PEROX-A-MINT SOLUTION- hydrogen peroxide mouthwash
Sage Products, LLC

Perox-A-Mint® Solution

Active ingredient

Hydrogen Peroxide 1.5%

Purpose

Oral Debriding Agent

Use

Aids in the removal of secretions associated with sore mouth

Warnings

Do not use:

- For more than 7 days unless told to do so by a doctor.
- Avoid swallowing.

Stop use and ask a doctor if:

- Sore mouth symptoms do not improve in 7 days.
- Swelling, rash or fever develops.
- Irritation, pain or redness persists of worsens.

Keep out of reach of children. If more than used for debriding is accidentally swallowed, get medical help or contact a Poison Control Center right away.

Directions

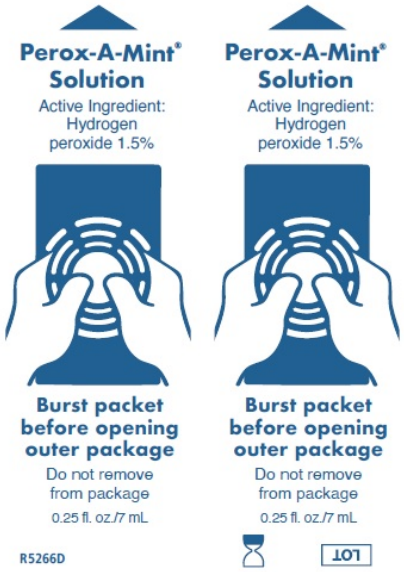
- Use up to 4 times daily or as directed by a dentist or doctor.
- Adults and children 3 years and over: rinse mouth for approx. one minute with 1/2 Tbsp. Instruct to expectorate.
- Children under 12 years of age: supervise use.
- Children under 3 years of age: consult a dentist or doctor.

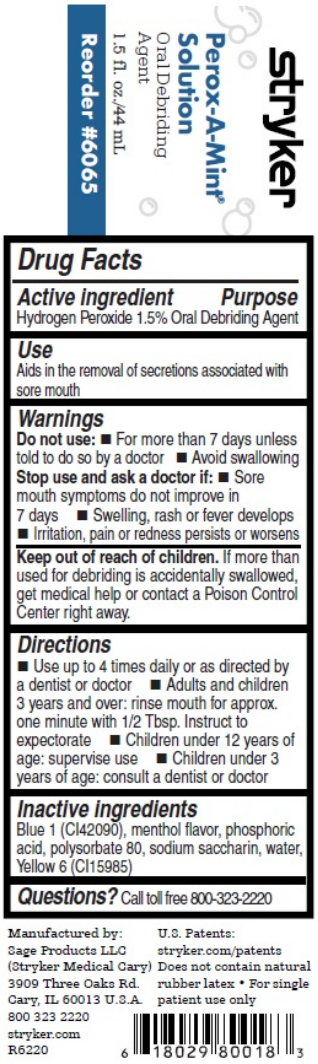
Inactive Ingredients

Blue 1 (CI42090), menthol flavor, phosphoric acid, polysorbate 80, sodium saccharin, Water, Yellow 6 (CI15985)

Questions?Call toll free 800-323-2220

Perox-A-Mint Solution





PEROX-A-MINT SOLUTION

hydrogen peroxide mouthwash

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53462-075
Route of Administration	BUCCAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROGEN PEROXIDE (UNII: BBX060AN9V) (HYDROGEN PEROXIDE - UNII:BBX060AN9V)	HYDROGEN PEROXIDE	15 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	

POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53462-075-30	44 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/21/1998	
2	NDC:53462-075-40	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/21/1998	08/20/2018
3	NDC:53462-075-60	7 mL in 1 PACKET; Type 0: Not a Combination Product	05/21/1998	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	05/21/1998	

Labeler - Sage Products, LLC (054326178)

Registrant - Sage Products, LLC (054326178)

Revised: 6/2026 Sage Products, LLC