

**OBLITERATE - benzoyl peroxide lotion
mybody**

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.

OBLITERATE

ACTIVE INGREDIENT BENZOYL PEROXIDE 5%

PURPOSE ACNE TREATMENT

USE FOR TREATMENT OF ACNE

WARNINGS

FOR EXTERNAL USE ONLY

DO NOT USE * IF YOU HAVE VERY SENSITIVE SKIN OR ARE SENSITIVE TO BENZOYL PEROXIDE.

ASK A DOCTOR OR PHARMACIST BEFORE USE IF * PREGNANT OR LACTATING. YOU ARE USING OTHER TOPICAL ACNE MEDICATIONS AT THE SAME TIME OR IMMEDIATELY FOLLOWING USE OF THIS PRODUCT. THIS MAY INCREASE DRYNESS OR IRRITATION OF THE SKIN. IF THIS OCCURS, ONLY ONE MEDICATION SHOULD BE USED UNLESS DIRECTED BY A DOCTOR.

WHEN USING THIS PRODUCT * AVOID CONTACT WITH EYES. IF CONTACT OCCURS, FLUSH THOROUGHLY WITH WATER. * KEEP AWAY FROM LIPS AND MOUTH.

KEEP OUT OF REACH OF CHILDREN * IF SWALLOWED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

DIRECTIONS

* AM: APPLY SIDE 1 BENZOYL PEROXIDE TO CLEAN SKIN.

PM: APPLY SIDE 1 BENZOYL PEROXIDE TO CLEAN SKIN. ALLOW TO DRY. NEXT APPLY SIDE 2 WITH RETINOL. ALLOW TO DRY. DAILY USE OF SUNSCREEN IS RECOMMENDED. * IF BOTHERSOME DRYNESS OR PEELING OCCURS, REDUCE APPLICATION TO ONCE EVERY OTHER DAY.

OTHER INFORMATION

* AVOID STORAGE AT EXTREME TEMPERATURES (BELOW 40°F AND ABOVE 100°F).

INACTIVE INGREDIENTS CHAMBER A: WATER, POLYACRYLAMIDE, C13-14 ISOPARAFFIN, LAURETH-7, ACETYL CARBOXYMETHYL COCOYL GLYCINE, PHENOXYETHANOL, CAPRYLYL GLYCOL, DECYLENE GLYCOL, CARBOMER.

QUESTIONS OR COMMENTS?

CALL 877.423.1314 * WWW.LOVEMYBODY.COM

MYBODY

OBLITERATE

BOOSTER PACK: ACTIVE ACNE

MYBOOSTER * STEP 2



OBLITERATE®

2-in-1 Extreme
Refining Solution
5% Benzoyl Peroxide

mytreatment
STEP 2 • AM / PM

ACNE

1 FL OZ US / 30 mL

Maximum-strength multi-active solution effectively combines 5% Benzoyl Peroxide with myR2X™ Enhanced Retinol Complex to help obliterate acne on contact.

ACTIVE INGREDIENT: Benzoyl Peroxide 5%
PURPOSE: Acne treatment

Application and Usage

Intended for moderate to severe acne.

AM: Apply **SIDE 1** Benzoyl Peroxide to clean skin. PM: Apply **SIDE 1** Benzoyl Peroxide to clean skin. Allow to dry. Next apply **SIDE 2** with Retinol. Allow to dry. Daily use of sunscreen is recommended.

WARNINGS: For external use only. Do not use with another acne medication except on the advice of a doctor. If excessive skin irritation develops, stop use. If irritation persists, consult a doctor. When using this product, avoid unnecessary sun exposure and use sunscreen. Keep out of reach of children.

Distributed by mybody®, LLC, Phoenix, AZ 85018
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www.loveyourbody.com

Drug Facts for OBLITERATE

Active Ingredient Benzoyl Peroxide 5%
Purpose Acne Treatment

USE for treatment of acne

Warnings

For external use only

Do not use • if you have very sensitive skin or are sensitive to Benzoyl Peroxide.

Ask a doctor or pharmacist before use if • Pregnant or lactating. You are using other topical acne medications at the same time or immediately following use of this product. This may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.

When using this product • Avoid contact with eyes. If contact occurs, flush thoroughly with water. • Keep away from lips and mouth.

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

Directions

• AM: Apply **SIDE 1** Benzoyl Peroxide to clean skin.
PM: Apply **SIDE 1** Benzoyl Peroxide to clean skin. Allow to dry. Next apply **SIDE 2** with Retinol. Allow to dry. Daily use of sunscreen is recommended. • If bothersome dryness or peeling occurs, reduce application to once every other day.

Other Information

• Avoid storage at extreme temperatures (below 40° F and above 100° F).

Inactive Ingredients CHAMBER A: Water, Polyacrylamide, C13-14 Isoparaffin, Laureth-7, Acetyl Carboxymethyl Cocoyl Glycine, Phenoxyethanol, Caprylyl Glycol, Deoxyene Glycol, Carbomer.

Questions or Comments?

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Inactive Ingredients CHAMBER B: Water, Glycerin, Glyceryl Stearate SE, C 12-15 Alkyl Benzoate, Carthamus Tinctorius (Safflower) Seed Oil, Butyrospermum Perka (Shea Butter), Dimethicone, Cetyl Alcohol, Butylene glycol, Sucrose Diaurate, Retinol, Glyceryl Disteoate, Retinyl Palmitate, Myristoyl Nonapeptide-3, Camellia Sinensis Leaf Extract, Caffeine, Undecylenyl Phenylalanine, Pterisatium (Pea) Extract, Tocopheryl Acetate, Allantoin, Panthenol, Safflower Acids, Cocoglycosides, Safflower Glycosides, Lysolcithin, Acetyl Carboxymethyl Cocoyl Glycine, Polysorbate 20, Cetearyl Alcohol, Ceteareth-20, Acrylate/C10-30 Alkyl Acrylate Crosspolymer, Xanthan Gum, Citric Acid, Polyglyceryl-10 Laurate, Polysorbate 80, Dimethylmethoxy Chromanyl Palmitate, Steareth-21, Steareth-20, Cetyl Dimethicone, Disodium EDTA, BHT, Tromethamine, Caprylyl Glycol, Phenoxyethanol, Fragrance.

NDC: 49620-101-51



OBLITERATE®

Booster Pack : **ACTIVE ACNE**

mybooster • STEP 2

2-1.0 FL OZ US / 30 mL

OBLITERATE

benzoyl peroxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49520-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOIC ACID - UNII:8SKN0B0MIM)	BENZOYL PEROXIDE	5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POLYACRYLAMIDE (1500 MW) (UNII: 5D6TC4BRWV)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
LAURETH-7 (UNII: Z95S6G8201)	
ACETYL CARBOXYMETHYL COCOYL GLYCINE (UNII: 3TNX4P92J3)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
DECYLENE GLYCOL (UNII: S57M60MI88)	
CARBOMER HOMOPOLYMER (ALLYL SUCROSE CROSSLINKED) (UNII: Z135WT9208)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49520-101-51	1 in 1 BOX		
1	NDC:49520-101-11	15 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	04/24/2012	

Labeler - mybody (004460532)

Registrant - mybody (004460532)

Revised: 5/2012

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