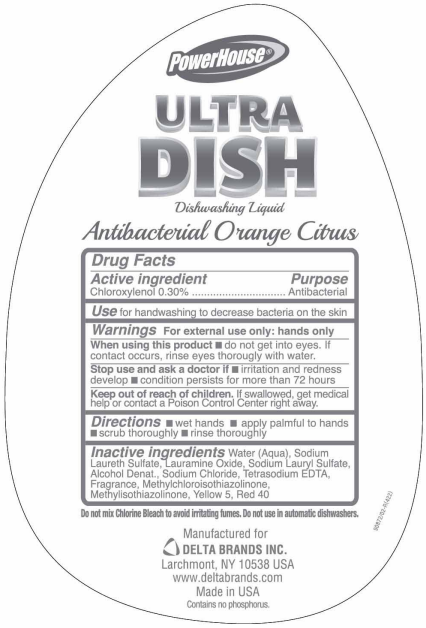
Directions

■ wet hands ■ apply palmful to hands ■ scrub thoroughly ■ rinse thoroughly

Inactive ingredients

Water (Aqua), Sodium Laureth Sulfate, Lauramine Oxide,Sodium Lauryl Sulfate, Alcohol Denat. Sodium Chloride, Tetrasodium EDTA, Fragrance, Methylchloroisothiazolinone, Methylisothiazolinone, Yellow 5, Red 40

Package Label



POWERHOUSE ULTRA DISH ANTIBACTERIAL

chloroxylenol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:20276-153
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.3 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
ALCOHOL (UNII: 3K9958V90M)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	

EDETATE SODIUM TETRAHYDRATE (UNII: L13NHD21X6)	
WATER (UNII: 059QF0KO0R)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:20276-153-25	740 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/01/2019	

Labeler - Delta Brands Inc (102672008)

Revised: 2/2019 Delta Brands Inc