HYPERCLN SANITARY WIPE- anti-microbial wipe cloth
Falcon Safety Products, Inc.

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.

Hypercln Sanitizing Wipes

Active ingredient

Benzalkonium chloride 0,13%

Purpose

Anti-bacterial

Indications

Hand sanitizer to help reduce bacteria on skin.

To use

To Use: open, unfold and apply. Sanitize hands or other affected area. Discard in trash after use. Do not flush.

Warnings

Warnings

For external use only.

When using this product avoid contact with eyes. If in eyes, flush thoroughly with water.

Ask a doctor if

Stop and ask a doctor if:

• irritation develops. • condition persists for more than 72 hours.

Keep out of the reaach of children

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

Inactive ingredients

Inactive Ingredients: Water, Decyl Glucoside, Glycerin, Sodium hydroxymethylglycinate, Citric Acid, PEG-12 Dimethicone, Fragrance, Sodium Benzoate, Potassium Sorbate, Allantoin, Tocopheryl Acetate, Aloe Barbadensis Leaf Water, Chamomilla Recutita (Matricaria) Flower Water, Panthenol, Tetrasodium EDTA, Benzyl Alcohol, Sodium Hydroxide

Principal Display Panel

HYPERCLN SANITARY WIPE

anti-microbial wipe cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78847-322
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0013

Inactive Ingredients

Ingredient Name	Strength
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CHAMOMILE (UNII: FGL3685T2X)	
CITRONITRILE (UNII: MS598KEL3M)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TETRAHYDROLINALOOL (UNII: UM4XS5M134)	

EUCALYPTOL (UNII: RV6J6604TK)	
3,3'-DIHYDROXYDIPROPYL ETHER (UNII: S1OQ81LMNA)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
PANTHENOL (UNII: WV9CM0O67Z)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
3,7-DIMETHYL-1-OCTANOL (UNII: DPY9K1927C)	
SODIUM HYDROXYMETHYLGLYCINATE (UNII: DIG6BWZ9XT)	
ALLANTOIN (UNII: 344S277G0Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PEG-12 DIMETHICONE (UNII: ZEL54N6W95)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78847-322-01	1 in 1 NOT APPLICABLE; Type 0: Not a Combination Product	07/13/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/13/2020	

Labeler - Falcon Safety Products, Inc. (150582500)

Registrant - Dan Mor (514523067)

Establishment

Name	Address	ID/FEI	Business Operations
Dan Mor		514523067	manufacture(78847-322)

Revised: 7/2020 Falcon Safety Products, Inc.