QUICK DRY ACNE TREATMENT- sulfur lotion
DR. THROWER'S 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.

ACTIVE INGREDIENTS:

SULFUR- 10.0%

ZINC OXIDE- 5.0%

PURPOSE:

TREATS ACNE AND SHAVE BUMPS

USES:

- DRIES UP EXCESS OIL
- CLEARS UP ACNE AND SHAVING BUMPS

DIRECTIONS:

USE EVENINGS. SHAKE TO ACTIVATE

APPLY TO CLEAN SKIN AFTER SPF 30 IN AM OR ALOE MOISTURIZING CREAM IN PM

APPLY DIRECTLY TO THE ACNE OR SHAVING BUMPS WITH A COTTON SWAB.

ALLOW TO DRY FOR APPROXIMATELY 5-10 MINTUES AND LEAVE ON OVERNIGHT.

INACTIVE INGREDIENTS:

ETHYL ALCOHOL, AQUA, SPIRAEA ULMARIA EXTRACT, CINNAMOMUM CAMPHORA OIL, MAGNESIUM ALUMINIUM SILICATE, PIGMENT YELLOW 42, PIGMENT BLACK11, PIGMENT RED 101, PEG-8 DIMETHICONE, GLYCERINE, OCTYLDODECANOL, RETINYL PALMITATE (VITAMIN A), TOCOPHERYL ACETATE (VITAMIN E), ASCORBIC ACID (VITAMIN C), PYRIDOXINE HCL (VITAMIN B6), SILICA, SODIUM PROPOXYHYDROXYPROPYL THIOSULFATE SILICA, CHOLECALCIFEROL

KEEP OUT OF REACH OF CHILDREN.

WARNINGS

FOR EXTERNAL USE ONLY

QUICK DRY ACNE TREATMENT

sulfur lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69299-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	10 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
FILIPENDULA ULMARIA WHOLE (UNII: 3LH0M209LN)	
CAMPHOR OIL (UNII: 75IZZ8Y727)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
GLYCERIN (UNII: PDC6A3C0OX)	
OCTYLDODECANOL (UNII: 461N1O614Y)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)	

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM PROPOXYHYDROXYPROPYL THIOSULFATE SILICA (UNII: 208G222332)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69299-101-51	1 in 1 BOX		
1	NDC:69299-101-11	30 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	11/17/2014	

Labeler - DR. THROWER'S SKINCARE, INC. (078711495)

Registrant - DR. THROWER'S SKINCARE, INC. (078711495)

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