SIGNATURE CARE ACNE MEDICATION- benzoyl peroxide gel SAFEWAY, 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.

SIGNATURE CARE 10% BENZOYL PEROXIDE ACNE MEDICATION

Active ingredient

Benzoyl Peroxide 10%

Purpose

Acne Medication

Uses

for the treatment of acne

Warnings

IFor external use only

Do not use if you • have very sensitive skin • are sensitive to benzoyl peroxide

When using this product

- avoid contact with the eyes, lips, and mouth
- avoid unnecessary sun exposure and use a sunscreen
- 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.
- 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.

□Stop use and ask a doctor if□ • irritation becomes severe

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

Directions

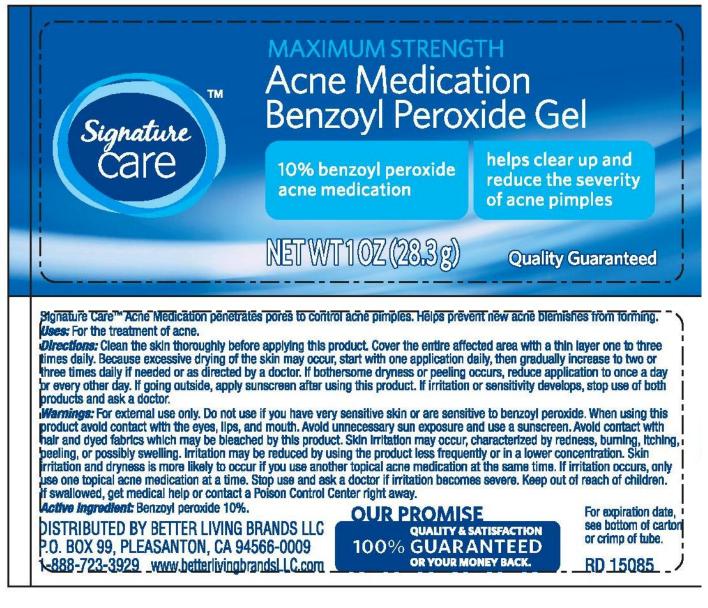
- Clean the skin thoroughly before applying this product
- Cover the entire affected area with a thin layer one to three times daily
- Becasue excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- 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.

Inactive ingredients

carbomer, disodium EDTA, hydroxypropyl methylcellulose, laureth-4, sodium hydroxide, water FACE VALUES ACNE MEDICATION 10% BENZOYL PEROXIDE GEL

