

DAB- benzalkonium chloride solution

Three Kings 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.

Active Ingredient

Benzalkonium Chloride 0.12%

Purpose

Antiseptic

Uses

- For handwashing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

- **For external use only**

When using this product

- Do not use in or near eyes

Stop use and ask a doctor if irritation or redness develop.

Keep out of reach of children. If swallowed, seek medical attention or contact Poison Control Center immediately.

Directions

- receive a small amount of foam into palm of hand
- rub thoroughly over all surfaces of both hands
- rub hands together until dry

Other information

- Store in a cool dry place below 104 F (40 C)

Inactive ingredients

Water, cocamidopropyl betaine, glycerin, poloxamer 124, phenoxyethanol, benzyl alcohol, dehydroacetic acid, sodium hydroxide, citric acid, D&C Green No. 8

Questions?

Contact Three Kings Corp. at 888-872-2350 Mon-Fri 9 am - 4pm CST

dab

HAND PROTECTANT + SANITIZER


MICROBIOLOGICAL DEFENSE SPECTRUM

1.2L (1.26 QT) 40.5 GL OZ

daproducts.com/healthcare

| | |
|--|----------------|
| Drug Facts | |
| Active Ingredient | Purpose |
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Manufactured for:
Three Kings Corp
Corinth, MS 38834
1-888-872-2350



105-08806-001
8 60000 59135 5



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DAB

benzalkonium chloride solution

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:72898-001 |
| Route of Administration | TOPICAL | | |

| Active Ingredient/Active Moiety | | | | |
|--|------------------|---|--------------------------|--------------------|
| | | Ingredient Name | Basis of Strength | Strength |
| | | BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 0.12 g in 100 mL |
| Inactive Ingredients | | | | |
| | | Ingredient Name | Strength | |
| | | GLYCERIN (UNII: PDC6A3C0OX) | | |
| | | PHENOXYETHANOL (UNII: HIE492ZZ3T) | | |
| | | DEHYDROACETIC ACID (UNII: 2KAG279R6R) | | |
| | | CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) | | |
| | | BENZYL ALCOHOL (UNII: LKG8494WBH) | | |
| | | SODIUM HYDROXIDE (UNII: 55X04QC32I) | | |
| | | WATER (UNII: 059QF0K00R) | | |
| | | COCAMIDOPROPYL BETAINE (UNII: 5OCF3011KX) | | |
| | | POLOXAMER 124 (UNII: 1S66E28KXA) | | |
| | | D&C GREEN NO. 8 (UNII: I2W85YOX9L) | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:72898-001-01 | 45 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product | 02/10/2020 | |
| 2 | NDC:72898-001-02 | 250 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product | 02/10/2020 | |
| 3 | NDC:72898-001-03 | 1200 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product | 02/10/2020 | |
| 4 | NDC:72898-001-04 | 946352 mL in 1 DRUM; Type 0: Not a Combination Product | 02/10/2020 | |
| Marketing Information | | | | |
| Marketing Category | | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph not final | | part333A | 02/10/2020 | |

Labeler - Three Kings Corp (080906833)

Registrant - Best Sanitizers, Inc (627278224)

Establishment

| Name | Address | ID/FEI | Business Operations |
|----------------------|---------|-----------|------------------------|
| Best Sanitizers, Inc | | 627278224 | manufacture(72898-001) |