HUMANE ACNE WASH- benzoly peroxide cream
Humane Consumer, LLC

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humane acne wash

Drug Facts

Active Ingredient

Benzoyl Peroxide 10%

Purpose

Acne Treatment

use

for the treatment of acne.

Warnings

For external use only.

Do not use

- If you are sensitive to Benzoyl Peroxide or have very sensitive skin.
- Using other topical acne drugs at the same time or right after use of the product may increase dryness, redness or irritation of the skin.
- Because excessive drying of the skin may occur, start with one application daily, then gradually increase to two to three times daily if neede. If bothersome dryness or peeling occurs, reduce application to once per day.
- Do not use for prolonged periods without contacting a doctor.

When using this product

- Avoid contact with eyes. of contact occurs, flush thoroughly with water. Keep away from lips and mouth.
- Avoid unnecessary sun exposure and use a sunscreen.
- Avoid Product contact with hair and dyed fabrics, including carpets and clothing which may be bleached by this Product.

Stop use and ask a doctor

if excessive irritation occurs.

Keep out of reach of children.

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

Directions

Cleanse Skin thoroughly before applying medication. Apply even layer over affected area. Rinse well. DO not use on broken skin or large parts of the body. For best results, apply one to three times per day.

Inactive ingredients

Aqua (Deionized Water), Carbomer, Cocamidopropyl Betaine, Gluconolactone, Sodium Benzoate, Sodium Hydroxide.

Product Label

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HUMANE ACNE WASH

benzoly peroxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69919-021
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69919-021-01	1 in 1 CARTON	02/08/2017	
1		237 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	05/28/2015	

Labeler - Humane Consumer, LLC (079845933)

Revised: 11/2023 Humane Consumer, LLC