HAND SANITIZER WIPES- juniper clean cloth
KYNC Design LLC

Hand Sanitizer Wipes

SPL UNCLASSIFIED SECTION

Active Ingredient(s)

Benzalkonium Chloride (CAS NO: 68424-85-1), 0.125% .

Purpose

Antimicrobial

Use

Hand sanitizing wipes help remove bacteria from hands and kill 99.99% of germs. Removes dirt to help with health and cleanliness.

WARNINGS

Use on hands only.

Do not use

If you are allergic to any of the ingredients.

Stop use and ask a doctor

if irritation or rash occurs.

Do not use in or near the eyes.

If contact occurs rinse with plenty of water. Remove contact lenses, if present and easy to do. Continue rinsing. If eye irritation persists, get medical advice/attention.

Keep out of reach of children.

If swallowed, get medical help or contact a poison control center immediately.

Directions

To Open Package: Flip open dispensing cap. Locate wipe at center of roll and pull through small openning in lid. For best results dispense wipes at an angle. Wet hands thoroughly with product and allow to dry. When finished, snap lid cap shut to retain moisture. Discard wipe in trash receptacle after use. Do not flush.

Inactive ingredients

Water(Aqua), Ethanol, Decyl Glucoside, Glycerine, Aloe Vera Extract, D-Panthenol, Fragrance (Perfume)

Package Label - Principal Display Panel

110 mL NDC: 76557-001-01

245 mL NDC: 76557-001-01

HAND SANITIZER WIPES

juniper clean cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76557-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.125 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
CLOVE (UNII: K48IKT5321)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
DEXPANTHENOL (UNII: 1O6C93RI7Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76557-001-04	75 in 1 CANISTER	06/04/2020	
1		3.14 mg in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
2	NDC:76557-001-03	35 in 1 CANISTER	06/04/2020	
2		3.14 mg in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
3	NDC:76557-001-05	72 in 1 PACKAGE	06/04/2020	
3		3.14 mg in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/04/2020	

Labeler - KYNC Design LLC (039933298)