

HYDROGEN PEROXIDE- hydrogen peroxide solution
CVS Pharmacy, Inc

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.

CVS 871.001-871AA Hydrogen Peroxide

Active ingredient

Hydrogen peroxide (stabilized) 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

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

Distributed by: CVS Pharmacy, Inc

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principal display panel

CVS Health

Topical Solution USP

HYDROGEN PEROXIDE

FIRST AID ANTISEPTIC

For treatment of minor cuts + abrasions

8 FL OZ (236 mL)



Topical Solution USP

Hydrogen Peroxide

FIRST AID ANTISEPTIC

For treatment of minor cuts & abrasions



8 FL OZ (236 mL)

L0012354FC

HYDROGEN PEROXIDE

hydrogen peroxide solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-871
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
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1	NDC:59779-871-34	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	
2	NDC:59779-871-43	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	
3	NDC:59779-871-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	
4	NDC:59779-871-50	710 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/1992	
5	NDC:59779-871-99	237 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/15/1992	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/1992	

Labeler - CVS Pharmacy, Inc (062312574)

Registrant - Vi Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(59779-871)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(59779-871)

Revised: 10/2023

CVS Pharmacy, Inc