

**DANDRUFF- pyrithione zinc shampoo**  
**Target Corporation**

*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.*

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**Top Care 281.007 281AK**

**claims**

up & up dry scalp relief dandruff shampoo with almond oil protects the scalp's natural moisture and leaves hair soft, shiny and manageable. It is gentle and pH balanced for everyday use.

**Active ingredient**

Pyrithione zinc 1%

**Purpose**

Anti-dandruff

**use**

helps prevent recurrence of flaking and itching associated with dandruff

**warnings**

For external use only

**When using this product**

- do not get into eyes. If contact occurs, rinse eyes thoroughly with water.

**Stop use and ask a doctor if**

- condition worsens or does not improve after regular use as directed

**Keep out of reach of children**

If swallowed, get medical help or contact a Poison Control Center right away

**Directions**

- shake well
- for maximum dandruff control, use every time you shampoo
- wet hair, massage onto scalp, rinse, repeat if desired
- for best results use at least twice a week or as directed by a doctor

**Inactive ingredients**

water, sodium lauryl sulfate, sodium laureth sulfate, glycol distearate, sodium chloride, zinc carbonate, sodium xylenesulfonate, amodimethicone, cocamidopropyl betaine, fragrance, sodium benzoate, guar hydroxypropyltrimonium chloride, magnesium carbonate hydroxide, citric acid, Prunus amygdalus dulcis (sweet almond) oil, methylchloroisothiazolinone, methylisothiazolinone

**Questions?**

Call 1-800-910-6874

This product is not manufactured or distributed by Procter & Gamble, distributor of Head & Shoulders Dry Scalp Care Shampoo

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Ingredient Name	Basis of Strength	Strength
<b>PYRITHIONE ZINC</b> (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ZINC CARBONATE (UNII: EQR32Y7H0M)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
AMODIMETHICONE (800 CST) (UNII: 363Z2T48P7)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
MAGNESIUM CARBONATE HYDROXIDE (UNII: YQO029V1L4)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALMOND OIL (UNII: 66YXD4DKO9)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-281-35	700.9 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/18/2002	
2	NDC:11673-281-15	420 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/18/2002	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	07/18/2002	

**Labeler** - Target Corporation (006961700)

**Registrant** - Vi-Jon, LLC (790752542)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(11673-281)

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