ABSOCLEAN-S SANITIZING MIST- 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.

AbsoClean-S Sanitizing Mist

ACTIVE INGREDIENT

HYPOCHLOROUS ACID -- 0.006%

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

Purpose

-For personal sanitzing to decrease bacteria on skin

-Recommended for repeated use

Warnings

For External Use

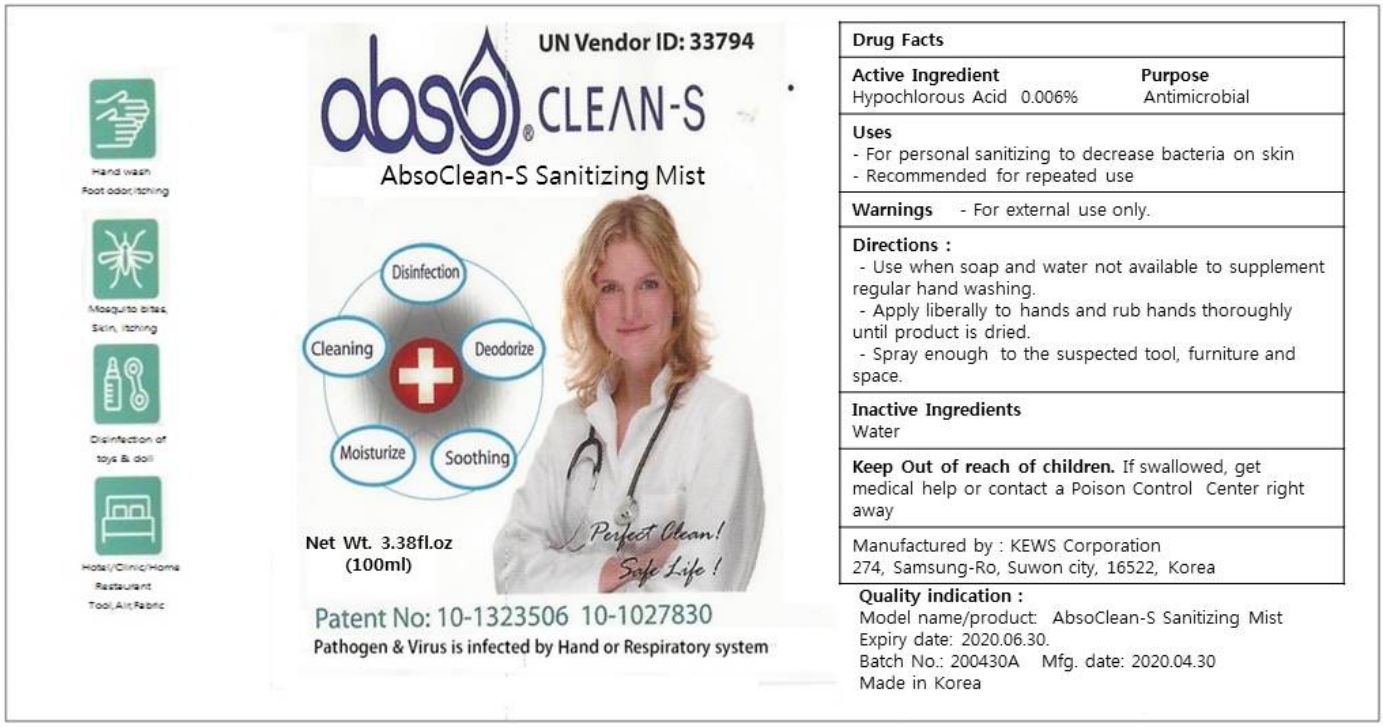
Keep out of reach of children.

