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

	CERTUS MEDICAL Guality Brands. Superior Service.
Celox Ultra	a
Foam Waterless Antise Hand Rinse with Lotion	
	Drug Facts Active Ingredient Purpose thyl Acabal 62% s/s.
	Use for hand-washing to decrease bacteria on the skin if hands are visibly salled, use regular scop and water recommended for repeated use
	Warnings Flammable, keep away from fire and flames For external use only
	When using this product do not get into eyes it contact occurs, rinse eyes thoroughly with water
Honofactured for Certus Medical, Inc. P. D. Box 16247 Atlanto, GA 30321 www.certastnetical.com	Stop use and ask a doctor if initiation and redness develop
1000 ML 33.8 FL. OZ.	Keep out of reach of children If swallowed, get medical help or cantact a Paison Cantrol Center right away
GEOGMBHM	Directions apply to hands and rub lightly until dry without viging or rinsing
	Inactive Ingredients water, DEA-C3-18 perfluercelkylethyl phosphate, propylene glycal

6968M8PM.jpg Celox Ultra 1000 ml

CI	ELOX ULTRA						
lc	ohol liquid						
P	roduct Information	ı					
Pr	oduct T ype		HUMAN OTC DRUG	Item Code (Source)	NDC:7599	0-007
Ro	oute of Administration	1	TOPICAL				
	···· T 1						
A	ctive Ingredient/A		-			-	
		Ingredient Name Basis of Strength			Strength		
AL	COHOL (UNII: 3K9958	V90M) (ALC	COHOL - UNII:3K9958V90M)		ALCOHOL	0.7	mL in 1 mL
In	active Ingredients	,					
			Ingredient Name				Strength
			0				
W	ATER (UNII: 059QF0KO	0 R)	5				
PR	OPYLENE GLYCOL (U	JNII: 6DC9Q	167V3)				
PR	OPYLENE GLYCOL (U	JNII: 6DC9Q	J. J	SPHATE (UNI	:4J55VM509S)		
PR	OPYLENE GLYCOL (U	JNII: 6DC9Q	167V3)	SPHATE (UNI	:4J55VM509S)		
PR	OPYLENE GLYCOL (U	JNII: 6DC9Q	167V3)	SPHATE (UNI	:4J55VM509S)		
PR DI	OPYLENE GLYCOL (U	JNII: 6DC9Q	167V3)	SPHATE (UNI	:4J55VM509S)		
PR DI	OPYLENE GLYCOL (U ETHANOLAMINE BIS(JNII: 6DC9Q C 8-C18 PER	167V3)			Marketin	g End Date
PR DI Pa	OPYLENE GLYCOL (U ETHANOLAMINE BIS(O Ickaging	JNII: 6DC9Q C 8-C18 PER	167V3) FLUOROALKYLETHYL)PHC ackage Description			Marketin	
PR DII Pa #	OPYLENE GLYCOL (U ETHANOLAMINE BIS(ckaging Item Code	JNII: 6DC9Q C8-C18 PER	167V3) FLUOROALKYLETHYL)PHC ackage Description X			Marketin	

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		
Μ	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.