

**SANI MANGO ANTISEPTIC HAND AND BODY WASH- chloroxylenol soap
ABC Compounding Co., Inc.**

Sani Mango Antiseptic Hand and Body Wash 6875 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 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, sodium chloride, methylchloroisothiazolinone, methylisothiazolinone, DMDM hydantoin, fragrance, aloe barbadensis, acid red 1

Sani Mango Antiseptic Hand and Body Wash 6875

SANI MANGO ANTISEPTIC HAND AND BODY WASH

chloroxylenol soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62257-875
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)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ACID RED 1 (UNII: 3365R6427R)	
METHYLCHLOROISOTHIAZOLINONE (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:62257-875-06	1 in 1 BOX	01/04/2017	
1		800 mL in 1 BAG; Type 0: Not a Combination Product		
2	NDC:62257-875-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	
3	NDC:62257-875-24	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	11/29/2025
4	NDC:62257-875-01	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/04/2017	11/29/2025
5	NDC:62257-875-03	350 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/04/2017	11/29/2025
6	NDC:62257-875-05	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	11/29/2025
7	NDC:62257-875-07	700 mL in 1 BAG; Type 0: Not a Combination Product	01/04/2017	11/29/2025
8	NDC:62257-875-09	2000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/04/2017	11/29/2025
9	NDC:62257-875-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/04/2017	11/29/2025

10	NDC:62257-875-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	11/29/2025
11	NDC:62257-875-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	01/04/2017	11/29/2025
12	NDC:62257-875-13	800 mL in 1 BAG; Type 0: Not a Combination Product	01/04/2017	11/29/2025
13	NDC:62257-875-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	
14	NDC:62257-875-15	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	
15	NDC:62257-875-28	149 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	11/29/2025
16	NDC:62257-875-27	800 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/04/2017	11/29/2025
17	NDC:62257-875-55	208200 mL in 1 DRUM; Type 0: Not a Combination Product	01/04/2017	11/29/2025
18	NDC:62257-875-08	1 in 1 BOX	01/04/2017	11/29/2025
18		1000 mL in 1 BAG; Type 0: Not a Combination Product		
19	NDC:62257-875-16	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	11/29/2025
20	NDC:62257-875-18	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2017	11/29/2025
21	NDC:62257-875-19	18900 mL in 1 CONTAINER; Type 0: Not a Combination Product	01/04/2017	11/29/2025
22	NDC:62257-875-20	75600 mL in 1 DRUM; Type 0: Not a Combination Product	01/04/2017	11/29/2025
23	NDC:62257-875-35	132500 mL in 1 DRUM; Type 0: Not a Combination Product	01/04/2017	11/29/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/04/2017	

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

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(62257-875)

Revised: 12/2025

ABC Compounding Co., Inc.