

**SUDS HIGH FOAMING ANTISEPTIC HAND AND BODY WASH-
chloroxylenol soap
Pro chem, Inc.**

Suds High Foaming Antiseptic Hand and Body Wash 6876 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, DMDM

hydantoin, fragrance, methylchloroiso-thiazolinone, methylisothiazolinone, aloe barbadensis, acid red 1

Suds High Foaming Antiseptic Hand & Body Wash 6876



An effective antibacterial hand soap for use in hotels, athletic facilities, correctional facilities, business/office buildings, dressing rooms, factories, institutional facilities, locker rooms, nurseries, shower and bath areas, public facilities, public rest rooms, schools, day care centers, and camp grounds.

This formulation effectively reduces the bacterial flora of the skin. When tested via the Time Kill Test, it demonstrates 99% or greater kill against *Staphylococcus aureus*, *E. Coli*, and *Pseudomonas aeruginosa*. Formulated with skin conditioners for extra mildness.



Contains ALOE VERA!

Suds Hand SOAP

High Foaming
Antiseptic Hand and Body Wash

NET CONTENTS: 18 OZ. (532 ml)

Drug Facts

Active Ingredient	Purpose
Chloroxylenol 0.3%	Antiseptic

Uses for hand-washing to decrease bacteria on the skin

Warnings
For external use 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 develop

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions ■ wet hands and forearms
■ apply 5 milliliters (teaspoonful) or palmful to hands and forearms ■ scrub thoroughly for 1 minute and rinse

Inactive Ingredients water, decyl glucoside, sodium laureth sulfate, cocamide MIPA, propylene glycol, DMDM hydantoin, fragrance, methylchloroiso-thiazolinone, methylisothiazolinone, aloe barbadensis, acid red 1

Product Code: 2182




Made In U.S.A.

Sold by PRO CHEM, INC.
1475 BLUEGRASS LAKES PARKWAY
ALPHARETTA, GA 30004
TO REORDER CALL: 1-800-241-8180
www.procheminc.com

BATCH NO.: 687618P994.121018

Suds High Foaming Antiseptic Hand & Body Wash

SUDS HIGH FOAMING ANTISEPTIC HAND AND BODY WASH			
chloroxylenol soap			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63830-876
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)			
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:63830-876-06	1 in 1 BOX	01/10/2017	11/20/2025
1		800 mL in 1 BAG; Type 0: Not a Combination Product		
2	NDC:63830-876-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	
3	NDC:63830-876-24	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	11/20/2025
4	NDC:63830-876-01	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/10/2017	11/20/2025
5	NDC:63830-876-03	350 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/10/2017	11/20/2025
6	NDC:63830-876-05	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	11/20/2025
7	NDC:63830-876-07	700 mL in 1 BAG; Type 0: Not a Combination Product	01/10/2017	11/20/2025
8	NDC:63830-876-09	2000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/10/2017	11/20/2025
9	NDC:63830-876-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/10/2017	
10	NDC:63830-876-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	11/20/2025
11	NDC:63830-876-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	01/10/2017	11/20/2025
12	NDC:63830-876-13	800 mL in 1 BAG; Type 0: Not a Combination Product	01/10/2017	11/20/2025
13	NDC:63830-876-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	
14	NDC:63830-876-15	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	11/20/2025
15	NDC:63830-876-28	149 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	11/20/2025
16	NDC:63830-876-27	800 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/10/2017	11/20/2025
17	NDC:63830-876-55	208200 mL in 1 DRUM; Type 0: Not a Combination Product	01/10/2017	11/20/2025
18	NDC:63830-876-08	1 in 1 BOX	01/10/2017	11/20/2025
18		1000 mL in 1 BAG; Type 0: Not a Combination Product		
19	NDC:63830-876-16	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	11/20/2025
20	NDC:63830-876-18	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	11/20/2025
21	NDC:63830-876-19	18900 mL in 1 CONTAINER; Type 0: Not a Combination Product	01/10/2017	11/20/2025
22	NDC:63830-876-20	75600 mL in 1 DRUM; Type 0: Not a Combination Product	01/10/2017	11/20/2025
23	NDC:63830-876-35	132500 mL in 1 DRUM; Type 0: Not a Combination Product	01/10/2017	11/20/2025

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	M003	01/10/2017	
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Labeler - Pro chem, Inc. (061396065)

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

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(63830-876)

Revised: 12/2025

Pro chem, Inc.