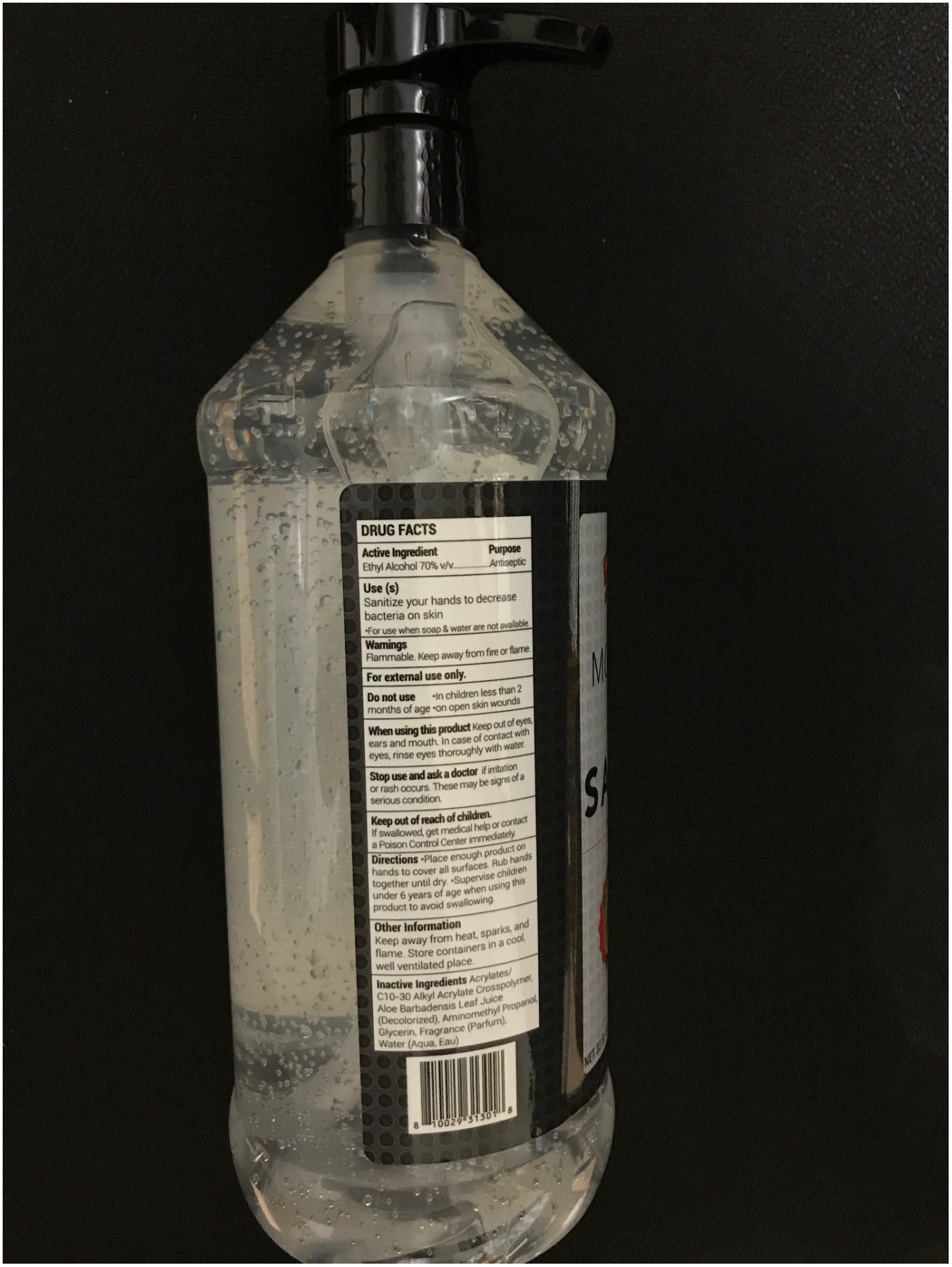
SANISPIRE HAND SANITIZER- ethyl alcohol gel
RNA PHARMA, LLC

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.


