WILLIAMS HAND SANITIZER- alcohol spray
Combe Incorporated

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.

Williams Hand Sanitizer Spray

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Drug Facts

Active ingredient

Ethyl Alcohol 70% w/w

Purpose

Antimicrobial

Use

to help reduce bacteria on the skin. For use when soap and water are not available.

Warnings

Flammable. Keep away from fire or flame.

For external use only.

When using this product,

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash appears. Do not use on broken skin.

Keep out of reach of children.

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

Directions

- Spray enough products on hands to cover all surfaces. Rub hands together briskly until dry.
- Rub hands together briskly until dry.
- Children should be supervised when using this product.

Other Information

- Store below 86°F (30°C)
- May discolor certain fabrics or surfaces

Questions?

Call **1-800-431-2610** weekdays 9 am to 5 pm EST

Inactive ingredients

water, glycerin, aloe barbadensis leaf juice powder.

PRINCIPAL DISPLAY PANEL

Williams

Hand Sanitizer Spray

Kills 99.9% of many common bacteria

Active Antibacterial Ingredient: 70% Alcohol

Moisturizing Glycerin + Aloe

Fragrance Free

1 fl. oz. (29.5 mL)

WILLIAMS HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11509-0401

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)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11509-0401-1	29.5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	05/18/2020	

Labeler - Combe Incorporated (002406502)

Revised: 5/2020 Combe Incorporated