BP WASH- benzoyl peroxide emulsion CINTEX SERVICES, LLC

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.

BP Wash 10% (benzoyl peroxide 10%)

Active ingredient

Benzoyl peroxide 10%

Purpose

Acne treatment

Use

For the treatment of acne

Warnings

For external use only

Do not use if you

- have very sensitive skin
- are sensitive to benzoyl peroxide

When using this product

- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.
- avoid unnecessary sun exposure and use a sunscreen
- avoid contact with the eyes, lips, and mouth
- avoid contact with hair and dyed fabrics, which may be bleached by this product
- skin irritation may occur, characterized by redness, burning, itching, peeling, or possibly swelling. Irritation may be reduced by using the product less frequently or in a lower concentration.

Stop use and ask a doctor if irritation becomes severe.

KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.

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

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer and rinse thoroughly one to three times daily
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day.
- if going outside, apply sunscreen after using this product. If irritation or sensitivity develops, stop

use of both products and ask a doctor.

Other information

Store at 20°C to 25°C (68°F to 77°F), excursions permitted between 15°C and 30°C (between 59°F and 86°F).

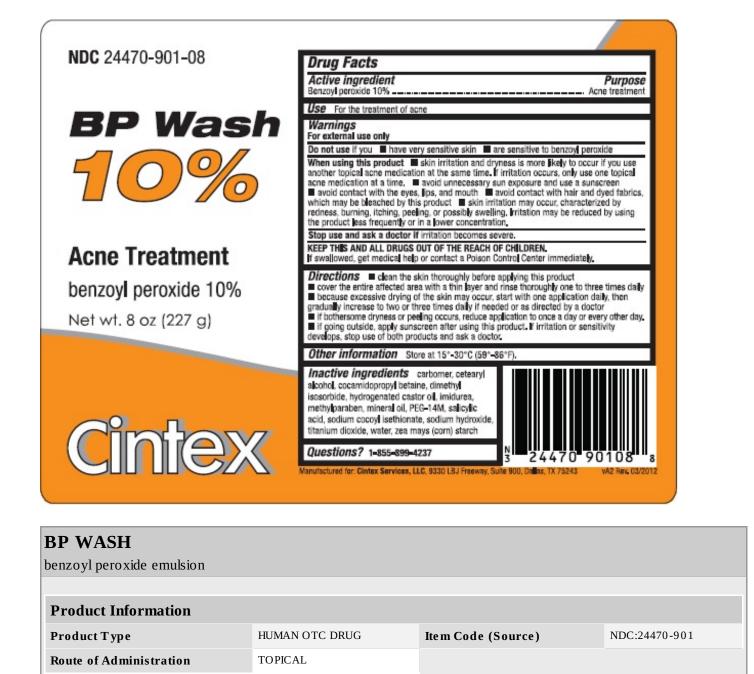
Inactive ingredients

carbomer, cetearyl alcohol, cocamidopropyl betaine, dimethyl isosorbide, hydrogenated castor oil, imidurea, methylparaben, mineral oil, PEG-14M, salicylic acid, sodium cocyl isethionate, sodium hydroxide, titanium dioxide, water, zea mays (corn) starch

Questions?

1-855-899-4237





| Active Ingredient/Active Moiety | | | | | | | |
|---|-------------------|---------------------|--|--|--|--|--|
| Ingredient Name | Basis of Strength | Strength | | | | | |
| BENZOYL PEROXIDE (UNII: W9 WZN9 A0 GM) (BENZOYL PEROXIDE - UNII:W9 WZN9 A0 GM) | BENZOYL PEROXIDE | $100\ mg$ in $1\ g$ | | | | | |
| BENZOYL PEROXIDE (UNII: W9 WZN9A0GM) (BENZOYL PEROXIDE - UNII: W9 WZN9A0GM) | BENZOYL PEROXIDE | 100 mg in | | | | | |

| Inactive Ingredients | | | | |
|--|----------|--|--|--|
| Ingredient Name | Strength | | | |
| COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX) | | | | |
| MINERAL OIL (UNII: T5L8T28FGP) | | | | |
| POLYETHYLENE OXIDE 600000 (UNII: 2126FD486L) | | | | |
| HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY) | | | | |
| WATER (UNII: 059QF0KO0R) | | | | |
| CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) | | | | |
| IMIDUREA (UNII: M629807ATL) | | | | |

| STARCH, CORN (UNII: |) (28232NY3SJ | | | | | |
|---|--|-------------------|----------------------|----------------------|-----------|----------|
| CARBOXYPOLYMETH | | A5MM307FC) | | | | |
| DIMETHYL ISOSORBI | DE (UNII: SA6A | 6V432S) | | | | |
| METHYLPARABEN (UN | (II: A2I8C7HI9T) | • | | | | |
| SALICYLIC ACID (UNII | | | | | | |
| SO DIUM HYDRO XIDE (| UNII: 55X04QC | 32I) | | | | |
| TITANIUM DIO XIDE (U | NII: 15FIX9V2JF | 2) | | | | |
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| | | | | | | |
| Product Character | ristics | | | | | |
| Color | | white | Score | | | |
| Shape | | | Size | | | |
| Flavor | | | Imprint Code | | | |
| Contains | | | | | | |
| | | | | | | |
| | | | | | | |
| Packaging | | | | | | |
| # Item Code | Package Description | | Marketing Start Date | Marketing | End Date | |
| 1 NDC:24470-901-05 | 42 g in 1 BOTTLE; Type 0: Not a Combination Product | | 0 1/15/20 12 | | | |
| | 227 g in 1 BOTTLE; Type 0: Not a Combination Product | | | 0 1/15/20 12 | | |
| 2 NDC:24470-901-08 2 | | | | | | |
| 2 NDC:24470-901-08 2 | | | | | | |
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| | rmation | | | | | |
| Marketing Info | | | | | | |
| 2 NDC:24470-901-08 2 Marketing Info Marketing Category OTC monograph final | | n Number or Monog | raph Citation | Marketing Start Date | Marketing | End Date |

Labeler - CINTEX SERVICES, LLC (078304114)

Revised: 12/2017

CINTEX SERVICES, LLC