1 OZ (28.3g)

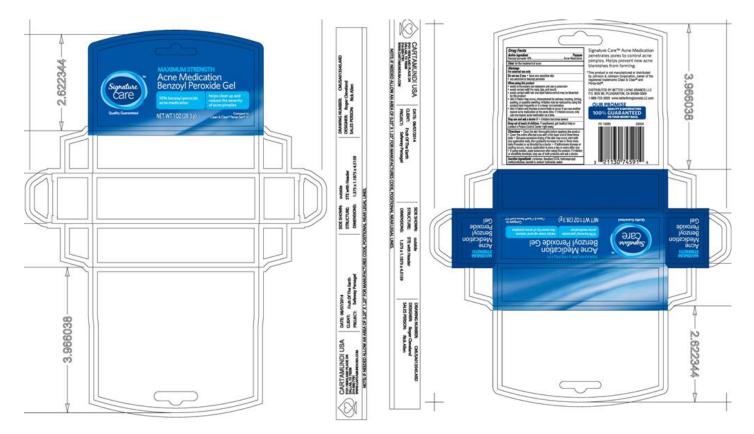
NDC 21130-730-16



FACE VALUES ACNE MEDICATION 10% BENZOYL PEROXIDE GEL

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NDC 21130-730-17



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P	roduct Informa	tion					
Pr	oduct T ype		HUMAN OTC DRUG	Item Code (Source)	N	NDC:21130	-730
Ro	ute of Administra	ition	TOPICAL				
Ad	tive Ingredien	t/Active Moi	ety				
		In	igredient Name		Basis of Str	rength	Strength
BE	NZOYL PEROXID	E (UNII: W9 WZN9	A0GM) (BENZOYL PEROXIDE ·	- UNII:W9WZN9A0GM)	BENZOYL PER	ROXIDE	100 mg in 1 §
In	active Ingredie	ents					
			Ingredient Name				Strength
CA	RBOMER INTERP	OLYMER TYPE	A (ALLYL SUCROSE CROSSI	L INKED) (UNII: 59TL3W	/G5CO)		Strength
CA ED	RBOMER INTERP(ETATE DISODIUM	OLYMER TYPE I (UNII: 7FLD91C8	A (ALLYL SUCROSE CROSSL 36K)	. INKED) (UNII: 59TL3W	/G5CO)		Strength
CA ED HY	RBOMER INTERP ETATE DISODIUM PROMELLOSE 29	OLYMER TYPE [(UNII: 7FLD91C8 10 (4000 MPA.S	A (ALLYL SUCROSE CROSSI	L INKED) (UNII: 59TL3W	/G5CO)		Strength
CA ED HY LA	RBOMER INTERP ETATE DISODIUM PROMELLOSE 29 URETH-4 (UNII: 6 H	OLYMER TYPE [(UNII: 7FLD91C8 10 (4000 MPA.S [Q855798J]	A (ALLYL SUCROSE CROSSL 36K) 5) (UNII: RN3152OP35)	. INKED) (UNII: 59TL3W	/G5CO)		Strength
CA ED HY LA SO	RBOMER INTERP ETATE DISODIUM PROMELLOSE 29	OLYMER TYPE [(UNII: 7FLD91C8 10 (4000 MPA.S [Q855798J) E (UNII: 55X04Q0	A (ALLYL SUCROSE CROSSL 36K) 5) (UNII: RN3152OP35)	L INKED) (UNII: 59TL3W	/G5CO)		Strength
CA ED HY LA SO	RBOMER INTERP ETATE DISODIUM PROMELLOSE 29 URETH-4 (UNII: 6 H DIUM HYDROXID)	OLYMER TYPE [(UNII: 7FLD91C8 10 (4000 MPA.S [Q855798J) E (UNII: 55X04Q0	A (ALLYL SUCROSE CROSSL 36K) 5) (UNII: RN3152OP35)	J INKED) (UNII: 59TL3W	/G5CO)		Strength
CA ED HY LA SO	RBOMER INTERP ETATE DISODIUM PROMELLOSE 29 URETH-4 (UNII: 6 H DIUM HYDROXID)	OLYMER TYPE [(UNII: 7FLD91C8 10 (4000 MPA.S [Q855798J) E (UNII: 55X04Q0	A (ALLYL SUCROSE CROSSL 36K) 5) (UNII: RN3152OP35)	J INKED) (UNII: 59TL3W	/G5CO)		Strength
CA ED HY LA SO W	RBOMER INTERP ETATE DISODIUM PROMELLOSE 29 URETH-4 (UNII: 6 H DIUM HYDROXID)	OLYMER TYPE [(UNII: 7FLD91C8 10 (4000 MPA.S [Q855798J) E (UNII: 55X04Q0	A (ALLYL SUCROSE CROSSL 36K) 5) (UNII: RN3152OP35)	. INKED) (UNII: 59TL3W	/G5CO)		Strength
CA ED HY LA SO W	RBOMER INTERP ETATE DISODIUM PROMELLOSE 29 URETH-4 (UNII: 6 H DIUM HYDROXIDI ATER (UNII: 059QF	OLYMER TYPE [(UNII: 7FLD91C8 10 (4000 MPA.S [Q855798J] E (UNII: 55X04Q6 DKO0R)	A (ALLYL SUCROSE CROSSL 36K) 5) (UNII: RN3152OP35)	LINKED) (UNII: 59TL3W		Marketin	Strength

1 NDC:21130-730-16	28.3 g 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 final	part333D	12/08/2014							
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Labeler - SAFEWAY, INC. (009137209)

Registrant - FRUIT OF THE EARTH, INC. (079559467)

Establishment

Name	Address	ID/FEI	Business Operations
FRUIT OF THE EARTH, INC.		080086802	manufacture(21130-730)

Revised: 5/2020

SAFEWAY, INC.