INHIBIT II HIGH FOAMING ANTISEPTIC HAND CLEANER- chloroxylenol soap
Select Specialty Products

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.

Inhibit II High Foaming Antiseptic Hand Cleaner 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, methylchloroisothiazolinone, methylisothiazolinone, aloe barbadensis, acid red 1

Inhibit II High Foaming Antiseptic Hand Cleaner

INHIBIT II HIGH FOAMING ANTISEPTIC HAND CLEANER

chloroxylenol soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67996-876
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:67996-876-06	1 in 1 BOX	01/12/2017	
1		800 mL in 1 BAG; Type 0: Not a Combination Product		
2	NDC:67996-876-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	
3	NDC:67996-876-24	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	
4	NDC:67996-876-01	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/12/2017	
5	NDC:67996-876-03	350 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/12/2017	
6	NDC:67996-876-05	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	
7	NDC:67996-876-07	700 mL in 1 BAG; Type 0: Not a Combination Product	01/12/2017	
8	NDC:67996-876-09	2000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/12/2017	
9	NDC:67996-876-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/12/2017	
10	NDC:67996-876-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	

11	NDC:67996-876-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	01/12/2017	
12	NDC:67996-876-13	800 mL in 1 BAG; Type 0: Not a Combination Product	01/12/2017	
13	NDC:67996-876-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	
14	NDC:67996-876-15	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	
15	NDC:67996-876-28	149 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	
16	NDC:67996-876-27	800 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/12/2017	
17	NDC:67996-876-55	208200 mL in 1 DRUM; Type 0: Not a Combination Product	01/12/2017	
18	NDC:67996-876-08	1 in 1 BOX	01/12/2017	
18		1000 mL in 1 BAG; Type 0: Not a Combination Product		
19	NDC:67996-876-16	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	
20	NDC:67996-876-18	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/12/2017	
21	NDC:67996-876-19	18900 mL in 1 CONTAINER; Type 0: Not a Combination Product	01/12/2017	
22	NDC:67996-876-20	75600 mL in 1 DRUM; Type 0: Not a Combination Product	01/12/2017	
23	NDC:67996-876-35	132500 mL in 1 DRUM; Type 0: Not a Combination Product	01/12/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/12/2017	

Labeler - Select Specialty Products (121819445)

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

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(67996-876)

Revised: 12/2018 Select Specialty Products