HYDROGEN PEROXIDE- hydrogen peroxide solution
Target Corp

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.

UP & UP
871.001-871AA
Drug Facts

Active ingredient

Hydrogen peroxide (stablized) 3%

Purpose

First Aid Antiseptic

Use

first aid to help prevent the risk of infection in minor cuts, scrapes and burns

Warnings

For external use only

Do not use

- in the eyes or apply over large areas of the body
- longer than one week

Ask a doctor before use if you have

deep or puncture wounds, animal bites or serious burns

Keep out of reach of children.

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

Directions

- clean the affected area
- spray a small amount of product on the affected area 1 to 3 times a day
- may be covered with a sterile bandage
- if bandaged, let dry first

Other information

keep tightly closed and at controlled room temperature. Do not shake bottle. Hold sprayer tip away from face when opening.

Inactive ingredients

purified water

Dist. by Target Corp., Mpls., MN 55403

Made in U.S.A. with U.S. and foreign components

Questions? Call 1-800-910-6874

principal display panel

Hydrogen

peroxide

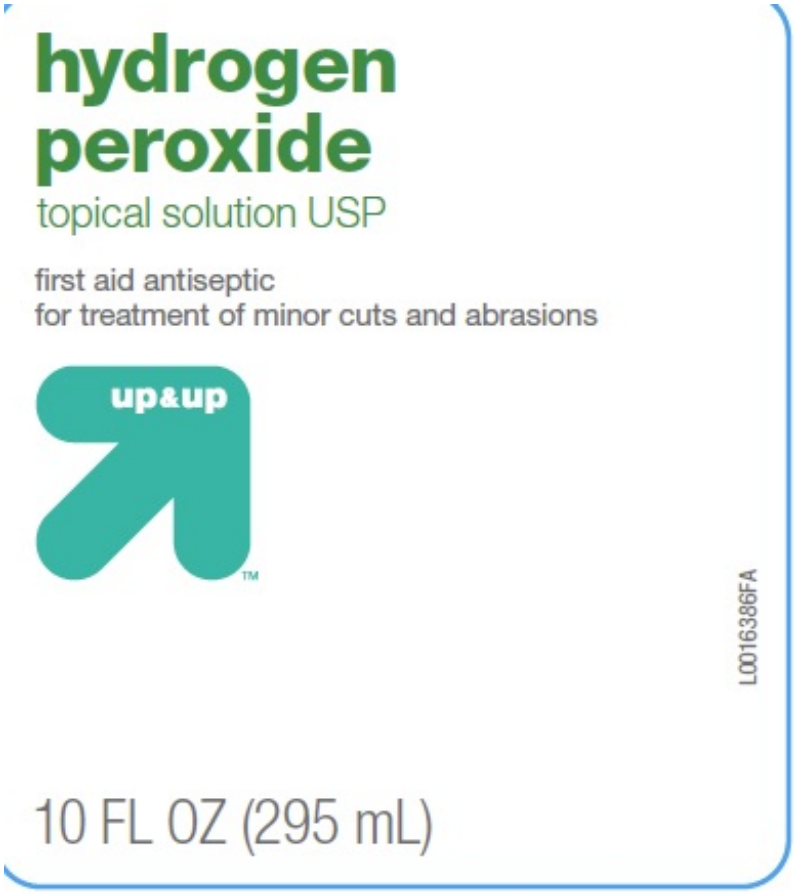
topical solution USP

first aid antiseptic

for treatment of minor cuts and abrasions

up & up

10 FL OZ (295 mL)



HYDROGEN PEROXIDE

hydrogen peroxide solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-871
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROGEN PEROXIDE (UNII: BBX060AN9V) (HYDROGEN PEROXIDE - UNII:BBX060AN9V)	HYDROGEN PEROXIDE	30 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:11673-	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a	07/15/1989	

1	871-43	Combination Product	07/15/1989	
2	NDC:11673-871-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1989	
3	NDC:11673-871-72	295 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/15/1989	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/1989	

Labeler - Target Corp (006961700)

Registrant - Vi Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(11673-871)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(11673-871)

Revised: 5/2022 Target Corp