# 3% HYDROGEN PEROXIDE- hydrogen peroxide liquid LJ Pharma

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## **Active Ingredient**

Hydrogen Peroxide (Stabilized) 3%

## **Purpose**

FIRST AID ANTISEPTIC ORAL DEBRIDING AGENT

#### Uses

First aid to help prevent the risk of infection in minor cuts, Scrapes & Burns. Aids in the removal of phlegm, mucus or other secretions associated with occasional sore mouth

## Warnings

- For external use only
- IF taken internally, Serious gastric disturbances will result. **Flammable**, **Keep away** from fire or flame.
- Use only in a well-ventilated area: Fumes may be toxic.

**Ask a doctor before use if you have**deep or puncture wounds, Animal bites or serious burns.

## When using this product:

- Do not Apply over large areas of the body.
- Do not use longer than one week unless directed by doctor

Stop use and ask doctor if conditions persist or get worse

**Keep out of reach of children** In case of ingestion, get medical help or contact a poison. control center right away.

#### **Directions**

- Clean the affected Area
- Apply Small amount of this product on the area 1-3 times daily
- Let the alcohol dry on skin before using bandages.
- May be covered with a sterile bandag.

#### Other information

- Store ate room temperature
- Do not shake bottle Hold away from face when opening.
- Does not contain, nor is intended as a substitute for grain or ethyl alchol.

## **Inactive Ingredients**

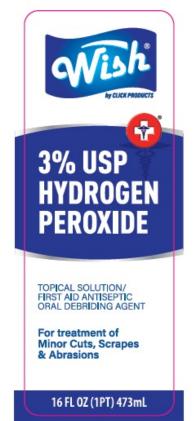
**Purified Water** 

#### **Product label**



48 x 130 mm







60 x 65 mm





## **3% HYDROGEN PEROXIDE**

hydrogen peroxide liquid

## **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:84809-005

Route of Administration TOPICAL

### **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
HYDROGEN PEROXIDE (UNII: BBX060AN9V) (HYDROGEN PEROXIDE - UNII: BBX060AN9V)	HYDROGEN PEROXIDE	3 mL in 100 mL

## **Inactive Ingredients**

Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84809- 005-01	12 in 1 CARTON	06/30/2025	
1		236 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
2	NDC:84809- 005-02	12 in 1 CARTON	06/30/2025	
2		473 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
3	NDC:84809- 005-03	12 in 1 CARTON	08/04/2025	
3		946 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/30/2025	

## **Labeler -** LJ Pharma (914078633)

Establishment				
Name	Address	ID/FEI	Business Operations	
LJ Pharma		914078633	manufacture(84809-005)	

Revised: 8/2025 LJ Pharma