

ANTISEPTIC FOAMING HAND SANITIZER- benzalkonium chloride liquid
Target Corporation

Up & Up 812.000/812AA
Foaming Hand Sanitizer Alcohol Free

Claims

- Made without parabens, phthalates, phenoxyethanol or sulfates**
- **Formulated without sulfated surfactants
- Not tested on animals

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Use

- to decrease bacteria on the skin

Warnings

For external use only - hands only

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin

Stop use and ask a doctor if

- skin irritation develops
- 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
- not recommended for infants

Inactive ingredients

water, glycerin*, diglycerin, disodium cocoamphodiacetate, lauramine oxide, Aloe barbadensis leaf juice*, sodium benzoate, cetrimonium chloride, citric acid*, methoxy PEG/PPG-7/3 aminopropyl dimethicone, propylene glycol, tetrasodium EDTA, Chamomilla recutita (matricaria) flower extract*, Pyrus malus (apple) fruit extract*, Citrus grandis (grapefruit) peel oil*, Citrus nobilis (mandarin orange) peel oil*, limonene

Claims

*Plant-derived ingredients

†Based on laboratory testing against certain bacteria

Adverse Reaction

Dist. by Target Corporation

Minneapolis, MN 55403

Made in U.S.A. with U.S. and foreign components.

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Target Brands, Inc.

Guest Services

1-800-910-6874

Principal display panel

NDC 82442-812-20

Alcohol-Free

Foaming Hand Sanitizer

Citrus Scent

- Hypoallergenic
- Made with plant-derived ingredients*
- Kills 99.9% certain germst†
- Dermatologist tested

up&up™

2-1.5 FL OZ (44 mL) TOTAL 3 FL OZ (88 mL)



ANTISEPTIC FOAMING HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:82442-812

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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIGLYCERIN (UNII: 3YC120743U)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHOXY PEG/PPG-7/3 AMINOPROPYL DIMETHICONE (UNII: 4M7P1JZ2V2)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CHAMOMILE (UNII: FGL3685T2X)	
APPLE (UNII: B423VGH5S9)	
CITRUS MAXIMA FRUIT RIND OIL (UNII: 8U3877WD44)	
MANDARIN OIL (UNII: NJO720F72R)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-812-20	2 in 1 CARTON	10/11/2024	
1		88 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/11/2024	

Labeler - Target Corporation (006961700)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091520	manufacture(82442-812)

Revised: 2/2026

Target Corporation