

ZO SKIN HEALTH ACNE CONTROL BENZOYL PEROXIDE- benzoyl peroxide cream
ZO Skin Health, Inc.

ZO[®] Skin Health Acne Control 10% Benzoyl Peroxide

National Drug Code 42851-087-60

Drug Facts

Active Ingredient

Benzoyl Peroxide 10%

Purpose

Acne Medication

Uses

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 out of reach of children.

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

Directions

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.

- Clean the skin thoroughly before applying this product.
- Cover the entire affected area with a thin layer 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

- Protect this product from excessive heat and direct sun.

Inactive Ingredients

Aqua/Water/Eau, Aspalathus Linearis Leaf Extract, Camellia Sinensis Leaf Extract, Carbomer, Diethylhexyl Sodium Sulfosuccinate, Dimethicone, Disodium EDTA, Ethylhexylglycerin, Glycerin, Melaleuca Alternifolia (Tea Tree) Leaf Oil, PEG-40 Stearate, PEG-8 Laurate, Phenoxyethanol, Pinus Strobus Bark Extract, Polyglyceryl-6 Isostearate, Propanediol, Propylene Glycol, Silica, Sodium Citrate, Sodium Hydroxide, Sorbitan Stearate, Vitis Vinifera (Grape) Seed Extract, Xanthan Gum.

Dist. by ZO Skin Health, Inc. Irvine, CA 92618

PRINCIPAL DISPLAY PANEL - 60 mL Tube Carton

ZO[®] SKIN HEALTH
BY ZEIN OBAGI MD

NDC 42851-087-60

ACNE CONTROL
10% Benzoyl Peroxide

60 mL / 2 Fl. Oz.



ACNE

ZO[®] SKIN HEALTH

BY ZEIN OBAGI MD

NDC 42851-087-60



ACNE CONTROL

10% Benzoyl Peroxide

60 mL / 2 Fl. Oz.

Maximum-strength acne-fighting formula that controls surface oil, neutralizes bacteria, and prevents future breakouts.

Drug Facts (Continued)

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Dist. by ZO Skin Health, Inc. Irvine, CA 92618
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ZO SKIN HEALTH ACNE CONTROL BENZOYL PEROXIDE

benzoyl peroxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42851-087
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)	
ASPALATHUS LINEARIS LEAF (UNII: H7UGK1GJCU)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
DOCUSATE SODIUM (UNII: F05Q2T2JA0)	
DIMETHICONE (UNII: 92RU3N3Y10)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
TEA TREE OIL (UNII: VIF565UC2G)	
PEG-40 STEARATE (UNII: ECU18C66Q7)	
PEG-8 LAURATE (UNII: 762O8IWA10)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PINUS STROBUS BARK (UNII: 8PJ4761097)	
POLYGLYCERYL-6 ISOSTEARATE (UNII: 5VKGO1003G)	
PROPANEDIOL (UNII: 5965N8W85T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
VITIS VINIFERA SEED (UNII: C34U15ICXA)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42851-087-60	1 in 1 CARTON	05/25/2023	
1		60 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	05/25/2023	

Labeler - ZO Skin Health, Inc. (826468527)

Revised: 10/2024

ZO Skin Health, Inc.