WALGREENS HAND SANITIZER COASTAL BREEZE SCENT- alcohol gel
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.

Walgreens Hand Sanitizer Coastal Breeze Scent

Drug Facts

Active ingredient

Alcohol 69% v/v

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only

Flammable, keep away from fire or flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

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 thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision

Other information

- do not store above 105F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

Water, Glycerin, Propylene Glycol, Carbomer, Aloe Barbadensis Leaf Extract, Aminomethyl Propanol, Fragrance, Tocopheryl Acetate, Denatonium Benzoate, Blue 1, Yellow 5.

Company Information

DISTRIBUTED BY: WALGREEN CO.,

200 WILMOT RD., DEERFIELD, IL 60015

Product Packaging

Walgreens

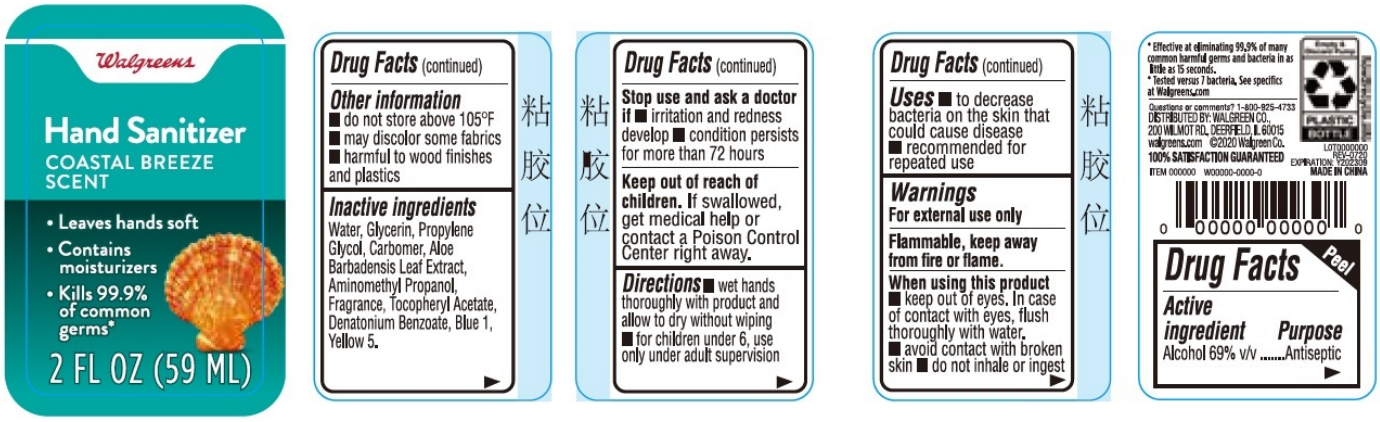
Hand Sanitizer

COASTAL BREEZE

SCENT

- Leaves hands soft
- Contains moisturizers
- Kills 99.9% of common germs

2 FL OZ (59 ML)



WALGREENS HAND SANITIZER COASTAL BREEZE SCENT

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-9006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	69 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0363-9006-01	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/11/2020	

Labeler - Walgreen Company (008965063)

Revised: 8/2020 Walgreen Company