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

100mL PDP

100mL NDC 75644-002-01

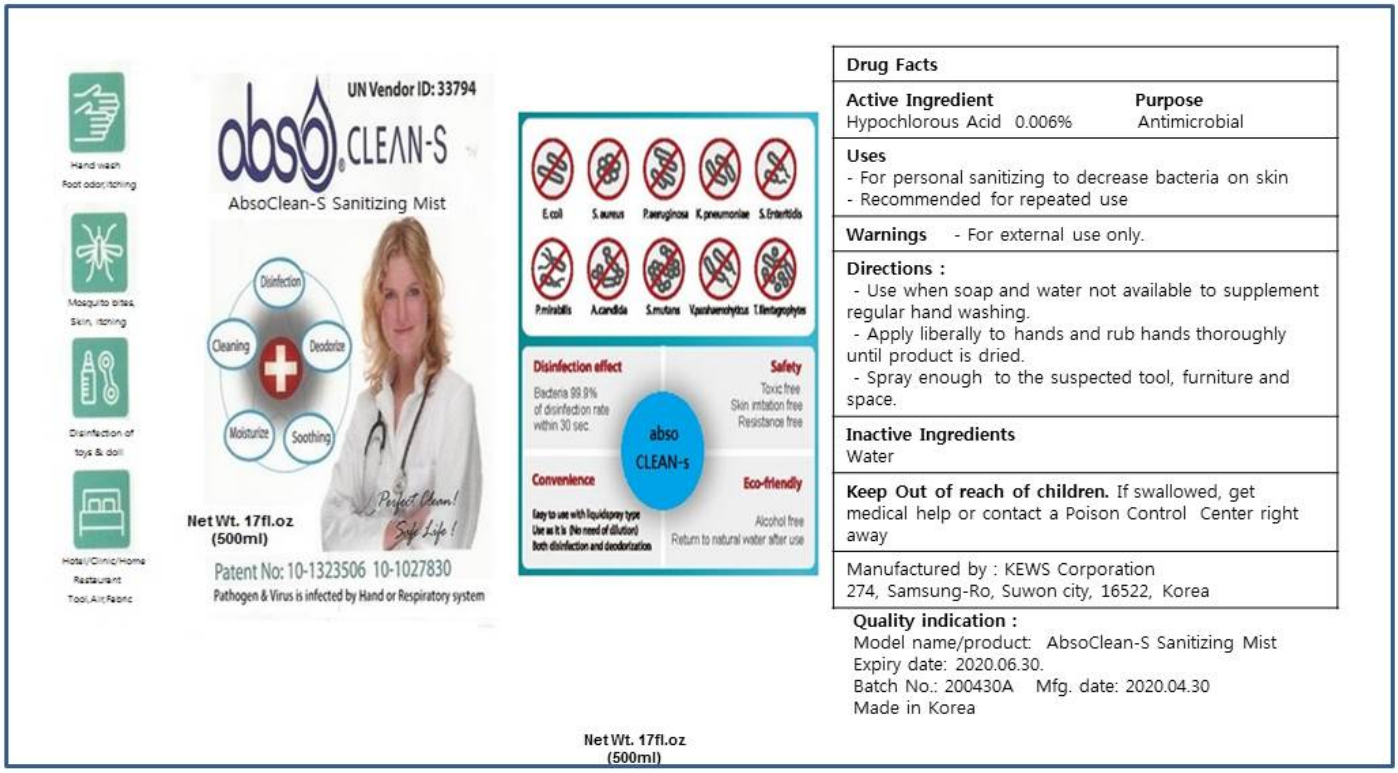
AbsoClean-S Sanitizing Mist 100ml



500mL PDP

500mL NDC 75644-002-02

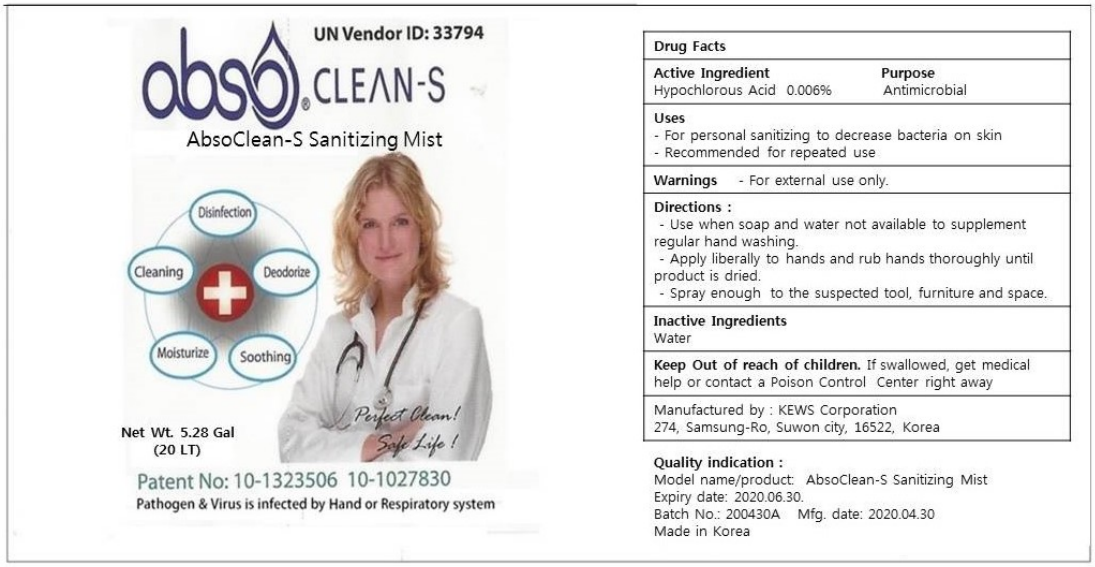
AbsoClean-S Sanitizing Mist 500ml



20LT PDP

20LT NDC 75644-002-03

AbsoClean-S Sanitizing Mist 20 LT



ABSOCLEAN-S SANITIZING MIST

hypochlorous acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75644-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYPOCHLOROUS ACID (UNII: 712K4CDC10) (HYPOCHLOROUS ACID - UNII:712K4CDC10)	HYPOCHLOROUS ACID	0.006 mg in 1 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:75644-002-01	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/30/2020	
2	NDC:75644-002-02	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/30/2020	
3	NDC:75644-002-03	20000 mL in 1 JUG; Type 0: Not a Combination Product	04/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/30/2020	

Labeler - KEWS CORPORATION (688640317)

Registrant - KEWS CORPORATION (688640317)

Establishment

Name	Address	ID/FEI	Business Operations
KEWS CORPORATION		688640317	manufacture(75644-002) , pack(75644-002) , label(75644-002)