

**QUALITY CHOICE DANDRUFF 2 IN 1- pyrithione zinc lotion/shampoo  
CHAIN DRUG MARKETING ASSOCIATION, INC.**

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**Dandruff 2 in 1 shampoo QC**

**Drug Facts**

***Active Ingredient***

Pyrithione zinc

***Purpose***

Antidandruff

***Use***

helps prevent recurrence of scalp 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 out of reach of children***

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

***Directions***

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

***Other information***

store away from direct sunlight, excessive heat, and freezing conditions

***Inactive ingredients***

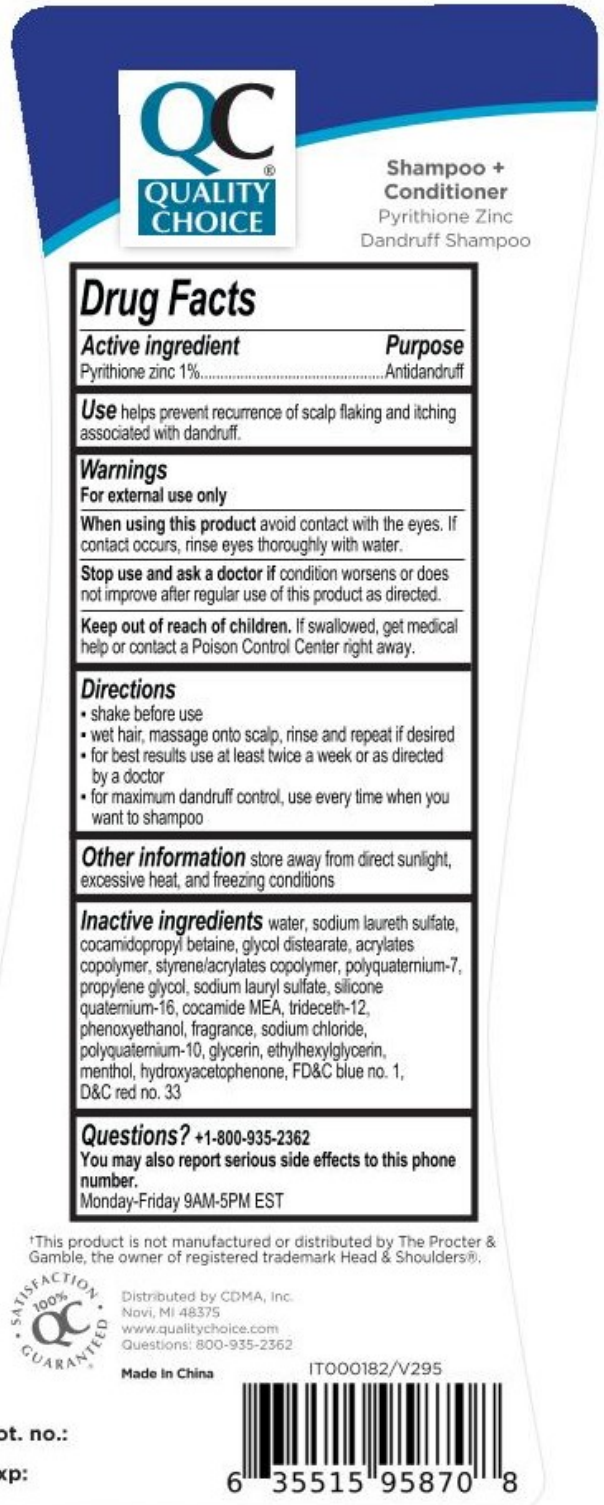
water, sodium laureth sulfate, cocamidopropyl betaine, glycol distearate, acrylates copolymer, styrene/acrylates copolymer, polyquaternium-7, propylene glycol, sodium lauryl sulfate, silicone quaternium-16, cocamide MEA, trideceth-12, phenoxyethanol, fragrance, sodium chloride, polyquaternium-10, glycerin, ethylhexylglycerin, menthol, hydroxyacetophenone, FD&C blue no. 1, D&C red no. 33

***Questions?***

**+1-800-935-2362**

**You may also report serious side effects to this phone number.**

Monday-Friday 9AM-5PM EST



**Drug Facts**

Active ingredient	Purpose
Pyrrithione zinc 1%.....	Antidandruff

**Use** helps prevent recurrence of scalp 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.

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**Directions**

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

**Other information** store away from direct sunlight, excessive heat, and freezing conditions

**Inactive ingredients** water, sodium laureth sulfate, cocamidopropyl betaine, glycol distearate, acrylates copolymer, styrene/acrylates copolymer, polyquaternium-7, propylene glycol, sodium lauryl sulfate, silicone quaternium-16, cocamide MEA, trideceth-12, phenoxyethanol, fragrance, sodium chloride, polyquaternium-10, glycerin, ethylhexylglycerin, menthol, hydroxyacetophenone, FD&C blue no. 1, D&C red no. 33

**Questions?** +1-800-935-2362  
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\*This product is not manufactured or distributed by The Procter & Gamble, the owner of registered trademark Head & Shoulders®.



Distributed by CDMA, Inc.  
Novi, MI 48375  
www.qualitychoice.com  
Questions: 800-935-2362

Made in China

IT000182/V295

Lot. no.:

Exp:



**QUALITY CHOICE DANDRUFF 2 IN 1**

pyrrithione zinc lotion/shampoo

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83324-321
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

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
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	
<b>POLYQUATERNIUM-10 (1000 MPA.S AT 2%)</b> (UNII: GMR4PEN8PK)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW)</b> (UNII: JZ1374NL9E)	
<b>POLYQUATERNIUM-7</b> (UNII: 0L414VCS5Y)	
<b>QUATERNIUM-16</b> (UNII: K4H0FG9I2T)	
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)	
<b>TRIDECETH-12</b> (UNII: YFY3KG5Y7O)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>HYDROXYACETOPHENONE</b> (UNII: G1L3HT4CMH)	
<b>GLYCOL DISTEARATE</b> (UNII: 13W7MDN21W)	
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)	
<b>COCAMIDE MEA</b> (UNII: C80684146D)	
<b>SODIUM LAURETH SULFATE</b> (UNII: BPV390UAP0)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	

**Product Characteristics**

<b>Color</b>	blue	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-321-13	399 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/23/2025	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	01/23/2025	

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**Labeler** - CHAIN DRUG MARKETING ASSOCIATION, INC. (011920774)

**Registrant** - Medical Group Care LLC (085137089)

Revised: 1/2025

CHAIN DRUG MARKETING ASSOCIATION, INC.