# SANIISWAB- isopropyl alcohol liquid IDO PHARM

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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isopropyl alcohol 70%

Water

two-step method to clean and sanitize the skin of the inner nostrils

■ cleans ■ sanitizes

Keep out of reach of children

Adults and children over 12 years of age
Step 1. Cleaning. Open Step 1 packet and hold handle, inserting
one swab tip into each nostril. Swab lower interior lining inside the
nose in a circular motion 7 times. Handle guard protects against
insertion of swab tip beyond the nostrils. Discard.
Step 2. Sanitizing. Repeat Step 1 with Step 2 packet. Discard

Flammable, keep away from fire and flames For external use only

For external use only











### WHY THIS WORKS

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SURGICALLY PROVEN

"Your nose is the main entry point for infection—which is why daily nasal care is as important as washing your hands.
My Z-step cleaning and santizing method—used in over 2,000 nasal procedures with ZERO infections—is now available for anytime, anywhere use...
SaniiSwab"."









### WASH AWAY IRRITANTS & ALLERGENS

Open Step 1 packet. Remove and hold SaniiSwab<sup>III</sup> handle between thumb and

### WIPE OUT BACTERIA & VIRUSES



PATENTED\* PROCESS

### **Drug Facts** Purpose

Stop use and ask a doctor if redness, pain, or irritation appear and lasts more than 72 hours. **Keep out of reach of children** 



### **SANIISWAB**

isopropyl alcohol liquid

### **Product Information**

HUMAN OTC DRUG **Item Code (Source)** NDC:77039-026 **Product Type** 

NASAL **Route of Administration** 

### **Active Ingredient/Active Moiety**

**Basis of Ingredient Name** Strength Strength ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL -**ISOPROPYL** 70 mL UNII:ND2M416302) ALCOHOL in 100 g

Inactive	Ingredients
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**Ingredient Name** Strength

WATER (UNII: 059QF0KO0R)

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:77039-026- 01	0.3 g in 1 POUCH; Type 0: Not a Combination Product	09/26/2023		

Marketing In	keting Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	09/20/2023			

## **Labeler -** IDO PHARM (694853523)

Revised: 9/2023 IDO PHARM