ULTRA DEFENSE SANI SMART HAND SANITIZING WIPES- benzalkonium chloride cloth
K7 Design Group 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.

Ultra Defense Sani Smart Hand Sanitizing Wipes

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

- For hand sanitizing to decrease bactera on the skin.
- Apply topically to the skin to help prevent cross contamination.
- Recommended for repeated use.
- Dries in seconds.

Warnings

For external use only.

When using this product

do not use in or contact the eyes.

Stop use and ask a doctor if

too much skin irritation or sensitivity develops or increases.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- Remove lid and discard seal from container.
- Pull wipe from center of roll and thread through opening in lid.
- Do not push finger through opening.
- Replace lid, pull wipe up, and then out at a 45º angle. The next wipe dispenses automatically.
- Close lid to retain moisture.

Other information

- Lot No. Manufacture date and Expiration date can be found on canister.

Inactive ingredients

Water, Phenoxyethanol, Ethyl Alcohol, DMDM Hydantoin, Glycerin, Propylene Glycol, Cocamidopropyl Betaine, Decyl glucoside, Aloe Barbadensis Leaf Extract

Company Information

YOU MAY REPORT A SERIOUS ADVERSE REACTION TO THIS PRODUCT TO REPORT REACTION, LLC, PO BOX 22, PLAINSBORO, NJ 08536

MANUFACTURED FOR & DISTRIBUTED BY K7 DESIGN GROUP INC. 2433 KNAPP STREET, BROOKLYN, NY 11235

Product Packaging - 70 COUNT

ULTRA DEFENSE

SANI+SMART

HAND

SANITIZING

WIPES

70 COUNT

COTTON SOFT WIPES

KILLS 99.99% OF GERMS

5.71 in x 8.07 in (14.5cm x 20.5cm)

ULTRA DEFENSE SANI SMART HAND SANITIZING WIPES

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74177-850
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 g

Inactive Ingredients	
Ingredient Name	**Strength**
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74177-850-70	70 in 1 CANISTER	08/25/2020	
1		191 g in 1 CANISTER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/25/2020	

Labeler - K7 Design Group Inc. (080357784)