

SONO HAND SANITIZER- benzalkonium chloride solution
Advanced Ultrasound Solutions Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

SONO Foaming Hand Sanitizer

Active Ingredient

Benzalkonium chloride.....0.13%

Purpose

Antiseptic Hand Sanitizer

Uses

- Hand sanitizer to help reduce bacteria on skin
- Recommended for repeated use

Warnings

For external use only

Do not use in ears or mouth

When using this product

Avoid contact with eyes. In case of contact, flush eyes with water.

Stop use and ask a doctor

If redness or irritation develops and persists for more than 72 hours.

Keep out of reach of children

Children should be supervised when using this product

Directions

- Apply a small amount onto hand, then spread on both hands
- Rub hands together until dry

Inactive ingredients

Glycerin, PEG/PPG-18/18 Dimethicone, water, Organic Citrus Aurantium Bergamia Fruit Oil.

Sono Foaming Hand Sanitizer



Questions or Comments
Call 1-855-USWIPES (879-4737)
SONO HEALTHCARE
www.sonohealthcare.com





FOAMING HAND SANITIZER

WITH MOISTURIZER AND BERGAMOT ESSENTIAL OIL



Kills 99.99% of most common germs

1.7 FL OZ (50 ml)

Drug Facts

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Drug Facts (continued)

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Manufactured for SONO HEALTHCARE
4715 Elm Street, #102, Murrieta, CA 92562

SONO HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77677-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
WATER (UNII: 059QF0KO0R)	
BERGAMOT OIL (UNII: 39W1PKE3JI)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77677-001-02	50 g in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2020	
2	NDC:77677-001-19	550 g in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/17/2020	

Labeler - Advanced Ultrasound Solutions Inc. (079552984)

Registrant - Advanced Ultrasound Solutions Inc. (079552984)

Establishment

Name	Address	ID/FEI	Business Operations
Morgan Gallacher Inc. DBA Custom Chemical Formulators Inc.		028311595	manufacture(77677-001) , api manufacture(77677-001) , pack(77677-001)

Revised: 12/2023

Advanced Ultrasound Solutions Inc.