HUMANE ACNE WASH- benzoyl peroxide liquid
Apprendista, LLC

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 this 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 needed. If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

When using this product

- Avoid contact with and near eyes. If 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.

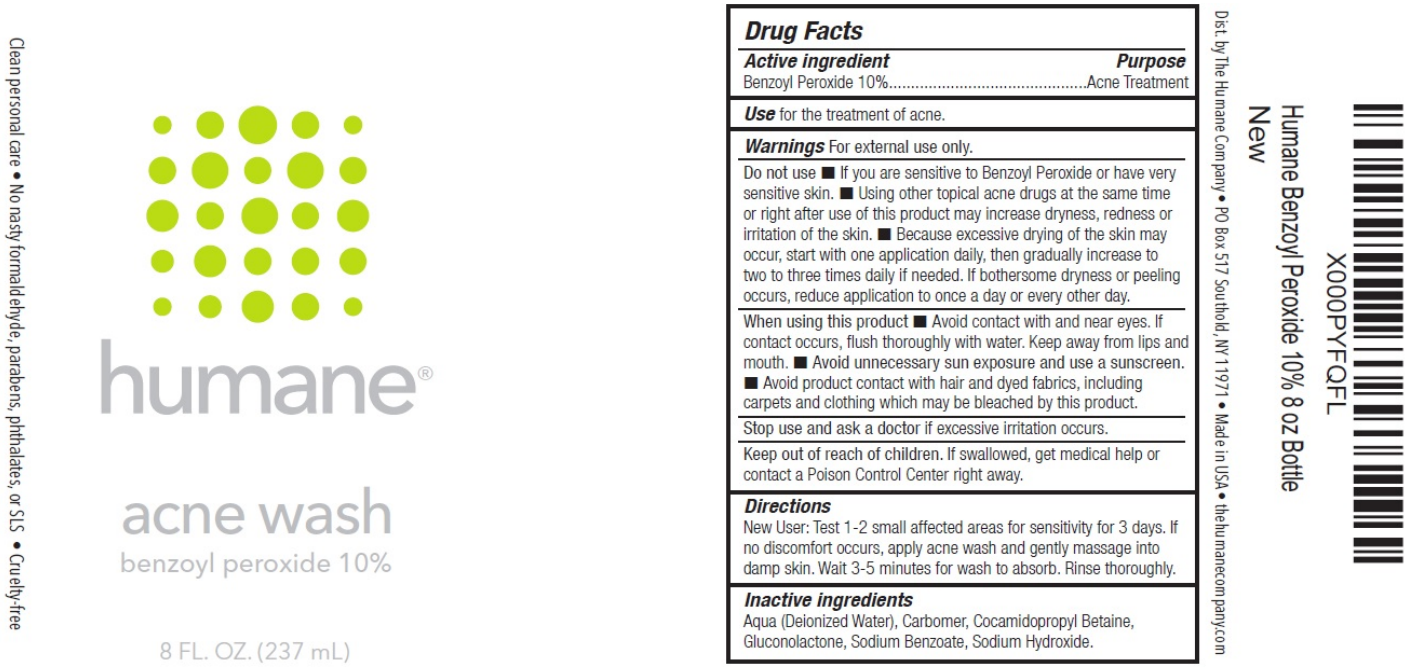
Directions

New User: Test 1-2 small affected areas for sensitivity for 3 days. If no discomfort occurs, apply acne wash and gently massage into damp skin. Wait 3-5 minutes for wash to absorb. Rinse thoroughly.

Inactive ingredients

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

Package Labeling:



HUMANE ACNE WASH

benzoyl peroxide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73010-900
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)	

CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
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:73010-900-00	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	04/01/2019	

Labeler - Apprendista, LLC (116995922)

Revised: 11/2023 Apprendista, LLC