PHARMACYS PRESCRIPTION HYDROGEN PEROXIDE- hydrogen peroxide liquid American Consumer Products 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.

Pharmacy's Prescription 16OZ Hydrogen Peroxide

Active Ingredient

Active ingredient (by volume)

Hydrogen peroxide 3%

Purpose

First aid antiseptic/oral debriding agent.

Warnings

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For external use only.

Stop Use

Stop use and ask a doctor if: the condition persists or gets worse, sore mouth symptoms do not improve within 7 days, irritation, pain or redness persists or worsens, swelling, rash or fever develops.

Keep out of reach of children

Keep this and all drugs out of the reach of children. If swallowed, seek professional or contact a Poison Control Center immediately.

Directions

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First aid antiseptic:

Clean affected area; apply small amount of product on affected area 1-3 times a day; may be covered with a sterile bandage; if bandage, let dry first.

Oral debriding agent (oral rinse):

Adults and children 2 years of age and over:

Mix with an equal amount of water; swish around in the mouth over affected area for at least 1 minute and then spit out; use up to 4 times daily after meals and at bedtime or as directed by a dentist or doctor.

Inactive Ingredients

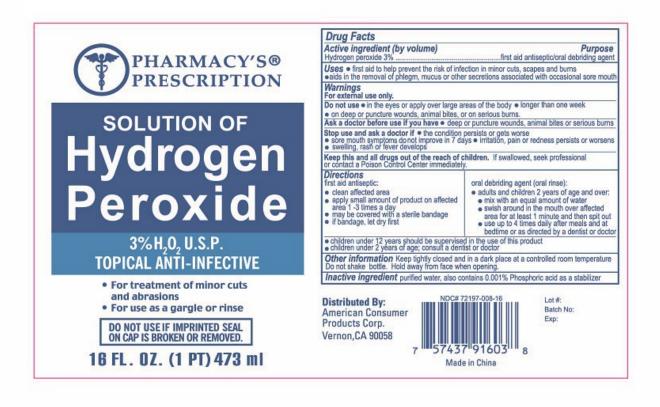
Inactive ingredient

Purified water, also contains 0.001% Phosphoric acid as a stabilizer.

Indications & Usage Section

Do not use

In the eyes or apply over large areas of the body; longer than one week; on deep or puncture wounds, animal bites, or on serious burns.



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hydrogen peroxide liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72197-008	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
HYDROGEN PEROXIDE (UNII: BBX060AN9V) (HYDROGEN PEROXIDE - UNII:BBX060AN9V)	HYDRO GEN PERO XIDE	3 mL in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHO SPHO RIC ACID (UNII: E4GA8884NN)	

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:72197-008- 16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/31/2018		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	08/31/2018		

 ${f Labeler}$ - American Consumer Products Corp (081101181)

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