RECENS- alcohol gel
Certus Medical, Inc.

Recens Gel 6966 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

Ethyl Alcohol 62%

Drug Facts Box OTC-Purpose Section

Antiseptic

Drug Facts Box OTC-Indications & Usage Section

for hand-washing to decrease bacteria on the skin, only when water is not available

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

wet hands thoroughly with product and allow to dry without wiping

Drug Facts Box OTC-Inactive Ingredient Section

water, diisopropylamine, carbomer, propylene glycol, DMDM hydantoin, tocopheryl acetate, aloe barbadensis

Recens Gel 6966 1000ml

RECENS

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75990-605
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)	
CARBOMER 934 (UNII: Z135WT9208)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75990-605-24	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/25/2020	
2	NDC:75990-605-01	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	06/25/2020	
3	NDC:75990-605-03	350 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	06/25/2020	
4	NDC:75990-605-05	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/25/2020	
5	NDC:75990-605-07	700 mL in 1 BAG; Type 0: Not a Combination Product	06/25/2020	
6	NDC:75990-605-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	06/25/2020	
7	NDC:75990-605-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/25/2020	
8	NDC:75990-605-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	06/25/2020	
9	NDC:75990-605-13	800 mL in 1 BAG; Type 0: Not a Combination Product	06/25/2020	
10	NDC:75990-605-28	149 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/25/2020	
11	NDC:75990-605-27	800 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	06/25/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	06/25/2020	

Labeler - Certus Medical, Inc. (118806847)

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

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(75990-605)

Revised: 1/2025 Certus Medical, Inc.