DR. THROWERS HYDROTET- tretinoin, hydrocortisone cream
DR. THROWER'S SKINCARE, INC.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

DIRECTIONS:

APPLY MEDICATION TO OILY AREA ONLY OF THE FACE IN VERY THIN LAYER. LEAVE ON OVERNIGHT.

PURPOSE:

RX STRENGTH TOPICAL RETINOID FOR TREATMENT OF ACNE, SHAVING BUMPS, AND SPECIFIC BUMPY SKIN PROBLEMS. LEAVES SKIN SMOOTH, CLEAR AND NEW.

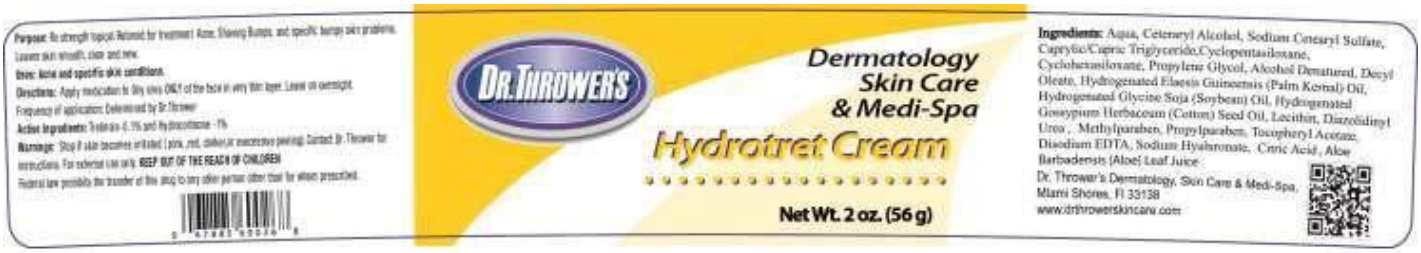
uSES: ACNE AND SPECIFIC SKIN CONDITIONS.

INGREDIENTS:

Aqua, Ceteraryl Alcohol, Sodium Cetearyl Sulfate, Caprylic/Capric Triglyceride,Cyclopentasiloxane, Cyclohexasiloxane, Propylene Glycol, Alcohol Denatured, Decyl Oleate, Hydrogenated Elaesis Guineensis (Palm Kernal) Oil, Hydrogenated Glycine Soja (Soybean) Oil, Hydrogenated Gossypium Herbaceum (Cotton) Seed Oil, Lecithin, Diazolidinyl Urea , Methylparaben, Propylparaben, Tocopheryl Acetate, Disodium EDTA, Sodium Hyaluronate, Citric Acid , Aloe Barbadensis (Aloe) Leaf Juice

WARNINGS

STOP USE IF SKIN BECOMES IRRITATED (PINK, RED DARKER, OR EXCESSIVE PEELING). cONTACT DR. THROWER FOR INSTRUCTIONS. FOR EXTERNAL USE ONLY. KEEP OUT OF REACH OF CHILDREN. FEDERAL LAW PROHIBITS THE TRANSFER OF THIS DRUG TO ANY OTHER PERSON OTHER THAN FOR WHOM IT IS PRESCRIBED.



DR. THROWERS HYDROTET

tretinoin, hydrocortisone cream

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:69299-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRETINOIN (UNII: 5688UTC01R) (TRETINOIN - UNII:5688UTC01R)	TRETINOIN	0.05 g in 100 g
HYDROCORTISONE (UNII: WI4X0X7BPJ) (HYDROCORTISONE - UNII:WI4X0X7BPJ)	HYDROCORTISONE	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
DECYL OLEATE (UNII: ZGR06DO97T)	
HYDROGENATED PALM KERNEL OIL (UNII: FM8D1RE2VP)	
HYDROGENATED SOYBEAN OIL (UNII: A2M91M918C)	
HYDROGENATED COTTONSEED OIL (UNII: Z82Y2C65EA)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69299-102-22	56 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/15/2015	

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

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

Revised: 5/2015 DR. THROWER'S SKINCARE, INC.