## HAND-AID- alcohol gel Pro Chem, 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.

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## Hand-Aid 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, DMDM hydantoin, diidopropylamine, carbomer, propylene glycol, tocopheryl acetate, aloe barbadensis,

## Hand-Aid 6605 Drug Facts and Label



## Drug Facts (Continued)

## 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

660525L984.071620

#### Hand-Aid 6605 Label

#### HAND-AID

alcohol gel

#### **Product Information**

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:63830-535

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: 6 DC9 Q167V3)		
ALOE VERA LEAF (UNII: ZY81Z83H0 X)		
.ALPHATO COPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)		

DIISO PRO PYLAMINE (UNII: BR9 JLI40 NO)	
DMDM HYDANTO IN (UNII: BYR0 546 TOW)	

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:63830-535- 21	2.5 mL in 1 DOSE PACK; Type 0: Not a Combination Product	07/27/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/27/2020	

## **Labeler -** Pro Chem, Inc. (061396065)

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

# Establishment Name Address ID/FEI Business Operations ABC Compounding Co., Inc. 003284353 manufacture(63830-535)

Establishment			
Name	Address	ID/FEI	Business Operations
Tekna Fill, Inc.		081509076	repack(63830-535)

Revised: 7/2020 Pro Chem, Inc.