

GREEN GUARD ANTISEPTIC WIPES- benzalkonium chloride swab
GREEN GUARD ANTIDEPTIC WIPES- benzalkonium chloride swab
Uniforst First Aid Corporation

Green Guard Antiseptic Wipes

Drug Facts

Active ingredient

Benzalkonium chloride 0.133% w/v

Purpose

First aid antiseptic

Uses

First aid to help prevent infection in minor cuts, scrapes and burns.

Warnings

Do not use

- in the eyes or apply over large area of the body
- in case of deep puncture wounds, animal bites or serious burns, consult a doctor
- longer than 1 week unless directed by a doctor

Stop use and ask a doctor if

- the condition persists or gets worse

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- tear open packet, unfold and clean the affected area
- discard after use
- may be covered with a sterile bandage
- if bandaged, let dry first

Other information

- store at room temperature 59°-86°F (15°-30°C)
- tamper evident sealed packets
- do not use any open or torn packets

Inactive ingredients

sodium bicarbonate, water

Questions or comments?

1-800-869-6970

Green Guard Antiseptic Wipes Label

Antiseptic wipes

First aid to help prevent infection

in minor cuts, scrapes and burns

Benzalkonium chloride 0.133% w/v

First Aid Antiseptic

Green Guard®

25 Extra Large Wipes Per Box

Order # 3025

| | | | | | | | | | | | | | | | | | | |
|--|--|--|---|--|---|--|---|--|--|--|---|--|--|--|---|--|---|--|
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Green Guard Antiseptic Wipes Extra Large Label

Antiseptic wipes

First aid to help prevent infection

in minor cuts, scrapes and burns

Green Guard®

25 Extra Large Wipes Per Box

Order # 3025

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|---|---|--|--|--|---|--|---|--|---|--|--|--|---|--|--|--|---|--|
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GREEN GUARD ANTISEPTIC WIPES

benzalkonium chloride swab

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:47682-045 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-----------------------|-------------------|
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 1.3 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| SODIUM BICARBONATE (UNII: 8MDF5V39QO) | |
| WATER (UNII: 059QF0KO0R) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:47682-045-71 | 20 in 1 BOX | 08/21/2017 | 08/21/2017 |
| 1 | | 1.4 mL in 1 PACKET; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M003 | 08/21/2017 | |

GREEN GUARD ANTIDEPTIC WIPES

benzalkonium chloride swab

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:47682-055 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|-----------------|-------------------|----------|
|-----------------|-------------------|----------|

| | | |
|--|-----------------------|-------------------|
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 1.3 mg in 1 mL |
|--|-----------------------|-------------------|

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| SODIUM BICARBONATE (UNII: 8MDF5V39QO) | |
| WATER (UNII: 059QF0KO0R) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:47682-055-71 | 20 in 1 BOX | 08/21/2017 | 08/22/2017 |
| 1 | NDC:47682-055-99 | 2.1 mL in 1 PACKET; Type 0: Not a Combination Product | | |
| 2 | NDC:47682-055-73 | 25 in 1 BOX | 08/21/2017 | |
| 2 | | 2.1 mL in 1 PACKET; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M003 | 08/21/2017 | |

Labeler - Uniforst First Aid Corporation (832947092)

Revised: 9/2025

Uniforst First Aid Corporation