MEDPURA BENZOYL PEROXIDE- benzoyl peroxide liquid
Akron Pharma Inc.

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.

MEDPURA 5% and 10% Benzoyl Peroxide
Acne Treatment Wash

Drug Facts

Active ingredient

Benzoyl peroxide 5%

Benzoyl peroxide 10%

Purpose

Acne treatment

Use

for the treatment of acne

Warnings

For external use only

Do not use this medication if you have very sensitive skin or if you are sensitive to benzoyl peroxide.

When using this product

- avoid unnecessary sun exposure and use a sunscreen.
- avoid contact with eyes, lips, and mouth, eyelids and mucous membranes.
- avoid contact with hair or 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.
- 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.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

- Shake well
- Sensitivity Test for a New User. Apply product sparingly to one or two small affected areas during the first 3 days. If no discomfort occurs, follow the directions stated below.
- wet area to be cleansed
- apply acne wash and gently massage area for 1-2 minutes
- rinse thoroughly and pat dry
- 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 controlled room temperature, 15°- 30°C (59°- 86°F)

Inactive ingredients

citric acid, edetate disodium, germall 115, glycerin, phophomer x polymer, purified water, sodium olefin sulfonate, sodium hydroxide

Questions or comments?

call toll-free 1-877-225-6999

Manufactured for:

Akron Pharma, Inc.
Fairfield, NJ-07004
www.akronpharma.com

Drug Facts

Active Ingredient	Purpose
Benzoyl peroxide USP, 5%	Acne medication

Use for the treatment of acne

Warnings

For external use only

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Stop use and ask a doctor if ■ irritation becomes severe.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions ■ shake well

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Other information

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Inactive ingredients

citric acid, carbopol 940, edetate disodium, germall 115, glycerin, purified water, sodium olefin sulfonate, sodium hydroxide

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Akron Pharma
Manufactured for:
Akron Pharma, Inc.
Fairfield, NJ-07004
www.akronpharma.com
Rev.:11/21

NDC 71399-9234-5

MEDPURA™

Benzoyl Peroxide

5%

TOPICAL WASH

Acne medication

For Topical Use

Made in USA

NET WT. (5 Oz) 148g

Drug Facts

Active Ingredient	Purpose
Benzoyl peroxide USP, 5%	Acne medication

Use for the treatment of acne

Warnings

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Akron Pharma
Manufactured for:
Akron Pharma, Inc.
Fairfield, NJ-07004
www.akronpharma.com
Rev.:11/21

NDC 71399-9234-8

MEDPURA™

Benzoyl Peroxide

5%

TOPICAL WASH

Acne medication

For Topical Use

Made in USA

NET WT. (8 Oz) 237g

Drug Facts

Active Ingredient — **Purpose**

Benzoyl peroxide USP, 10% ..... Acne medication

Use for the treatment of acne

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Inactive ingredients

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N 3 71399 92365 3

Akron Pharma
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Akron Pharma, Inc.
Fairfield, NJ-07004
www.akronpharma.com
Rev.:11/21

NDC 71399-9236-5

MEDPURA™

Benzoyl Peroxide

10%

TOPICAL WASH

Acne medication

For Topical Use

Made in USA

NET WT. (5 Oz) 148g

Drug Facts

Active Ingredient — **Purpose**

Benzoyl peroxide USP, 10% ..... Acne medication

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

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Rev.:11/21

NDC 71399-9236-8

MEDPURA™

Benzoyl Peroxide

10%

TOPICAL WASH

Acne medication

For Topical Use

Made in USA

NET WT. (8 Oz) 237g

MEDPURA BENZOYL PEROXIDE

benzoyl peroxide liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71399-9234
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Benzoyl Peroxide (UNII: W9WZN9A0GM) (Benzoyl Peroxide - UNII:W9WZN9A0GM)	Benzoyl Peroxide	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
DOCUSATE SODIUM (UNII: F05Q2T2JA0)	
ALCOHOL (UNII: 3K9958V90M)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
2-METHACRYLOYLOXYETHYL PHOSPHORYLCHOLINE (UNII: 59RU860S8D)	
water (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-9234-5	148 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/04/2021	
2	NDC:71399-9234-8	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/04/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part333D	11/04/2021	

MEDPURA BENZOYL PEROXIDE

benzoyl peroxide liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71399-9236
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
DOCUSATE SODIUM (UNII: F05Q2T2JA0)	
ALCOHOL (UNII: 3K9958V90M)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
2-METHACRYLOYLOXYETHYL PHOSPHORYLCHOLINE (UNII: 59RU860S8D)	
water (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-9236-5	148 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/04/2021	
2	NDC:71399-9236-8	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/04/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part333D	11/04/2021	

Labeler - Akron Pharma Inc. (067878881)

Revised: 3/2023 Akron Pharma Inc.