

ALCO SAN- alcohol gel
Mid American Research Chemical Corp

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.

Alco San Instant Hand Sanitizer 6605 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

Alco San 6605 5oz

MARCO 

ALCO-SAN

Instant Hand Sanitizer

(GEL-TYPE) NDC 64233-221-28

This instant hand sanitizer is an ethyl alcohol hand sanitizer for use when water is unavailable.

- Kills 99.99% of E. coli, Salmonella enterica and Staphylococcus aureus (MRSA) in 15 seconds

Contains Vitamin E and Aloe Vera!

DANGER: FLAMMABLE
 KEEP OUT OF REACH OF CHILDREN.
 KEEP AWAY FROM FIRE OR FLAME.
 FOR EXTERNAL USE ONLY.
 See other cautions on opposite panel of label.

Net Contents: 5 oz. (149 ml) M-417

Drug Facts

Active Ingredient	Purpose
Ethyl Alcohol 62%	Antiseptic

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

Warnings

Flammable, keep away from fire and flames
 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 thoroughly with product and allow to dry without wiping

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

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P0515

Manufactured in USA For:
 Mid-American Research Chemical Corp.
 P. O. Box 927, Columbus, NE 68602-0927 | 800-228-8508

ALCO SAN

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64233-221
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: 059QF0K00R)	
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:64233-221-06	1 in 1 BOX	07/01/2015	
1		800 mL in 1 BAG; Type 0: Not a Combination Product		
2	NDC:64233-221-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2015	
3	NDC:64233-221-24	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2015	
4	NDC:64233-221-01	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/01/2015	
5	NDC:64233-221-03	350 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/01/2015	
6	NDC:64233-221-05	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2015	
7	NDC:64233-221-07	700 mL in 1 BAG; Type 0: Not a Combination Product	07/01/2015	
8	NDC:64233-221-09	2000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/01/2015	
9	NDC:64233-221-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/01/2015	
10	NDC:64233-221-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2015	
11	NDC:64233-221-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	07/01/2015	
12	NDC:64233-221-13	800 mL in 1 BAG; Type 0: Not a Combination Product	07/01/2015	

13	NDC:64233-221-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2015	
14	NDC:64233-221-15	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2015	
15	NDC:64233-221-28	149 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2015	
16	NDC:64233-221-27	800 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	07/01/2015	
17	NDC:64233-221-55	208200 mL in 1 DRUM; Type 0: Not a Combination Product	07/01/2015	
18	NDC:64233-221-08	1 in 1 BOX	07/01/2015	
18		1000 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/01/2015	

Labeler - Mid American Research Chemical Corp (051254092)

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

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(64233-221)

Revised: 1/2019

Mid American Research Chemical Corp