

HYDROGEN PEROXIDE- hydrogen peroxide solution
NDC, Inc.

Hydrogen Peroxide 3%

Drug Facts

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 and burns
- aids in removal of phlegm, mucous, or other secretions associated with occasional sore mouth

Warnings

For external use only

Do not use

- in the eyes or apply over large areas of the body
- longer than 1 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

**Oral Debriding Agent (oral
rinse):**

Adults and children 2 years of

First Aid Antiseptic

- clean the affected area
- apply a small amount of product on the area 1 to 3 times a day
- may be covered with a sterile bandage
- if bandaged, let dry first

age & over:

- mix with an equal amount of water
- swish around in the mouth over the 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

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- 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 doctor

Other Information

- Keep tightly closed in a cool dark place
- Do not shake bottle
- Hold away from face when opening

Inactive Ingredient

Purified Water

PRINCIPAL DISPLAY PANEL - 473 mL Bottle Label

REF: P903016

NDC 43128-001-16

Hydrogen

Peroxide

3% USP

Tamper evident cap for your protection. If ring-band is detached from cap or missing do not use.

16oz. (473mL)

Made in USA for:

NDC, Inc., 407 New Sanford Rd.

La Vergne, TN 37086

www.ProAdvantagebyNDC.com

Pro

ADVANTAGE®

by **NDC**

REF: **P903016**
NDC 43128-001-16



Hydrogen Peroxide 3% USP

Tamper evident cap for your protection. If ring-band is detached from cap or missing do not use.

Agua Oxigenada 3% USP

Manipulación evidente casquillo para su protección. No utilice banda anillo o si está separado de la tapa.

Made in the USA for:
Fabricado en Estados Unidos para:
NDC, Inc., 407 New Sanford Rd.
La Vergne, TN 37086
www.ProAdvantagebyNDC.com

16 fl. oz. (473 mL)

1-07-D0012N05F

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Directions

First Aid Antiseptic • clean the affected area • apply a small amount of product on the area 1 to 3 times a day • may be covered with a sterile bandage • if bandaged, let dry first

Oral Debriding Agent (oral rinse) Adults and children 2 years of age & over: • mix with an equal amount of water • swish around in the mouth over the 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 doctor

Other information • Keep tightly closed in a cool dark place • Do not shake bottle • Hold away from face when opening

Inactive Ingredient Purified Water

1-07-D0012N05B



M220P9030168X

HYDROGEN PEROXIDE

hydrogen peroxide solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43128-001
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
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WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43128-001-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2014	
2	NDC:43128-001-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/01/2014	

Labeler - NDC, Inc. (009831413)

Revised: 11/2023

NDC, Inc.