HANNA DISINFECTANTSPRAY- sodium hypochlorite liquid
Rainbow Co Ltd

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.

Drug Facts

sodium hypochlorite

WATER, peg-60 hydrogenated castor oil, aloe extract, camellia sinensis leaf extract, grapefruit seed extract, fragrance

Sterilization of hands and skin

KEEP OUT OF REACH OF THE CHILDREN

Apply an appropriate amount on your hands and rub well to dry.

1. Do not use on the following body parts. A wide range of body parts and damaged skin around the eyes and ears, in the oral cavity (may have irritating effects)
2. If the following symptoms appear, stop using them immediately and consult a doctor or pharmacist.
1) Hypersensitivity symptoms such as rash, erythema, itching, and edema
2) Skin irritation symptoms

for external use only

HANNA DISINFECTANTSPRAY

sodium hypochlorite liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74247-0014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM HYPOCHLORITE (UNII: DY38VHM5OD) (HYPOCHLORITE ION - UNII:T5UM7HB19N)	SODIUM HYPOCHLORITE	0.02 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
GRAPEFRUIT PEEL (UNII: 3582N05Q44)	
PEG-60 HYDROGENATED CASTOR OIL (UNII: 02NG325BQG)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74247-0014-1	55 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
2	NDC:74247-0014-2	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
3	NDC:74247-0014-3	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
4	NDC:74247-0014-4	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
5	NDC:74247-0014-5	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/13/2020	

Labeler - Rainbow Co Ltd (690423720)

Registrant - Rainbow Co Ltd (690423720)

Establishment

Name	Address	ID/FEI	Business Operations
Rainbow Co Ltd		690423720	manufacture(74247-0014)

Revised: 6/2020 Rainbow Co Ltd