ALCOHOL HAND SANITIZER- alcohol spray
Webb Business Promotion

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.

Antibacterial Hand Sanitizer

Directions:

Spray enough product on hands to cover all surfaces. Rub hands together until dry.

Active Ingridient

Ethyl Alcohol (62%).

Inactive Ingredients:

Deionized Water, Glycerin, Propylene Glycol, Tocopherol, Citrus Fragrance, Aloe Vera.

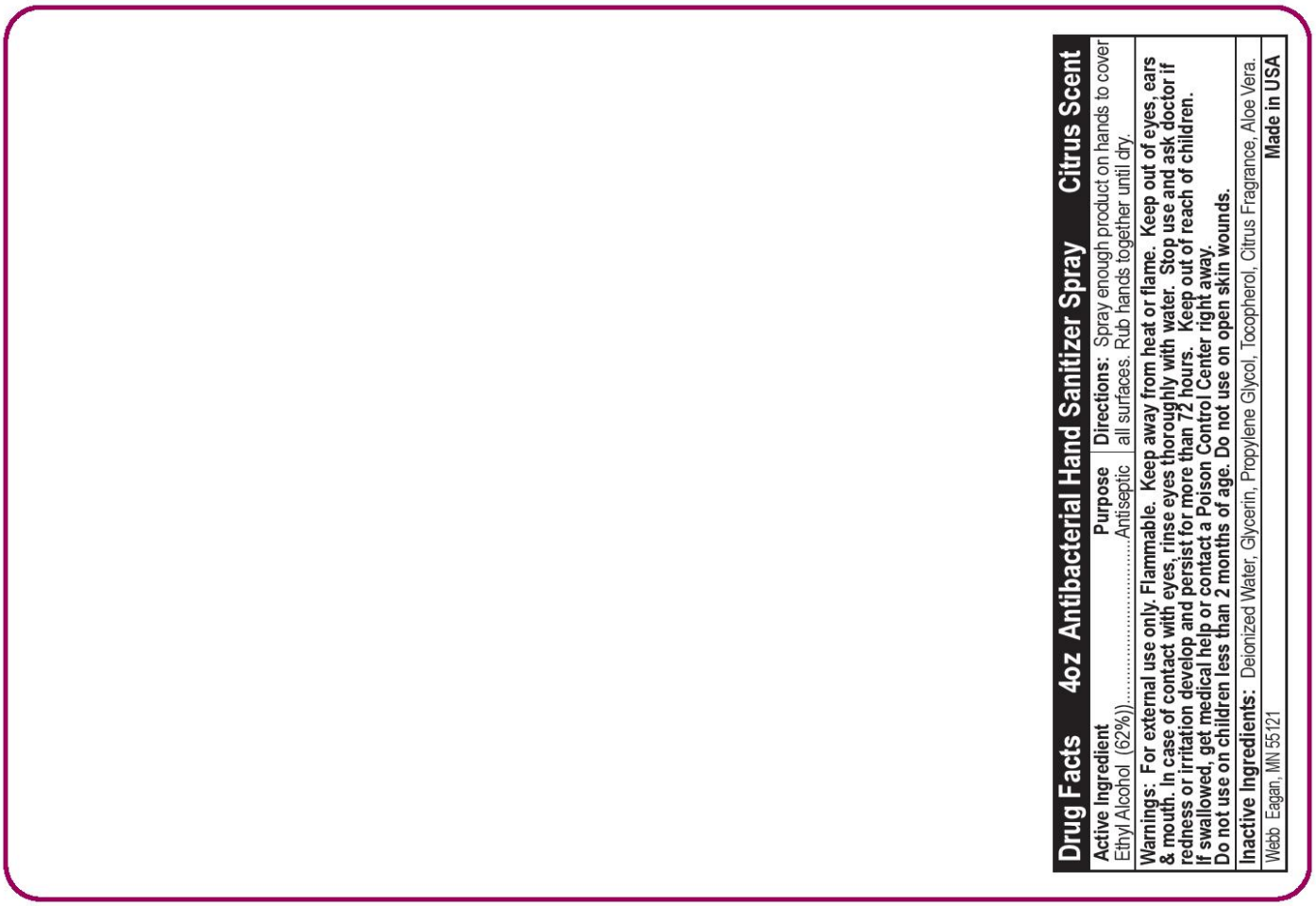
Keep out of reach of children.

Do not use on children less than 2 months of age. Do not use on open skin wounds.

For external use only. Flammable. Keep away from heat or flame. Keep out of eyes, eras & mouth. In case of contact with eyes, rinse eyes thoroughly with water. If swallowed, get medical help or contact a Poison Control Center right away.

Antibacterial hand sanitizer spray.

Antiseptic.



ALCOHOL HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70445-412
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
TOCOPHEROL (UNII: R0ZB2556P8)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70445-412-20	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
2	NDC:70445-412-22	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
3	NDC:70445-412-23	10 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/25/2020	
4	NDC:70445-412-26	20 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/25/2020	
5	NDC:70445-412-21	5 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/08/2020	

Labeler - Webb Business Promotion (154445647)

Establishment

Name	Address	ID/FEI	Business Operations
Webb Business Promotion		154445647	manufacture(70445-412)

Revised: 1/2022 Webb Business Promotion