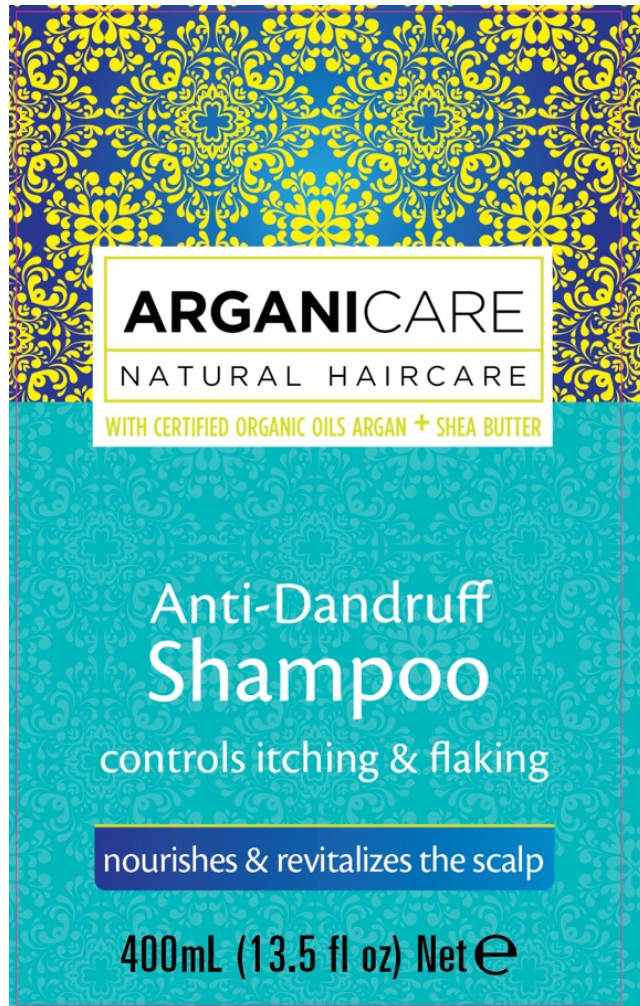


**ANTIDANDRUFFSHAMPOO- pyrithione zinc shampoo shampoo**  
**Peer Pharm Ltd.**

*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.*

**Arganicare Anti-Dandruff Shampoo**



Arganicare anti-dandruff shampoo is designed to control dandruff symptoms and to keep the skin of the scalp moisturized. The shampoo is enriched with Argan Oil and Shea Butter and an advanced complex of vitamins and microelements. This deeply nourishing formula leaves hair manageable and refreshed with a pleasantly clean sensation.

Drug Facts	
<b>Active ingredient</b>	<b>Purpose</b>
Zinc Pyrithione 1.0%	Anti-dandruff
<b>Use:</b> Controls itching and flaking associated with dandruff.	
<b>WARNINGS</b>	
For external use only.	
<b>Avoid contact with eyes</b> If contact occurs, rinse eyes thoroughly with water.	
<b>Stop use and ask a doctor if</b> condition worsens or does not improve after regular use as directed.	
<b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center right away.	
<b>Directions</b> Wet hair, apply shampoo and work into a lather. Rinse thoroughly with warm water. For best results, use at least twice a week or as directed by a doctor.	
<b>Inactive Ingredients</b> AQUA (WATER), COCAMIDOPROPYL HYDROXYSULTAINE, AMMONIUM LAURETH SULFATE, HAMAMELIS VIRGINIANA (WITCH HAZEL) WATER, GLYCERIN, COCAMIDE MEA, COCAMIDOPROPYL BETAINE, ARGANIA SPINOSA (ARGAN) KERNEL OIL, SILICONE QUATERNIUM-22, GLYCOL DISTEARATE, CARBOMER, SIMMONDSIA CHINENSIS (JUJOBA) SEED OIL, POLYQUATERNIUM-39, C12-15 ALKYL LACTATE, PEG-12 DIMETHICONE, POTASSIUM SORBATE, SODIUM BENZOATE, FRAGRANCE, DIMETHYL HYDANTOIN, GUAR, HYDROXYPROPYLTRIMONIUM CHLORIDE, ACRYLAMIDOPROPYLTRIMONIUM CHLORIDE/ACRYLAMIDE COPOLYMER, BUTYROSPERMUM PARKII (SHEA) OIL, PANTHENOL, LACTIC ACID, TETRASODIUM EDTA.	

www.arganicare.com  
Manufactured  
by Peer Pharm LTD.  
Made in Israel

905404 ITEM # 4657

7290104367397

Zinc Pyrithione 1.0%

Controls Dandruff

Keep out of reach of children.

Controls itching and flaking associated with dandruff

For external use only. Avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water. If condition worsens or does not improve after regular use of this product as directed, consult a doctor.

Wet hair, apply shampoo and work into a lather. Rinse thoroughly with warm water. For best results, use at least twice a week or as direct by a doctor.

**ANTIDANDRUFFSHAMPOO**

pyrithione zinc shampoo shampoo

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69435-1608
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	1 g in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
PANTHENOL (UNII: WV9CM0O67Z)	
WATER (UNII: 059QF0KO0R)	
AMMONIUM LAURETH-12 SULFATE (UNII: E1ZP93931S)	
GLYCERIN (UNII: PDC6A3C0OX)	
JOJOBA OIL (UNII: 724GKU717M)	
HAMAMELIS VIRGINIANA TOP WATER (UNII: NT00Y05A2V)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
POLYQUATERNIUM-39 (22.5/51/26.5 ACRYLIC ACID/ACRYLAMIDE/DADMAC; 160000 MW) (UNII: X2NH1K9F8K)	
C12-15 ALKYL LACTATE (UNII: GC844VRD7E)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
ACRYLAMIDOPROPYLTRIMONIUM CHLORIDE/ACRYLAMIDE COPOLYMER (400 MPA.S) (UNII: 2RX4MI2LCX)	
SHEANUT OIL (UNII: O88E196QRF)	
LACTIC ACID (UNII: 33X04XA5AT)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	
ARGAN OIL (UNII: 4V59G5UW9X)	
CARBOMER 1342 (UNII: 809Y72KV36)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69435-1608-1	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/23/2016	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	03/23/2016	

**Registrant** - Peer Pharm Ltd. (514678390)

**Establishment**

Name	Address	ID/FEI	Business Operations
Peer Pharm Ltd.		514678390	manufacture(69435-1608)

Revised: 6/2016

Peer Pharm Ltd.