MEDLINE PEROXI FRESH MOUTHWASH- hydrogen peroxide mouthwash
Medline Industries, LP

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086 Medline Peroxi Fresh

Active ingredient

Hydrogen Peroxide 1.5%

Purpose

Oral Debriding Agent

Use

- aids in the removal of phlegm, mucus, or other secretions associated with occasional sore mouth

Warnings

Do not use this product

- for more than 7 days unless directed by a dentist or physician.

When using this product

- do not swallow

Stop use and ask a doctor if

- 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 more than used for rinsing is accidentally swallowed, get medical help or contact a poison Control Center right away.

Directions

- Adults and Children 2 Years of Age or Over: Swish around in 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.
- Children Under 12 Years of Age: Should be supervised in the use of this product. Children Under 2 Years of Age: Consult a dentist or a doctor.

Other information

- avoid excessive heat and protect from freezing

Inactive ingredients

flavor, sodium lauryl sulfate, sodium saccharin, water

Manufacturing Information

Manufactured for:

Medline Industries, LP

Three Lakes Drive, Northfield, IL 60093 USA

Made in USA

www.medline.com

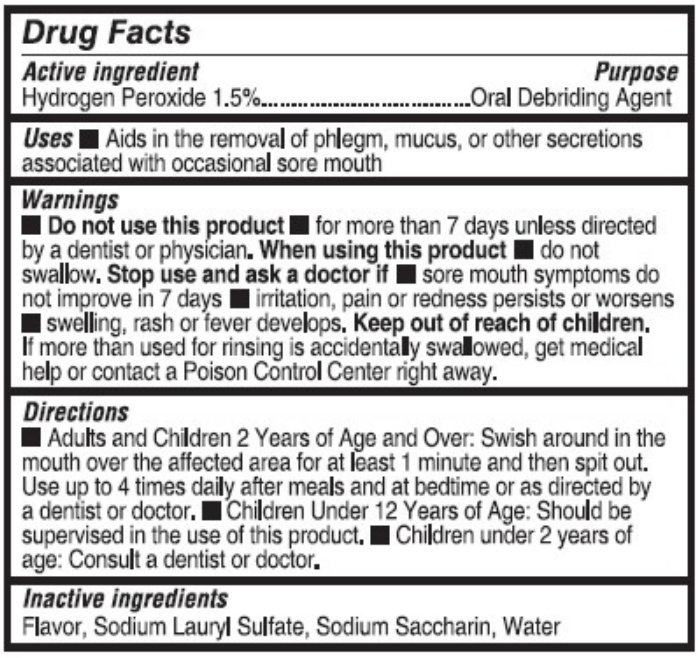
1-800-MEDLINE (633-5463)

REF: MDS096065HP

V1 RB22ANO

Package Label

Label



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MEDLINE PEROXI FRESH MOUTHWASH

hydrogen peroxide mouthwash

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53329-086
Route of Administration	DENTAL		

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
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-086-13	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2007	12/31/2017
2	NDC:53329-086-66	10 mL in 1 PACKET; Type 0: Not a Combination Product	01/01/2007	07/31/2021
3	NDC:53329-086-14	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/08/2017	
4	NDC:53329-086-95	7 mL in 1 PACKET; Type 0: Not a Combination Product	01/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	01/01/2007	

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 8/2026 Medline Industries, LP