ALLOUT NANO- isopropyl alcohol spray
SERENITY TECHNOLOGIES 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.

ALLOUT™ NANO

DRUG FACTS

ACTIVE INGREDIENTS

ISOPROPYL ALCOHOL 75% v/v.

PURPOSE

ANTISEPTIC

USES

HAND SANITIZER TO HELP REDUCE GERMS ON THE SKIN WHEN SOAP AND WATER ARE NOT AVAILABLE.

WARNINGS

FLAMMABLE, KEEP AWAY FROM FIRE AND FLAMES. FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT KEEP OUT OF EYES. IN CASE OF CONTACT WITH EYES RINSE EYES THOROUGHLY WITH WATER.

STOP USE AND ASK A DOCTOR IF IRRITATION AND REDNESS DEVELOP AND 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

APPLY PRODUCT TO HANDS THOROUGHLY, RUB HANDS TOGETHER UNTIL DRY.

OTHER INFO

- CONTAINS PPM LEVELS OF SUB-MICRON AND NANO PARTICLES OF COPPER, ZINC AND SILVER
- STORE BETWEEN 15-30C (59-86F)
- AVOID FREEZING AND EXCESSIVE HEAT ABOVE 40C (104F)

INACTIVE INGREDIENTS

ALOE BARBADENSIS LEAF EXTRACT, ALUM, COPPER, GLYCERINE, HYALURONIC ACID, SILVER, WATER, WHITE CAMPHOR OIL, ZINC.

QUESTIONS?

+1 (951)-587-3753

WWW.ALLOUTNANO.COM

Manufactured By:
Serenity Technologies Inc.
43320 Business Park Dr., B105
Temecula, CA 92590

PRINCIPAL DISPLAY PANEL - 59.1 ml Bottle Label

NANOTECHNOLOGY

ALLOUT™
NANO

ADVANCED
HAND SANITIZER

LONG LASTING
ANTIMICROBIAL PROTECTION
FORMULATED WITH
COPPER, SILVER AND ZINC

SPRAY

SKIN NOURISHING
WITH HYALURONIC ACID

CLEAN HANDS

COUNT

2 fl oz (59.1 ml)

ALLOUT NANO

isopropyl alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75814-103
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
POTASSIUM ALUM (UNII: 1L24V9R23S)	
SILVER (UNII: 3M4G523W1G)	
COPPER (UNII: 789U1901C5)	

PYRITHIONE ZINC (UNII: R953O2RHZ5)	
CAMPHOR OIL, WHITE (UNII: 26P3H26Z9X)	
HYALURONIC ACID (UNII: S270N0TRQY)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75814-103-01	29.6 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/04/2020	
2	NDC:75814-103-02	59.1 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/04/2020	
3	NDC:75814-103-06	177.4 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/04/2020	
4	NDC:75814-103-08	236.6 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/04/2020	
5	NDC:75814-103-34	1000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/04/2020	

Labeler - SERENITY TECHNOLOGIES INC. (829399240)

Revised: 8/2020 SERENITY TECHNOLOGIES INC.