
END-ZIT® ACNE CONTROL DRYING LOTION

Intended for use on blemishes only. This advanced formulation is designed to dry and aid in elimination of pimples. It is make-up quality for excellent coverage of the blemish. It is unbelievably effective, clinically tested, dermatologist approved and recommended.

DIRECTIONS: Shake bottle well before each use. Apply directly to blemish only, using applicator. Wait a few moments until slightly dry. Pat with fingertip or clean cotton swab to blend color to skin. Make-up may be applied if desired. Should remain on skin all day. Re-apply at night to clean skin to remain on skin overnight.

NOTE: PERSONS WITH KNOWN SENSITIVITY TO SULFUR SHOULD AVOID USE OF THIS PRODUCT. KEEP FROM EYES.

DRUG FACTS ACTIVE INGREDIENT SULFUR 5%

PURPOSE ACNE TREATMENT

WARNINGS:

• For external use only.

• Using other topical acne medications at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.

• Do not get into eyes. If excessive skin irritation deve lops or increases, discontinue use and consult a doctor.

• Keep out of reach of children.

OTHER INGREDIENTS: Isopropyl Alcohol, Water, Zinc Oxide, Propylene Glycol, Camphor, Talc, Sodium Laureth Sulfate, Titanium Dioxide, Diazolidinyl Urea, Methylparaben, Propylparaben. May contain Iron Oxides.

NDC #'s

Light/Medium 68605-2001-2 Medium/Dark 68605-2002-2 Acne Control Mask 68605-2010-2

----PACKAGE LABEL.PRINCIPAL DISPLY PANEL----

DIRECTIONS: SHAKE BOTTLE WELL.

APPLY TWICE DAILY DIRECTLY TO ACNE BREAKOUT. ALLOW TO DRY FOR 10 SECONDS. PAT TO BLEND. APPLY MAKE-UP IF DESIRED.

ACTIVE INGREDIENT: SULFUR 5%.

WARNINGS: AVOID IF ALLERGIC TO SULFUR AVOID CONTACT WITH EYES FOR EXTERNAL USE ONLY KEEP FROM CHILDREN

Manufactured by ABBE Laboratories, Inc. Farmingdale, NY 11735 Made in the U.S.A.

END-ZIT®

ACNE CONTROL DRYING LOTION

ABBE

0.62 OZ. (17.57 g)



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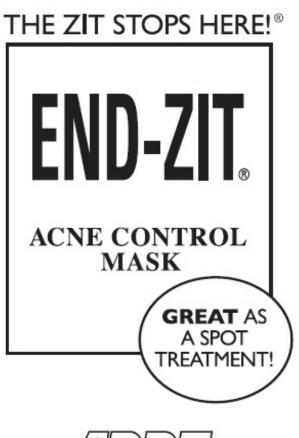
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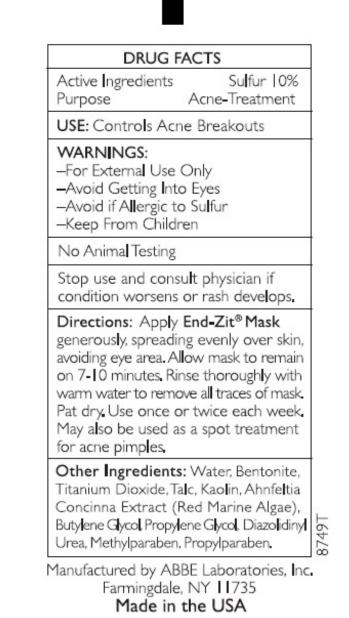
ABBE

0.62 OZ. (17.57 g)





NET WT. 2.5 OZ. (70g)



END-ZIT sulfur lotion, augmented					
Product Information					
Product Type	HUMAN OTC DRUG	ltem Co	ode (Source)	NDC	:68605-2001
Route of Administration	TOPICAL				
Active Ingredient/Active	Moiety				
Ingredie	nt Name		Basis of Streng	th	Strength
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)			SULFUR		.05 g in 1 mL

	Ingredient Name		Strength			
SOPROPYL ALCO						
WATER (UNII: 059	QF0KO0R)					
ZINC OXIDE (UNI	: SOI2LOH54Z)					
PROPYLENE GLY	COL (UNII: 6DC9Q167V3)					
CAMPHOR (SYNT	THETIC) (UNII: 5TJD82A1ET)					
TALC (UNII: 7SEV)	/J4R1U)					
SODIUM LAURET	H SULFATE (UNII: BPV390UAP0)					
TITANIUM DIOXI	DE (UNII: 15FIX9V2JP)					
	REA (UNII: H5RIZ3MPW4)					
METHYLPARABE	N (UNII: A2I8C7HI9T)					
PROPYLPARABE	(UNII: Z8IX2SC1OH)					
Product Cha	racteristics					
Color	brown (Light/Medium)	Score				
Shape		Size	Size			
Flavor		Imprint Code	Imprint Code			
Contains						
Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing En Date			
1 NDC:68605- 2001-2	14.78 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	04/20/2004				
	Information					
Marketing						
Marketing Marketing Category	Application Number or Monogra Citation	ph Marketing Start Date	Marketing End Date			

