MG217 PSORIASIS THERAPEUTIC CONDITIONER- salicylic acid shampoo Wisconsin Pharmacal Company

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 Ingredient

Salicylic Acid 3%

Purpose

Psoriasis, Seborrheic Dermatitis, Dandruff Treatment

Uses

For the temporary relief of these symptoms associate with psoriasis, seborrheic dermatitis or dandruff

- itching
- scaling
- flaking,
- reness
- irritaion of the scalp

Helps prevent recurrence.

Warnings

For external use only.

Ask a doctor before use

• if condition covers larger area of body, consult your doctor before using this product.

When using this product

• avoid contact with the eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if

• condition worsens or does not improve after regular use of this product as directed, consult a doctor.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center Immediately.

Directions

- wet hair then massage liberal amount of MG217 into the scalp and leave on for several minutes. Rinse thoroughly.
- for best results, use at least twice a week or as directed by a physician.

Inactive Ingredients

Purified water, Sodium laureth sulfate, Cocamidopropyl betaine, Lauryl glucoside, Sodium citrate, Coco-glucoside, Glyceryl oleate, Propanediol, Glycerin, Polyquaterinum-44, Sodium benzoate, Panthenol, Calendula officinalis flower extract, Chamomilla recutita (matricaria) flower extract, Propylene glycol, Citric acid, Chlorhexidine gluconate, Disodium EDTA





MG217 PSORIASIS THERAPEUTIC CONDITIONER salicylic acid shampoo

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:68093-7242 Route of Administration TOPICAL

Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	

Inactive Ingredients	
Ingredient Name	Strength
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
CHLORHEXIDINE GLUCONATE (UNII: MOR8 4MUD8 E)	
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390 UAPO)	
COCAMIDO PRO PYL BETAINE (UNII: 50 CF30 11KX)	
SO DIUM CITRATE (UNII: 1Q73Q2JULR)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
PANTHENOL (UNII: WV9CM0O67Z)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
MATRICARIA CHAMOMILLA FLOWERING TOPOIL (UNII: SA8AR2W4ER)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)	
LAURYL GLUCO SIDE (UNII: 76 LN7P7UCU)	
COCO GLUCOSIDE (UNII: ICS790225B)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
PROPANEDIOL (UNII: 5965N8W85T)	
POLYQUATERNIUM-44 (200000 MW) (UNII: TTD9 VOM830)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:68093-7242-1	1 in 1 CARTON	11/0 1/20 15	
1	283 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333D	11/0 1/20 15		

Labeler - Wisconsin Pharmacal Company (800873986)

Establishment					
Name	Address	ID/FEI	Business Operations		
Wisconsin Pharmacal Company		800873986	manufacture(68093-7242)		

Wisconsin Pharmacal Company Revised: 3/2016