

**HEAD AND SHOULDERS CLASSIC CLEAN DANDRUFF- pyrithione zinc shampoo**  
**Navajo Manufacturing Company Inc**

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**Head & Shoulders Classic Clean Dandruff Shampoo**

**Active ingredients**

Pyrithione zinc 1%

**Purpose**

Anti-dandruff

**Uses**

helps prevent recurrence of flaking and itching associated with dandruff

**Warnings**

**For external use only**

**When using this product**

- avoid contact with the 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 of this product as directed

**Keep this and all drugs out of reach of children.**

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

**Directions**

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

**Inactive ingredients**

water, sodium lauryl sulfate, sodium laureth sulfate, glycol distearate, zinc carbonate, sodium chloride, sodium xylenesulfonate, cocamidopropyl betaine, fragrance, dimethicone, sodium benzoate, guar hydroxypropyltrimonium chloride, magnesium carbonate hydroxide, methylchloroisothiazolinone, methylisothiazolinone, blue 1, red 33

**Questions or comments?**

1-800-723-9569

Head & Shoulders Classic Clean Dandruff Shampoo 50ml (67751-053-01)



**HEAD AND SHOULDERS CLASSIC CLEAN DANDRUFF**  
pyrithione zinc shampoo

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67751-053(NDC:37000-071)
Route of Administration	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		Strength
Ingredient Name		
WATER (UNII: 059QF0KO0R)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)		
GLYCOL DISTEARATE (UNII: 13W7MDN21W)		

<b>ZINC CARBONATE</b> (UNII: EQR32Y7H0M)
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)
<b>SODIUM XYLENESULFONATE</b> (UNII: G4LZF950UR)
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)
<b>GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE)</b> (UNII: B16G315W7A)
<b>MAGNESIUM CARBONATE HYDROXIDE</b> (UNII: YQO029V1L4)
<b>METHYLCHLOROISOTHIAZOLINONE</b> (UNII: DEL7T5QRPN)
<b>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67751-053-01	1 in 1 CARTON	11/10/2017	
1		50 mL 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 Drug	M032	05/31/2013	

**Labeler** - Navajo Manufacturing Company Inc (091917799)

### Establishment

Name	Address	ID/FEI	Business Operations
Navajo Manufacturing Company Inc		136941411	relabel(67751-053) , repack(67751-053)

Revised: 4/2024

Navajo Manufacturing Company Inc