END-ZIT sulfur lotion, augmented					
Product Information					
Product Type	HUMAN OTC DRUG	ltem Coc	le (Source)	NDC:6	8605-2002
Route of Administration	TOPICAL				
Active Ingredient/Active	Moiety				
•	ent Name		Basis of Stren	gth	Strength
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)			SULFUR		.05 g in 1 g

	Strength				
ISC					
WA	TER (UNII: 059	QF0KO)R)		
ZIN	IC OXIDE (UNII	: SOI2L	OH54Z)		
PR	OPYLENE GLY	COL (U	NII: 6DC9Q167V3)		
CA	MPHOR (SYNT	HETIC	(UNII: 5TJD82A1ET)		
TA	LC (UNII: 7SEV7	/J4R1U)			
so	DIUM LAURET	H SULF	ATE (UNII: BPV390UAP0)		
тіт		DE (UNI	: 15FIX9V2JP)		
DIA	ZOLIDINYL UP	REA (UN	II: H5RIZ3MPW4)		
ME	THYLPARABE	N (UNII:	A2I8C7HI9T)		
PR	OPYLPARABEN	(UNII:	Z8IX2SC1OH)		
Pr	oduct Chai	racte	ristics		
	oduct Chai Ior	racte	brown (Medium/Dark)	Score	
Co		racte		Score Size	
Co Sh	lor	racte			
Co Sh Fla	lor ape	racte		Size	
Co Sh Fla	lor ape ivor	racte		Size	
Co Sh Fla	lor ape ivor	racte		Size	
Co Sh Fla Co	lor ape ivor	racte		Size	
Co Sh Fla Co P a	lor ape ivor ntains	racte		Size	Marketing En Date
Co Sh Fla Co P a #	lor ape ivor ntains ackaging	17.57	brown (Medium/Dark)	Size Imprint Code Marketing Start	-
Co Sh Fla Co Pa #	lor ape ovor ntains ackaging Item Code NDC:68605-	17.57	brown (Medium/Dark) Package Description g in 1 BOTTLE, GLASS; Type 0: Not a	Size Imprint Code Marketing Start Date	-
Co Sh Fla Co Pa #	lor ape ovor ntains ackaging Item Code NDC:68605-	17.57 Combi	brown (Medium/Dark) Package Description g in 1 BOTTLE, GLASS; Type 0: Not a hation Product	Size Imprint Code Marketing Start Date	-
Co Sh Fla Co Pa #	lor ape nvor ntains ackaging item Code NDC:68605- 2002-2	17.57 Combi	brown (Medium/Dark) Package Description g in 1 BOTTLE, GLASS; Type 0: Not a hation Product	Size Imprint Code Marketing Start Date	-

END-ZII					
sulfur lotion, augmented					
Product Information					
Product Type	HUMAN OTC DRUG	ltem Cod	le (Source)	NDC:68	3605-2010
Route of Administration	TOPICAL				
Active Ingredient/Active	Moiety				
Ingredie	ent Name		Basis of Stren	gth	Strength
SULFUR (UNII: 70FD1KFU70) (SULF	UR - UNII:70FD1KFU70)		SULFUR		.05 g in 1 g

In	active Ingred	ients				
		lente	Ingredient Name			Strength
ISC	OPROPYL ALCOHO	Jucigu				
	ATER (UNII: 059QFC	-				
	NC OXIDE (UNII: SO		Ζ)			
PR	OPYLENE GLYCO	L (UNII: 6[DC9Q167V3)			
CA	MPHOR (SYNTHE	TIC) (UNI	: 5TJD82A1ET)			
ГΑ	LC (UNII: 7SEV7J4R	1U)				
50	DIUM LAURETH S	ULFATE	(UNII: BPV390UAP0)			
ТІТ		(UNII: 15F	X9V2JP)			
DI/	ZOLIDINYL UREA	UNII: H5	RIZ 3MPW4)			
ME	THYLPARABEN (U	JNII: A2180	7HI9T)			
PR	OPYLPARABEN (U	NII: Z8IX2	SC10H)			
Pr	oduct Charac	teristi	CS			
Color white (Mask) Score						
Shape			Size			
Fla	vor			Imp	rint Code	
Co	ntains					
Pa	ackaging					
#	Item Code		Package Description		Marketing Start Date	Marketing End Date
		70 g in 1 T Product	UBE; Type 0: Not a Combination	04	4/20/2004	
	arketing Ir	nform	ation			
	arketing Ir Marketing Category		ation ication Number or Monogra Citation	ph	Marketing Start Date	Marketing Enc Date

Labeler - ABBE Laboratories, Inc. (781745286)

Establishment					
Name	Address	ID/FEI	Business Operations		
ABBE Laboratories, Inc.		781745286	manufacture(68605-2001, 68605-2002, 68605-2010)		

Revised: 11/2023

ABBE Laboratories, Inc.