CLEAR MED 10%- benzoyl peroxide lotion
CONTROL CORRECTIVE SKINCARE 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.

CONTROL CORRECTION - CLEAR MED 10% (70764-107)

ACTIVE INGREDIENTS

BENZOYL PEROXIDE 10%

PURPOSE

ACNE TREATMENT

USE

HELPS REDUCE ACNE BREAKOUTS.

WARNINGS

FOR EXTERNAL USE ONLY. DO NOT USE IF YOU ARE ALLERGIC OR SENSITIVE TO BENZOYL PEROXIDE. WHEN USING THIS PRODUCT AVOID CONTACT WITH EYES, LIPS AND MOUTH. IF CONTACT OCCURS, FLUSH THOROUGHLY WITH WATER. AVOID UNNECESSARY SUN EXPOSURE. MAY BLEACH FABRIC. MILD IRRITATION MAY BE REDUCED BY USING THE PRODUCT LESS FREQUENTLY OR IN A LOWER CONCENTRATION. DISCONTINUE USE, IF IRRITATION OCCURS AND CONSULT YOUR DOCTOR. USING OTHER TOPICAL ACNE MEDICATION AT THE SAME TIME OR IMMEDIATELY FOLLOWING USE OF THIS PRODUCT MAY INCREASE DRYNESS OR IRRITATION OF THE SKIN.

DIRECTIONS

ALWAYS PATCH TEST A SMALL AREA BEFORE FIRST APPLICATION. APPLY SPARINGLY TO AFFECTED AREAS ONCE OR TWICE DAILY AS NEEDED, AVOIDING THE EYE AREA. DRYNESS AND LIGHT PEELING MAY OCCUR DURING THE FIRST FEW WEEKS OF USAGE.

INACTIVE INGREDIENTS

WATER, SULFUR 3%, BUTYLENE GLYCOL, CARBOMER, POTASSIUM SORBATE, PHENOXYETHANOL, SODIUM BENZOATE, SODIUM HYDROXIDE.

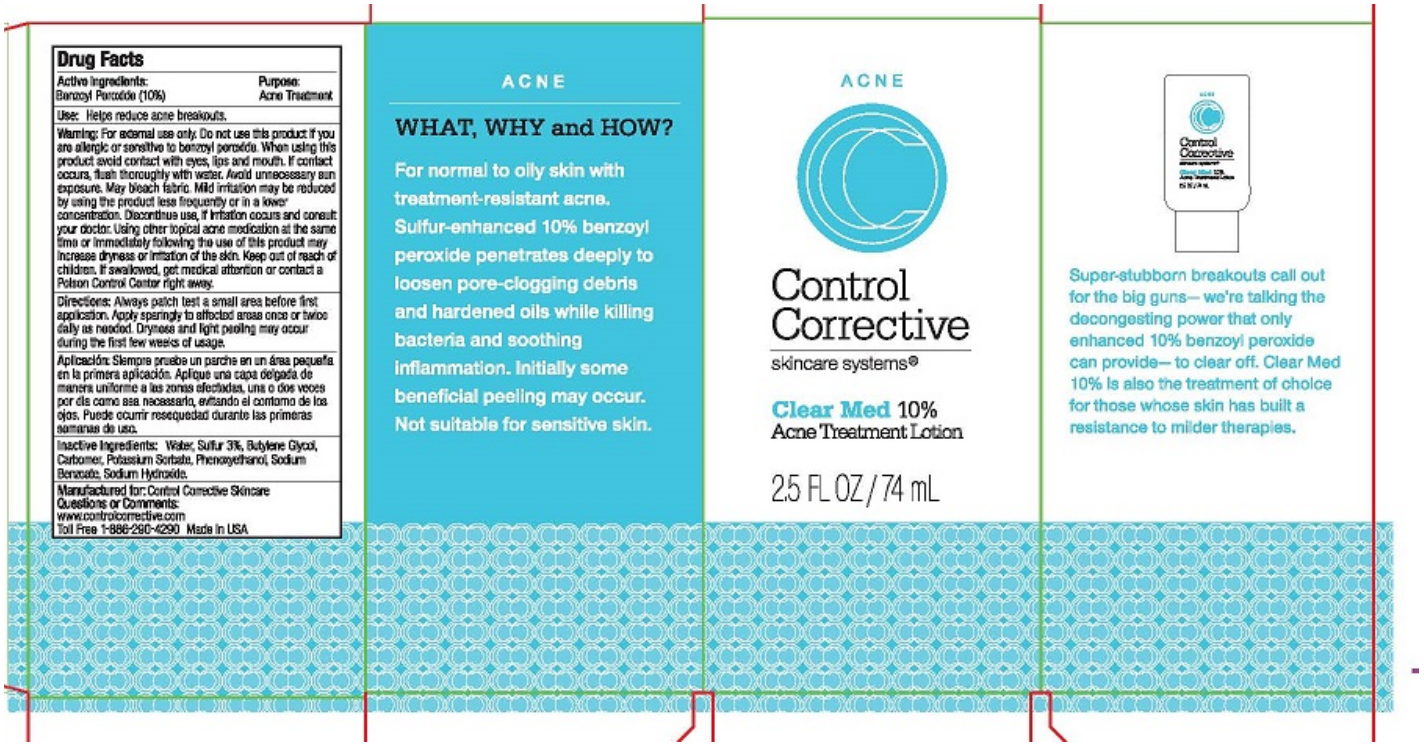
KEEP OUT OF REACH OF CHILDREN. IF SWALLOWED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

QUESTIONS OR COMMENTS:

WWW.CONTROLCORRECTIVE.COM

TOLL FREE 1-866-290-4290

MADE IN USA



CLEAR MED 10%

benzoyl peroxide lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70764-107
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	10 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SULFUR (UNII: 70FD1KFU70)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70764-107-51	1 in 1 BOX	06/09/2016	
1	NDC:70764-107-11	74 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 final	M006	06/09/2016	

Labeler - CONTROL CORRECTIVE SKINCARE INC (023999357)

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