DRUG FACTS
Active Ingredient
Ethyl Alcohol 70% v/v
Purpose
Antiseptic
Use (s)
Sanitize your hands to decrease bacteria on skin
•For use when soap & water are not available
Warnings
Flammable. Keep away from fire or flame.
For external use only.
Do not use
•In children less than 2 months of age •on open skin wounds
When using this product
Keep out of eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly with water.
Stop use and ask a doctor
if irritation or rash occurs. These may be signs of a serious condition.
Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center immediately.
Directions
•Place enough product on hands to cover all surfaces. Rub hands together until dry. •Supervise children under 6 years of age when using this product to avoid swallowing.
Other Information
Keep away from heat, sparks, and flame. Store containers in a cool, well ventilated place.
Inactive Ingredients
Acrylates/ C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Juice (Decolorized), Aminomethyl Propanol, Glycerin, Fragrance (Parfum), Water (Aqua, Eau)
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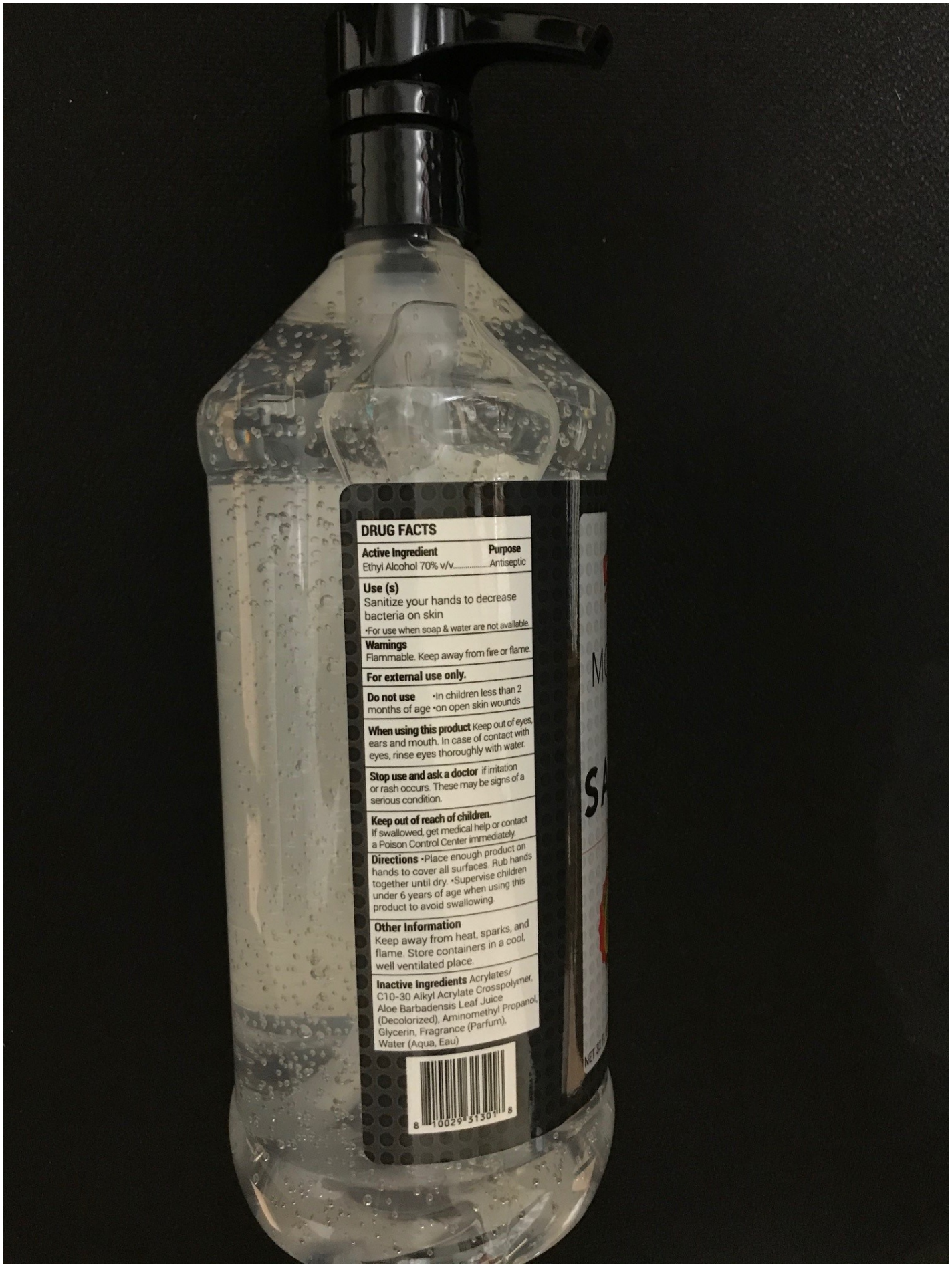
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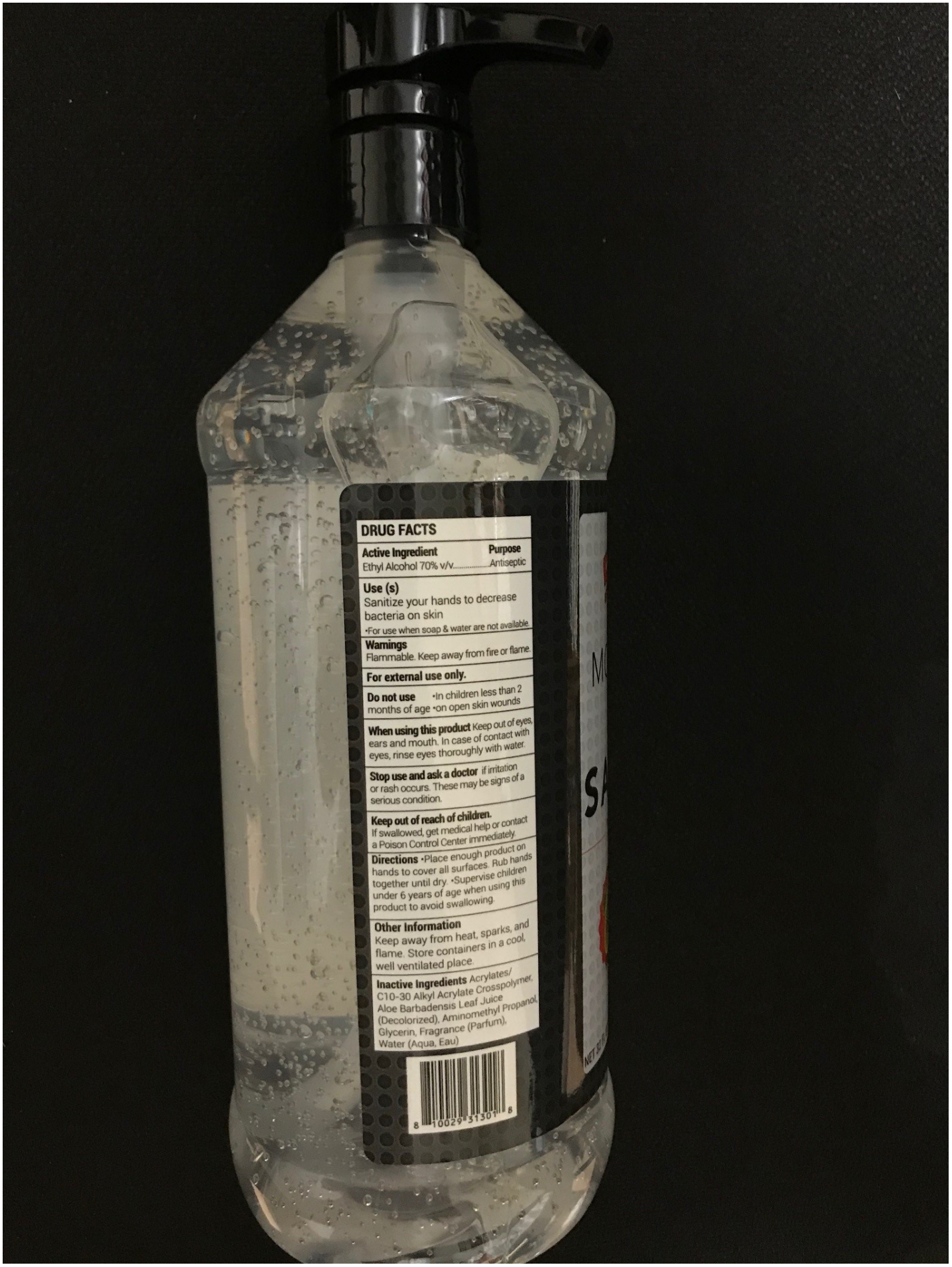
