

**WALGREENS FRESH ANTIBACTERIAL FOAMING HAND- benzalkonium chloride soap**  
**Walgreen Company**

*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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**Walgreens Fresh Antibacterial Foaming Hand Soap**

**Drug Facts**

**Active ingredient**

Benzalkonium Chloride 0.13%

**Purpose**

Antibacterial

**Use**

for handwashing to decrease bacteria on the skin

**Warnings**

For external use only: hands only

**When using this product**

- avoid contact with eyes. If contact occurs, rinse eyes with water.

**Stop use and ask a doctor if**

- irritation and redness develop
- condition persists for more than 72 hours

**Keep out of reach of children.**

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

**Directions**

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

**Inactive ingredients**

water, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamido-propylamine oxide, glycerin, fragrance, citric acid, tetrasodium EDTA, benzophenone-4, sodium benzoate, blue 1, red 33

**Questions or comments?**

1-800-925-4733

## Company Information

DISTRIBUTED BY:

WALGREEN CO.

200 WILMOT RD.

DEERFIELD, IL 60015

## Product Packaging

FRESH

ANTIBACTERIAL

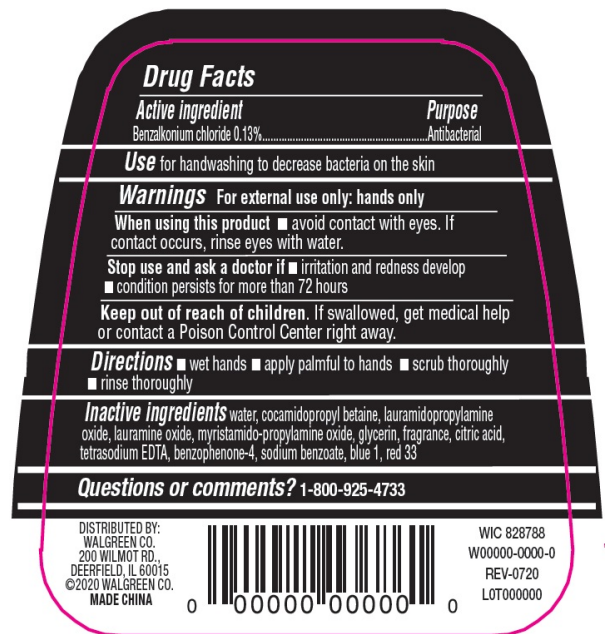
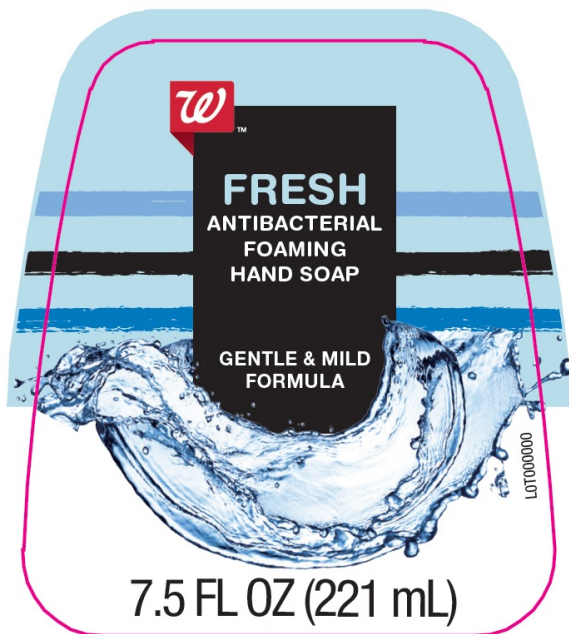
FOAMING

HAND SOAP

GENTLE & MILD

FORMULA

7.5 FL OZ (221 mL)



## WALGREENS FRESH ANTIBACTERIAL FOAMING HAND

benzalkonium chloride soap

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9000
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)		BENZALKONIUM CHLORIDE	1.3 mg in 1 mL	
<b>Inactive Ingredients</b>				
Ingredient Name		Strength		
WATER (UNII: 059QF0KO0R)				
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)				
LAURAMINE OXIDE (UNII: 4F6FC4M18W)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
SULISOBENZONE (UNII: 1W6L629B4K)				
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)				
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)				
GLYCERIN (UNII: PDC6A3C0OX)				
EDETATE SODIUM (UNII: MP1J8420LU)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-9000-01	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/10/2020	
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	08/10/2020		

**Labeler** - Walgreen Company (008965063)

Revised: 8/2020

Walgreen Company