

SCRUB- chlorhexidine gluconate solution
Bajaj Medical, LLC

Drug Facts

Active ingredient

chlorhexidine gluconate 0.75% solution

Purposes

Antisepetic

Uses

- **healthcare personnel handwash:** helps reduce bacteria that potentially can cause disease

Warnings

For external use only.

Do not use

- if you are allergic to chlorhexidine gluconate or any other ingredients
- in contact with meninges
- in the genital area
- as a preoperative skin preparation (especially on the head and face)
- on skin wounds
- general skin cleanser as surgical hand scrub

When using this product

- keep out of eyes, ears, and mouth. May cause serious and permanent eye injury if placed or kept in the eye during surgical procedures or may cause deafness when instilled in the middle ear through perforated eardrums.
- if solution should contact these areas, rinse out promptly and thoroughly with water

Stop use and ask a doctor if

irritation, sensitization or allergic reaction occurs. These may be signs of a serious condition.

Keep out of reach of children

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

Other safety information

This product contains a chemical known to the State of California to cause Cancer.

Directions

- use with care in premature infants and infants under 2 months of age. These products may cause irritation or chemical burns.

Healthcare personnel handwash:

- wet hands with water

- dispense about 5 ml of product into cupped hands and wash in a vigorous manner for 30 seconds
- rinse and dry thoroughly

Other information

- store at 20-25°C (68-77°F)
- avoid excessive heat above 40°C (104°F)

Inactive ingredients

cocamide DEA, FD&C yellow #5, FD&C red #4, fragrance, hydroxyethylcellulose, isopropyl alcohol, lauramine oxide, purified water, ricinoleamidopropyl trimethyl ammonium chloride

Questions of Comments?

Call 1-855-332-2525 Monday through Friday 7:00 AM to 3:30 PM

Package/Label Principal Display Panel

NDC 61037-412-01

SCRUB™

FROM HOSPITAL TO HOME

Antiseptic Handwash

Chlorhexidine Gluconate 0.75% Solution

FDA Approved Antiseptic Handwash

Contains: 0.75% Chlorhexidine Gluconate

Distributed by: Bajaj Medical, LLC

415 W. Pershing Rd.,

Chicago, IL 60609

FOR EXTERNAL USE ONLY

Net Contents: 4 fl oz (118 ml)

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Lift Here

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SCRUB

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Product Information

| | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:61037-412 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------------|------------------|
| CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L) | CHLORHEXIDINE GLUCONATE | .75 mg in 100 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| COCO DIETHANOLAMIDE (UNII: 92005F972D) | |
| HYDROXYETHYL CELLULOSE (140 CPS AT 5%) (UNII: 8136Y38GY5) | |
| ISOPROPYL ALCOHOL (UNII: ND2M416302) | |
| LAURAMINE OXIDE (UNII: 4F6FC4M18W) | |
| RICINOLEAMIDOPROPYLTRIMONIUM CHLORIDE (UNII: 93OU7D1C3U) | |
| WATER (UNII: 059QF0K00R) | |
| CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) | |
| FD&C YELLOW NO. 5 (UNII: I753WB2F1M) | |
| FD&C RED NO. 4 (UNII: X3W0AM1JLX) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|-----------------------------|----------------------|--------------------|
| 1 | NDC:61037-412-01 | 118 mL in 1 BOTTLE, PLASTIC | | |
| 2 | NDC:61037-412-02 | 237 mL in 1 BOTTLE, PLASTIC | | |
| 3 | NDC:61037-412-03 | 473 mL in 1 BOTTLE, PLASTIC | | |
| 4 | NDC:61037-412-04 | 946 mL in 1 BOTTLE, PLASTIC | | |
| 5 | NDC:61037-412-05 | 3785 mL in 1 JUG | | |
| 6 | NDC:61037-412-06 | 237 mL in 1 BOTTLE, PUMP | | |
| 7 | NDC:61037-412-07 | 237 mL in 1 BOTTLE, PUMP | | |
| 8 | NDC:61037-412-10 | 60 mL in 1 BOTTLE, PUMP | | |
| 9 | NDC:61037-412-11 | 60 mL in 1 BOTTLE, PLASTIC | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| NDA | NDA020111 | 09/30/2014 | |

Labeler - Bajaj Medical, LLC (078774921)

Registrant - Bajaj Medical, LLC (078774921)

Establishment

| Name | Address | ID/FEI | Business Operations |
|--------------------|---------|-----------|------------------------|
| Bajaj Medical, LLC | | 078774921 | manufacture(61037-412) |

Revised: 10/2014

Bajaj Medical, LLC