

PAULAS CHOICE CLEAR DAILY SKIN CLEARING TREATMENT- benzoyl peroxide lotion
Paula's Choice, LLC

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.

Paulas Choice Clear Daily Skin Clearing Treatment 2.5% Benzoyl Peroxide Extra Strength

Benzoyl Peroxide 5%

Acne Treatment

- Treats acne.
- Helps prevent new acne blemishes.
- Penetrates pores to reduce acne blemishes.

After cleansing with Paula's Choice CLEAR Pore Normalizing Cleanser and exfoliating with Anti-Redness Exfoliating Solution, cover the affected area with a thin layer 1 to 3 times a day. Because excessive drying of the skin may occur, start with 1 application daily and then gradually increase to 2 or 3 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, use a sunscreen. If sensitivity develops, discontinue use.

For external use only.

Do not use · on broken skin · on large areas of the body · if you are sensitive to benzoyl peroxide.

If too much skin irritation or skin sensitivity develops or increases.

· Avoid unnecessary sun exposure and use a sunscreen · Avoid contact with eyes, lips and mouth. If contact occurs, rinse with water · This product may bleach hair or dyed fabrics · Using other topical acne products at the same time or right after use of this product may increase dryness or irritation of the skin. If this occurs, only one drug should be used unless directed by a doctor.

If swallowed, get medical help and contact a Poison Control Center immediately.

If too much skin irritation or skin sensitivity develops or increases.

Store at 20-26°C (68-77°F). · You may report serious adverse reactions to 1030 SW 34th Street, Suite A, Renton, WA 98057.

Water, Propylene Glycol, Bisabolol, Allantoin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Hydroxyethylcellulose, Sodium Citrate, Laureth-4, Caprylyl Glycol, Hexylene Glycol, Sodium Hydroxide, Disodium EDTA, Phenoxyethanol.

PAULA'S CHOICE
Clear Daily Skin Clearing Treatment, Extra Strength, Travel Size
Item #: 6117
Part #: 162109
Version Code: r417
Artwork Created: 11/15/17
Tube: molded PMS 2198C, shiny
Shoulder: molded PMS 2198C, shiny
Cap: molded natural, shiny

INKS: 3



WHITE



BLACK



PMS 301C

Artwork that appears blue is white ink.

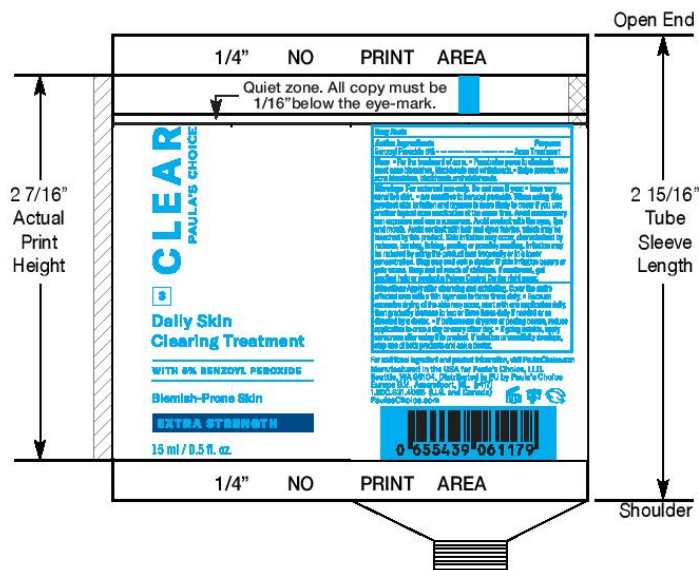
CCL TUBE	Tube Size: 1" (D) x 2 15/16" (H)		GENERAL NOTES
	Drawing No: 1X2-15-16-DP	Created: 1/31/14 RR	
	Shoulder Margin: 1/4"	Revisions: N/A	
	Open End Margin: 1/4"		
	Eye-mark Size: 1/8" x 1/4"		
	Scale: 100%		
	Die Line Type: Direct Print		



1/8" bleed for overlap
only if full wrap is desired



1/8" no copy area



FYI DIELINE HAS BEEN FLIPPED

Artwork that appears blue is white ink.

GENERAL NOTES		Tube Size: 1 3/16" (D) x 5 3/8" (H)
<ul style="list-style-type: none"> Measurements are in inches. All copy must be 1/16" below the eye-mark. Please include a PDF, list of colors, and convert text to outlines when submitting artwork. 	Revisions: 1/31/14 RR Revisions: N/A	1-3-16X5 Die Line No: Shoulder Margin: 1/4" Open End Margin: 1/4" Eye-mark Size: 1/8" x 1/4" Scale: 100% Die Line Type: Direct Print



PAULA'S CHOICE
 Clear Daily Skin Clearing Treatment, Extra Strength
 Item #: 6110
 Part #: 163268
 IL version: il62v1
 Formula #: 20TC(MH)070A
 Version Code: r417
 Artwork Created: 11/12/17, Revised 1/12/18

Tube: molded PMS 2198, matte
 Shoulder: molded PMS 2198, shiny
 Cap: Molded natural, shiny

INKS: 3



WHITE



BLACK



PMS 301C

PAULAS CHOICE CLEAR DAILY SKIN CLEARING TREATMENT

benzoyl peroxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76144-611
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
LEVOMENOL (UNII: 24WE03BX2T)	
ALLANTOIN (UNII: 344S277G0Z)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
HYDROXYETHYL CELLULOSE (5000 CPS AT 1%) (UNII: X70SE62ZAR)	
LAURETH-4 (UNII: 6HQ855798J)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76144-611-01	67 mL in 1 TUBE; Type 0: Not a Combination Product	09/07/2012	
2	NDC:76144-611-02	15 mL in 1 TUBE; Type 0: Not a Combination Product	09/07/2012	
3	NDC:76144-611-03	2 mL in 1 PACKET; Type 0: Not a Combination Product	09/07/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	09/07/2012	

Labeler - Paula's Choice, LLC (029583981)

Registrant - Paula's Choice, LLC (029583981)

Establishment

Name	Address	ID/FEI	Business Operations
Thibiant International, Inc.		083913913	manufacture(76144-611)

Revised: 1/2022

Paula's Choice, LLC