HYDROGEN PEROXIDE- hydrogen peroxide solution
Chain Drug Marketing Association

Quality Choice 871.001/871AA
Hydrogen Peroxide

Active ingredient

Hydrogen peroxide (stabilized) 3%

Purpose

First aid antiseptic

Uses

- first aid to help prevent the rise 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 1 week

Ask a doctor before use if

you have deep or puncture wounds, animal bites or serious burns

Stop use and ask a doctor if

• the condition persists or gets worse

Keep out of reach of children.

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

Directions

- clean the affected area
- apply 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 away from face when opening.

Inactive ingredient

purified water

Adverse reaction

Distributed by CDMA

Novi, MI 48375

www.qualitychoice.com

Questions: 800-935-2362

principal display panel

NDC 83324-176-10

QC ®

QUALITY CHOICE

Topical Solution USP

Hydrogen Peroxide

First Aid Antiseptic

For Treatment of Minor Cuts and Abrasions

10 FL OZ (236 mL)

HYDROGEN PEROXIDE

hydrogen peroxide solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-176
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:83324-176-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/02/2024	
2	NDC:83324-176-32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/02/2024	
3	NDC:83324-176-10	295 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/29/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	07/02/2024	

Labeler - Chain Drug Marketing Association (011920774)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(83324-176)

Revised: 2/2026 Chain Drug Marketing Association