

HYDROGEN PEROXIDE- hydrogen peroxide liquid

Medical Chemical Corporation

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.

Hydrogen Peroxide 3% Label

Usage: Intended for use as a general purpose topical antiseptic for first-aid treatment of minor cuts and abrasions. In case of deep or puncture wounds consult a physician. Discontinue use and get medical attention if redness, irritation or infection develops.

Active ingredient: hydrogen peroxide, 3% w/v in water.

For External Use Only. Do not use in eyes or over large areas of the body. Keep out of the reach of children.

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For oral hygiene, dilute with an equal volume of water and gargle for about one minute. Rinse with water. In case of accidental ingestion contact a physician or a poison control center.

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h2o2label.jpg

Manufactured by Medical
Chemical Corp. Torrance,
CA 90501

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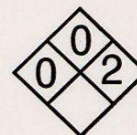
Hydrogen Peroxide, 3%

Lot #: Size:
Expiration:

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Catalog# 1202E
NDC 012745-1202



HYDROGEN PEROXIDE

hydrogen peroxide liquid

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:12745-202 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|---------------------|
| HYDROGEN PEROXIDE (UNII: BBX060AN9V) (HYDROGEN PEROXIDE - UNII:BBX060AN9V) | HYDROGEN PEROXIDE | 8.57 g in 100 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---------------------------------|----------|
| WATER (UNII: 059QF0KO0R) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:12745-202-01 | 59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 07/21/1980 | |

| | | | | |
|----------|------------------|---|------------|--|
| 2 | NDC:12745-202-02 | 118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 07/21/1980 | |
| 3 | NDC:12745-202-03 | 3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 07/21/1980 | |
| 4 | NDC:12745-202-04 | 208175 mL in 1 DRUM; Type 0: Not a Combination Product | 07/21/1980 | |
| 5 | NDC:12745-202-05 | 946250 mL in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK; Type 0: Not a Combination Product | 07/21/1980 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part333A | 07/21/1980 | |

Labeler - Medical Chemical Corporation (008496861)

Establishment

| Name | Address | ID/FEI | Business Operations |
|------------------------------|---------|-----------|------------------------|
| Medical Chemical Corporation | | 008496861 | manufacture(12745-202) |

Revised: 1/2022

Medical Chemical Corporation