PERSANI INSTANT HAND SANITIZER- alcohol spray
Wisconsin Pharmacal Company

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.

Persani Isntant Hand Sanitizer 6 OZ

Purpose

Hand sanitizer to help reduce germs on the skin that can potentially cause disease.

Drug Facts

Active Ingredient

Active ingredient	Purpose
Ethyl Alcohol 62%	Antimicrobial

Uses

■ Hand sanitizer to help reduce germs on the skin that can potentially cause disease. ■ Recommended for repeated use.

Warnings

For external use only.

■ **Flammable: Keep away from fire or flame**. ■ **Contents under pressure.**
Do not puncture or incinerate. Do not store at temperatures above 120F°(48C°).

When using this product ■ Keep out of eyes. In case of eye contact, flush eyes with water.

■ Avoid contact with broken skin.

Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.

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. ■ Rub hands together for 15 seconds, pay attention between fingers, fingernails and cuticles. ■ Allow to dry without wiping.

Inactive ingredients

Aloe Vera, FD&C Blue #1, FD&C Yellow #5, Fragrance, PEG/PPG-18/18 Dimethicone, Tocopheryl Acetate, Water

Package Label - Principal Display Panel

Kills 99.99% of Germs

Persani

Instant

Hand Sanitizer

Ethyl Alcohol 62%

with **Moisturizing**

Vitamin E & Aloe

SmartScent

Odor Eliminination Technology

Safe for the entire family

CONTINUOUS SPRAY

AT ANY ANGLE

NET WT 6 OZ (170.1 g)

PERSANI INSTANT HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68093-7116
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 g in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:68093-7116-1	170.1 g in 1 CAN; Type 0: Not a Combination Product	04/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/15/2020	

Labeler - Wisconsin Pharmacal Company (800873986)

Establishment

Name	Address	ID/FEI	Business Operations
Wisconsin Pharmacal Company		800873986	manufacture(68093-7116) , pack(68093-7116) , label(68093-7116)

Revised: 4/2020 Wisconsin Pharmacal Company