

HYDROGEN PEROXIDE- hydrogen peroxide solution
Vi-Jon, LLC

Swan Hydrogen Peroxide
871.001/871AA

Active ingredient

Hydrogen peroxide (stabilized) 3%

Purpose

First Aid Antiseptic

Uses

- 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
- sore mouth symptoms do not improve in 7 days
- irritation, pain or redness persists or worsens
- swelling, rash, or fever develops

Keep out of reach of children.

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

Directions

For First aid antiseptic:

- 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 away from face when opening.

Inactive ingredient

purified water

Adverse reaction

Distributed by: Vi-Jon, LLC.,

8515 Page Ave.

St. Louis, MO 63114

Principal Display Panel

Swan

Hydrogen Peroxide

Topical Solution USP

- First Aid Antiseptic/Oral Debriding Agent
- For treatment of minor cuts & abrasions

6 FL OZ (177 mL)



HYDROGEN PEROXIDE SPRAY

Topical Solution USP

- ▶ First Aid Antiseptic
- ▶ For Treatment of Minor Cuts & Abrasions

6 FL OZ (177 mL)

HYDROGEN PEROXIDE

hydrogen peroxide solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0869-0871
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:0869-0871-72	295 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/09/2017	
2	NDC:0869-0871-43	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/09/2017	
3	NDC:0869-0871-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/09/2017	
4	NDC:0869-0871-18	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/09/2017	04/12/2021
5	NDC:0869-0871-99	237 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/09/2017	
6	NDC:0869-0871-82	177 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/09/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	07/09/2017	

Labeler - Vi-Jon, LLC (088520668)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(0869-0871)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(0869-0871)

Revised: 4/2024

Vi-Jon, LLC