ULTRA DEFENSE SANI SMART HAND SANITIZER- alcohol spray
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 Sanitizer Spray

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 and 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 and persist 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

- Spray directly on hands and rub in thoroughly
- For children under 6, use adult supervision
- Not recommended for infants

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Water, Aloe Barbadensis Leaf Extract, Tocopheryl Acetate, Fragrance, Denatonium Benzoate.

Company Information

MANUFACTURED FOR & DISTRIBUTED BY K7 DESIGN GROUP LLC.
2433 KNAPP STREET, BROOKLYN, NY 11235
ORIGIN: CHINA

Product Packaging - 100 mL

ULTRA DEFENSE SANI SMART HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74177-954
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
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74177-954-01	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/14/2020	
2	NDC:74177-954-02	9 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/14/2020	
3	NDC:74177-954-03	3 in 1 PACKAGE	10/14/2020	
3		9 mL in 1 BOTTLE, SPRAY; 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	10/14/2020	

Labeler - K7 Design Group Inc. (080357784)

Revised: 10/2020 K7 Design Group Inc.