PRO ADVANTAGE INSTANT HAND SANITIZER- alcohol liquid
NDC National Distribution & Contracting, 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.

Pro Advantage Instant Hand Sanitizer

Drug Facts

ACTIVE INGREDIENT:

Alcohol, 62%

Purpose

Antiseptic

USE

To help reduce bacteria on the hands that can potentially cause disease.

WARNING:

• For External Use Only
• Flammable, Keep away from fire or flame.

When using this product

• Avoid contact with eyes, if this occurs rinse thoroughly with water and contact a physician.

Ask a doctor before use if you have

• deep wounds, animals bites or serious burns.

Stop use and ask a doctor if

• condition persists.

Keep out of reach of children.

• If swallowed get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Apply a liberal amount to hands and rub hands thoroughly until dry. Do not rinse or wipe off gel.

INACTIVE INGREDIENTS

Water, Glycerin, Fragrance, Carbomer, Triethanolamine, DMDM Hydantoin

REF: P779114 NDC 43128-114-03

Made in China
www.ProAdvantagebyNDC.com

Manufactured for NDC, Inc.
407 New Sanford Road, La Vergne, TN 37086

Pro Advantage by NDC Instant
Hand Sanitizer
4 FL. OZ.
(118 mL)
Kills Germs With 62% Alcohol
+M220P77911480

Product Labels

PRO ADVANTAGE INSTANT HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43128-114
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	620 mg in 1 mL

Inactive Ingredients

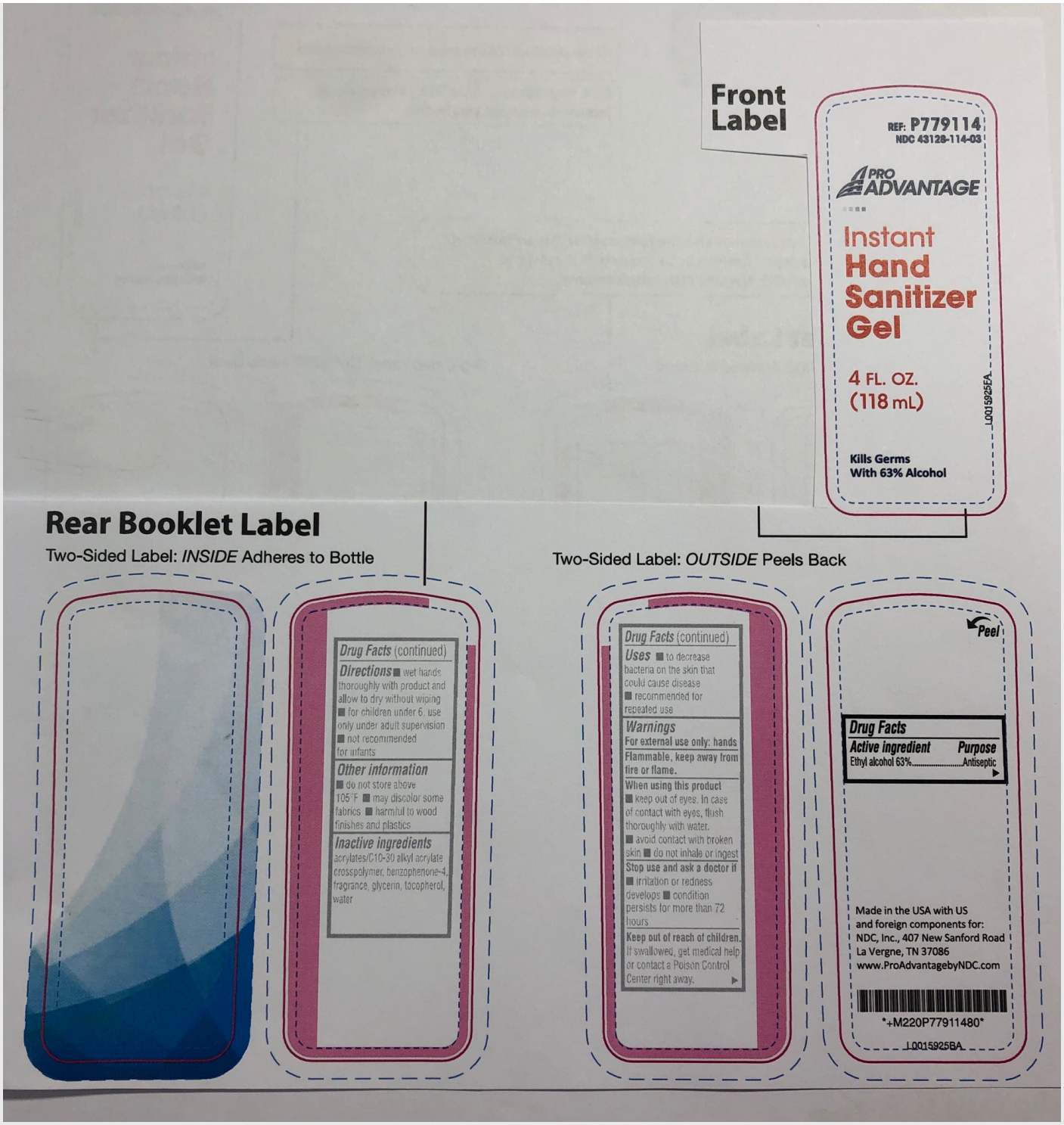
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROLAMINE (UNII: 9O3K93S3TK)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43128-114-03	1 in 1 PACKAGE	03/08/2013	
1		118 mL in 1 BOTTLE; 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	03/08/2013	

Labeler - NDC National Distribution & Contracting, Inc. (009831413)

Establishment

Name	Address	ID/FEI	Business Operations
Dukal		088520668	manufacture(43128-114)

Revised: 3/2023 NDC National Distribution & Contracting, Inc.