CELOX ULTRA - alcohol liquid
Certus Medical, 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.

Celox Ultra 6968 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

Ethyl Alcohol 62% v/v

Drug Facts Box OTC-Purpose Section

Antiseptic

Drug Facts Box OTC-Indications & Usage Section

for hand-washing to decrease bacteria on the skin

if hands are visibly soiled, use regular soap and water

recommended for repeated use

Drug Facts Box OTC-Warnings Section

FLAMMABLE, keep away from fire and flames

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

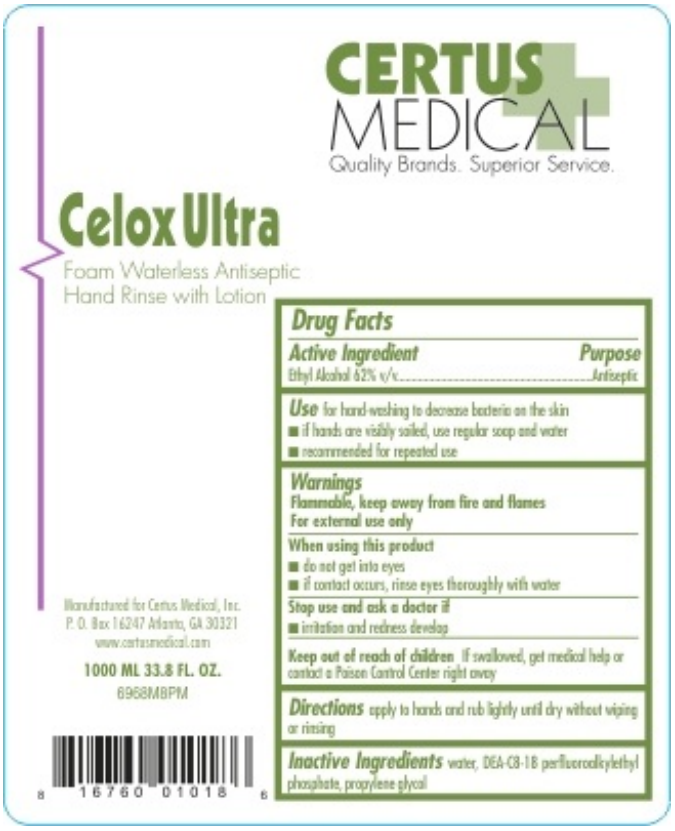
Drug Facts Box OTC-Dosage & Administration Section

apply to hands and rub lightly until dry without wiping or rinsing

Drug Facts Box OTC-Inactive Ingredient Section

water, DEA-C8-18 perfluoroalkylethyl phosphate, propylene glycol

Celox Ultra 6968 1000ml bag



6968M8PM.jpg Celox Ultra 1000 ml

CELOX ULTRA

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75990-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIETHANOLAMINE BIS(C8-C18 PERFLUOROALKYLETHYL)PHOSPHATE (UNII: 4J55VM509S)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:75990-007-06	1 in 1 BOX		
1		800 mL in 1 BAG		
2	NDC:75990-007-30	473 mL in 1 BOTTLE, PLASTIC		

3	NDC:75990-007-24	118 mL in 1 BOTTLE, PLASTIC		
4	NDC:75990-007-01	1200 mL in 1 CARTRIDGE		
5	NDC:75990-007-03	350 mL in 1 CARTRIDGE		
6	NDC:75990-007-05	540 mL in 1 BOTTLE, PLASTIC		
7	NDC:75990-007-07	700 mL in 1 BAG		
8	NDC:75990-007-09	2000 mL in 1 CARTRIDGE		
9	NDC:75990-007-10	1000 mL in 1 CARTRIDGE		
10	NDC:75990-007-11	1000 mL in 1 BOTTLE, PLASTIC		
11	NDC:75990-007-12	1000 mL in 1 BAG		
12	NDC:75990-007-13	800 mL in 1 BAG		
13	NDC:75990-007-14	3785 mL in 1 BOTTLE, PLASTIC		
14	NDC:75990-007-15	946 mL in 1 BOTTLE, PLASTIC		
15	NDC:75990-007-28	149 mL in 1 BOTTLE, PLASTIC		
16	NDC:75990-007-27	800 mL in 1 CARTRIDGE		
17	NDC:75990-007-55	208200 mL in 1 DRUM		
18	NDC:75990-007-08	1 in 1 BOX		
18		1000 mL in 1 BAG		
19	NDC:75990-007-16	236 mL in 1 BOTTLE, PLASTIC		
20	NDC:75990-007-18	50 mL in 1 BOTTLE, PLASTIC		
21	NDC:75990-007-17	532 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/25/2011	

Labeler - Certus Medical, Inc. (966433653)

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

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture

Revised: 4/2011 Certus Medical, Inc.