

**MANGO SELECT HIGH FOAMING ANTISEPTIC HAND AND BODY WASH 6876-
chloroxylenol soap**

Warsaw Chemical Holding, LLC

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.

Mango Select High Foaming Antiseptic Hand and Body Wash 6876

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

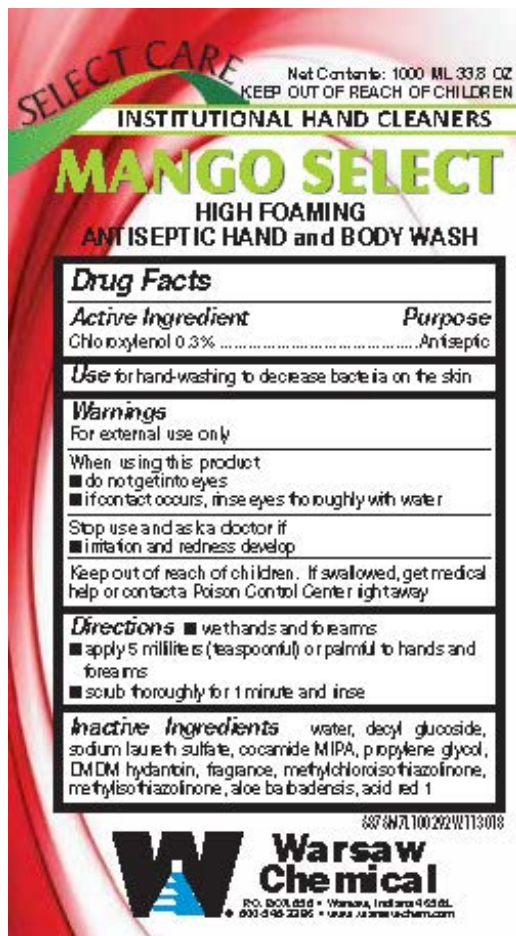
Drug Facts Box OTC-Dosage & Administration Section

- 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, DMDM hydantoin, fragrance, methylchloroisothiazolinone, methylisothiazolinone, aloe barbadensis, acid red 1

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72379-876
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)	
ACID RED 1 (UNII: 3365R6427R)	
METHYLCHLOROISO THIAZOLINONE (UNII: DEL7T5QRPN)	

METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72379-876-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	06/18/2018	
2	NDC:72379-876-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2018	
3	NDC:72379-876-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	06/18/2018	
4	NDC:72379-876-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2018	
5	NDC:72379-876-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/18/2018	

Labeler - Warsaw Chemical Holding, LLC (081244708)

Registrant - ABC Compounding Co., Inc. (003284353)

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(72379-876)

Revised: 12/2018

Warsaw Chemical Holding, LLC