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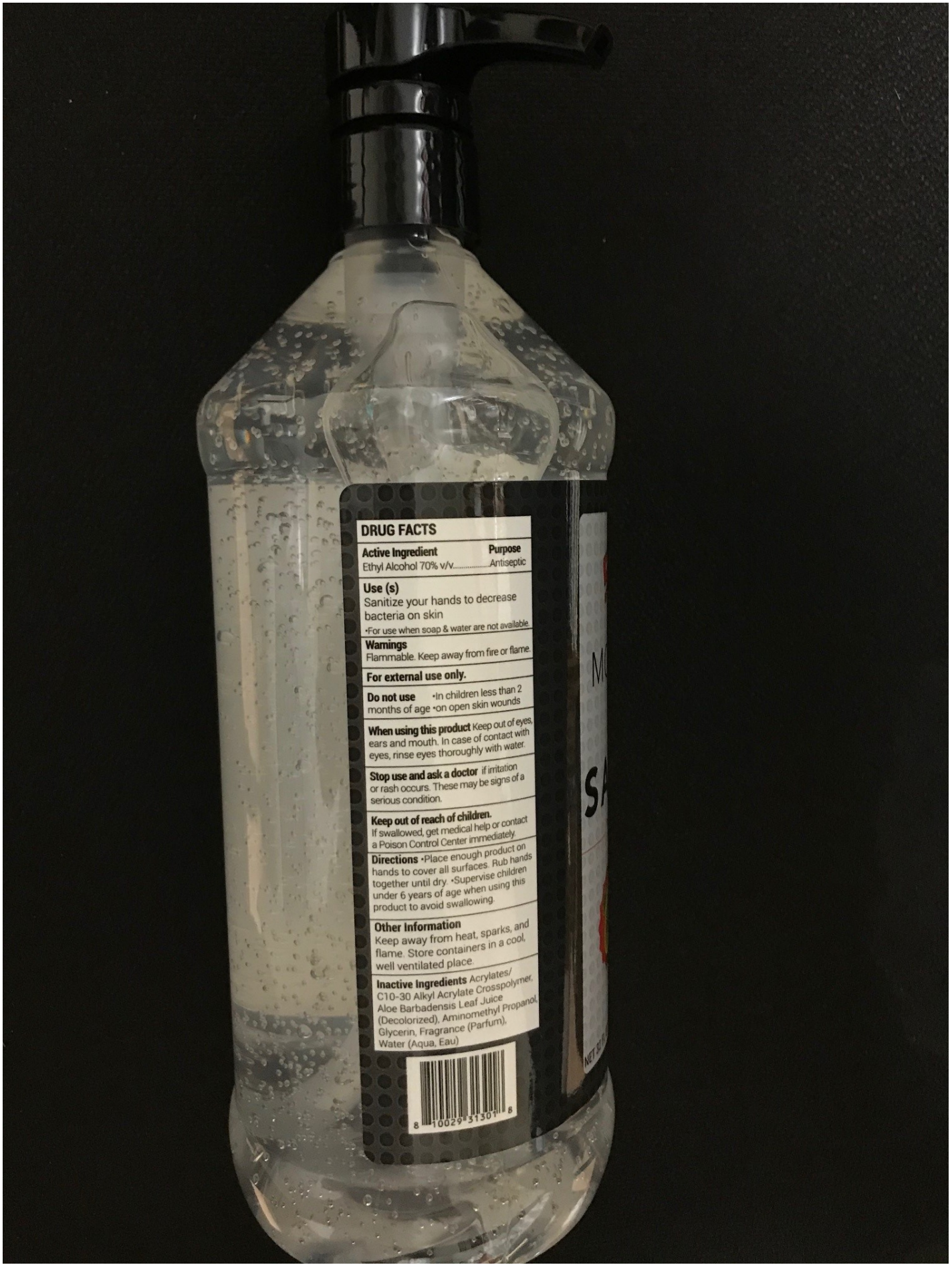
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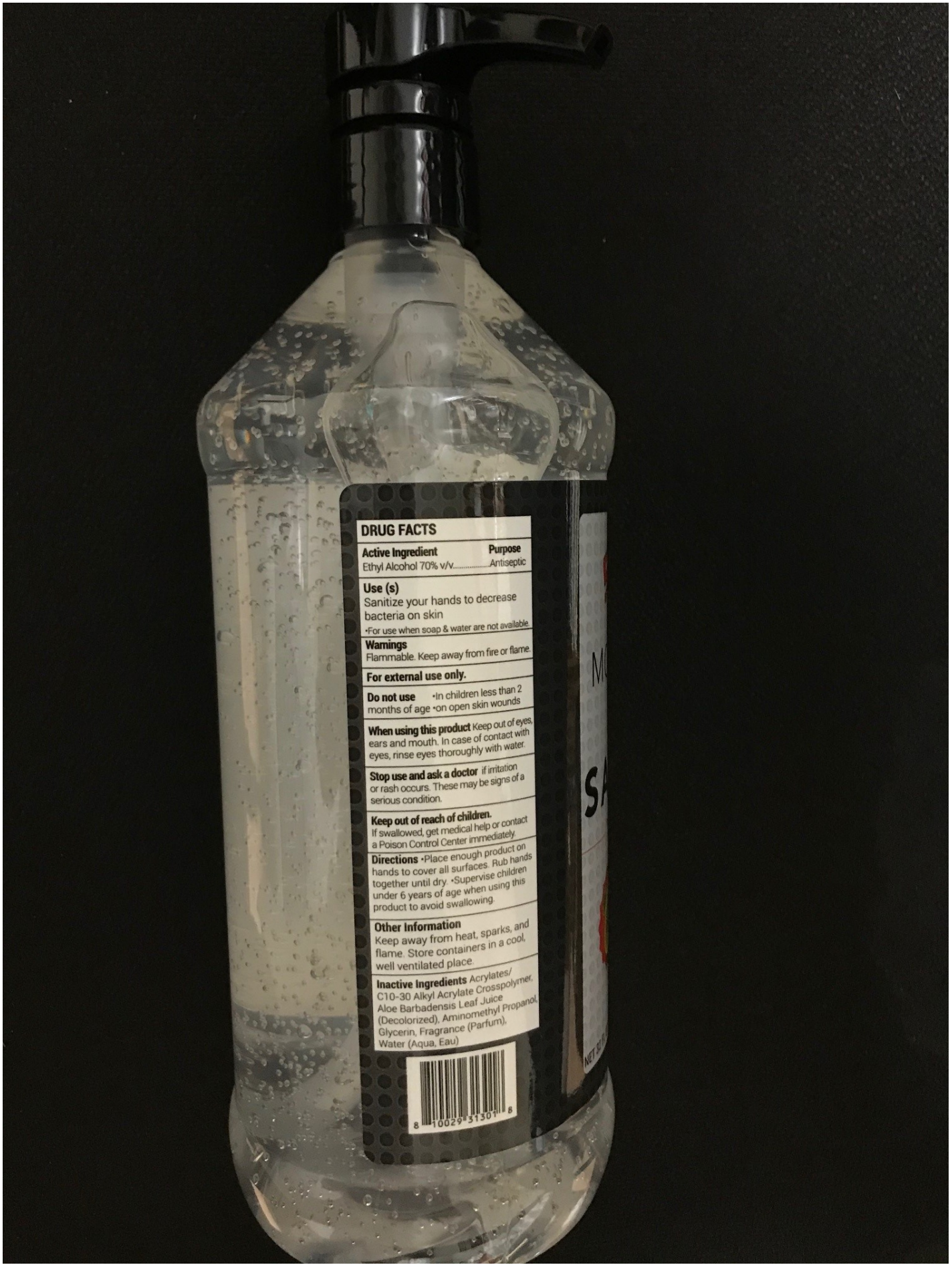
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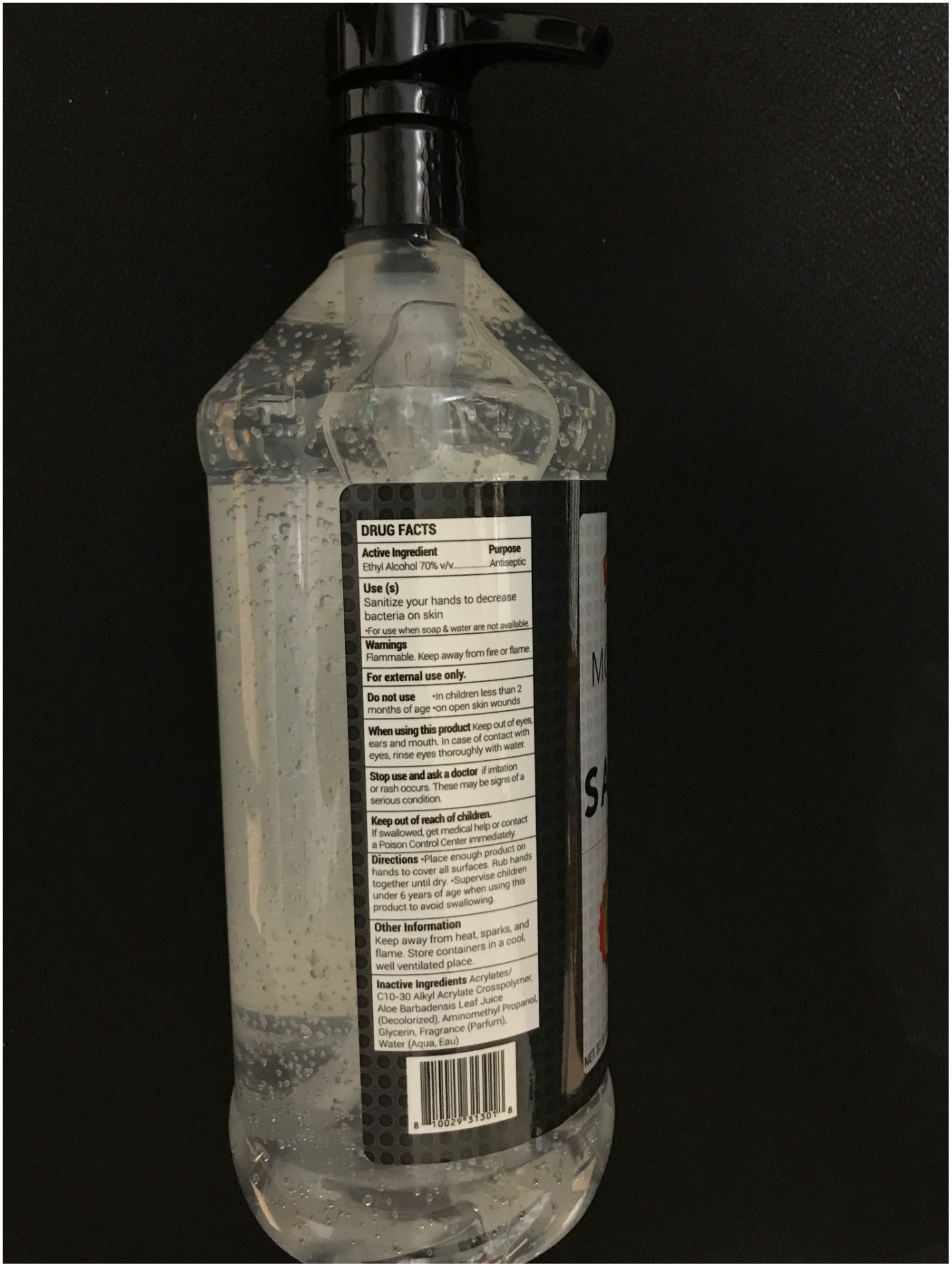
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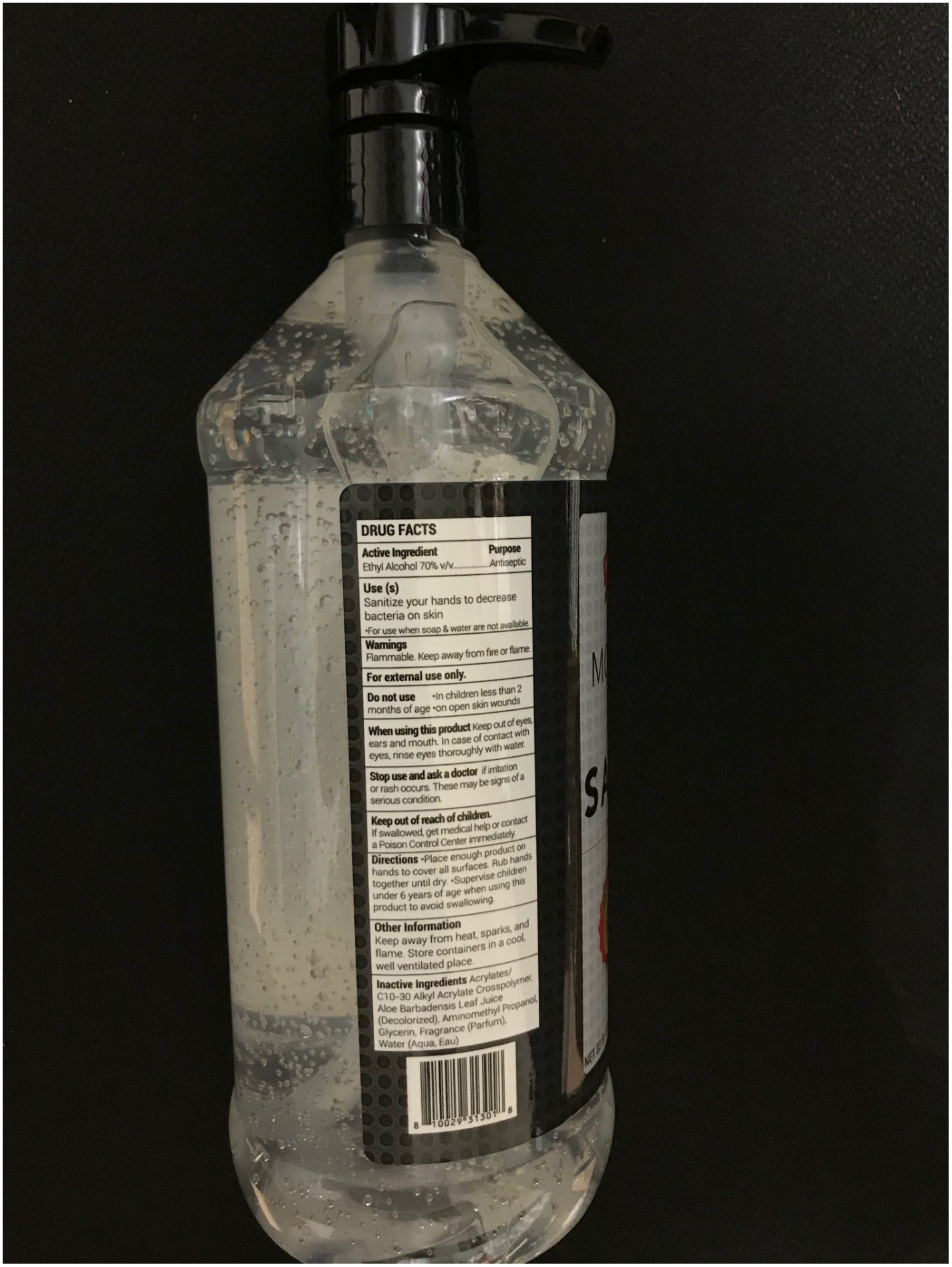
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SANI-SPIRE™
MOISTURIZING
HAND
SANITIZER
70% ETHYL ALCOHOL
Fresh Lemon Scent
MADE IN USA
MADE IN A FDA

SANISPIRE HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60717-811
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)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 132584PQMO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Product Characteristics			
Color		**Score**	
Shape	ROUND	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:60717-811-32	960 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/09/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/09/2021	

Labeler - RNA PHARMA, LLC (079103999)

Establishment

Name	Address	ID/FEI	Business Operations
RNA PHARMA, LLC		079103999	manufacture(60717-811)

Revised: 4/2021