HAND-AID- alcohol gel
ABC Compounding Co., Inc.

Hand-Aid 6605 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

Ethyl Alcohol 70% 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, 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, DMDM hydantoin, diidopropylamine, carbomer, propylene glycol, tocopheryl acetate, aloe barbadensis,

Hand-Aid 6605 Drug Facts and Label

800ml bag 1-15-2021

HAND-AID

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62257-122
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
CARBOMER 934 (UNII: Z135WT9208)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62257-122-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/15/2021	
2	NDC:62257-122-24	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/15/2021	11/19/2025
3	NDC:62257-122-01	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/15/2021	11/19/2025
4	NDC:62257-122-28	149 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/15/2021	11/19/2025
5	NDC:62257-122-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/15/2021	
6	NDC:62257-122-13	800 mL in 1 BAG; Type 0: Not a Combination Product	01/15/2021	
7	NDC:62257-122-47	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/15/2021	11/19/2025
8	NDC:62257-122-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/15/2021	
9	NDC:62257-122-55	208200 mL in 1 DRUM; Type 0: Not a Combination Product	01/15/2021	11/19/2025
10	NDC:62257-122-21	2.5 mL in 1 DOSE PACK; Type 0: Not a Combination Product	01/15/2021	11/19/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	01/15/2021	

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

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

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(62257-122)

Revised: 11/2025 ABC Compounding Co., Inc.