BELSON ALCOHOL FREE HAND SANIZITER- hypochlorous acid liquid KEWS CORPORATION

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

BELSON ALCOHOL FREE HAND SANITIZER

ACTIVE INGREDIENT

HYPOCHLOROUS ACID -- 0.007%

Inactive Ingredients

Water

Purpose

Antimicrobial

Directions for Use

- -Use when soap and water not available to supplement regular hand washing.
- -Apply liberally to hands and rub hands thorougly until product is dried
- -Spray enough to the suspected tool, furniture and space.

Intended Use

- -For personal sanitizing to decrease bacteria on skin
- -Recommended for repeated use

USES

- -For personal sanitzing to decrease bacteria on skin
- -Recommended for repeated use

Keep out of reach of children.

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

Warnings

For External Use

100mL PDP



PATENT NO.: 10-1323506, 10-1027830 UN VENDOR ID: 33794 Net Wt. 3.38 fl. oz (100ml)



100mL NDC 75644-003-01

500mL PDP



500mL NDC 75644-003-02

1LT PDP

1LT NDC 75644-003-03



99.9% ANTIBACTERIAL AND DEODORIZING

PATENT NO. : 10-1323506, 10-1027830 UN VENDOR ID : 33794

Net Wt. 33.8fl. oz (1LT)



4LT PDP

4LT NDC 75644-003-04



99.9% ANTIBACTERIAL AND DEODORIZING

PATENT NO.: 10-1323506, 10-1027830 UN VENDOR ID: 33794

Net Wt. 1.05 gal (4LT)



20LT PDP



99.9% ANTIBACTERIAL AND DEODORIZING

PATENT NO.: 10-1323506, 10-1027830 UN VENDOR ID: 33794 Net Wt. 52.9 gal (20LT)





20LT NDC 75644-003-05

BELSON ALCOHOL FREE HAND SANIZITER

hypochlorous acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75644-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
	HYPOCHLOROUS ACID	0.007 mg in 1 mL	

Inactive Ingredients			
	Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:75644- 003-01	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/19/2020		
2	NDC:75644- 003-02	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/19/2020		
3	NDC:75644- 003-03	1000 mL in 1 JUG; Type 0: Not a Combination Product	06/19/2020		
4	NDC:75644- 003-04	4000 mL in 1 JUG; Type 0: Not a Combination Product	06/19/2020		

5 NDC:75644- 003-05	20000 mL in 1 JUG; Type 0: Not a Combination of 06/19/2020		
Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
	Citation	Date 06/19/2020	Date

Labeler - KEWS CORPORATION (688640317)

Registrant - KEWS CORPORATION (688640317)

Establishment			
Name	Address	ID/FEI	Business Operations
KEWS CORPORATION		688640317	manufacture(75644-003), pack(75644-003), label(75644-003)

Revised: 1/2022 KEWS CORPORATION