QUALITY CHOICE DANDRUFF EVERYDAY CLEAN- pyrithione zinc lotion/shampoo
CHAIN DRUG MARKETING ASSOCIATION, INC.

Dandruff Everyday Clean 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

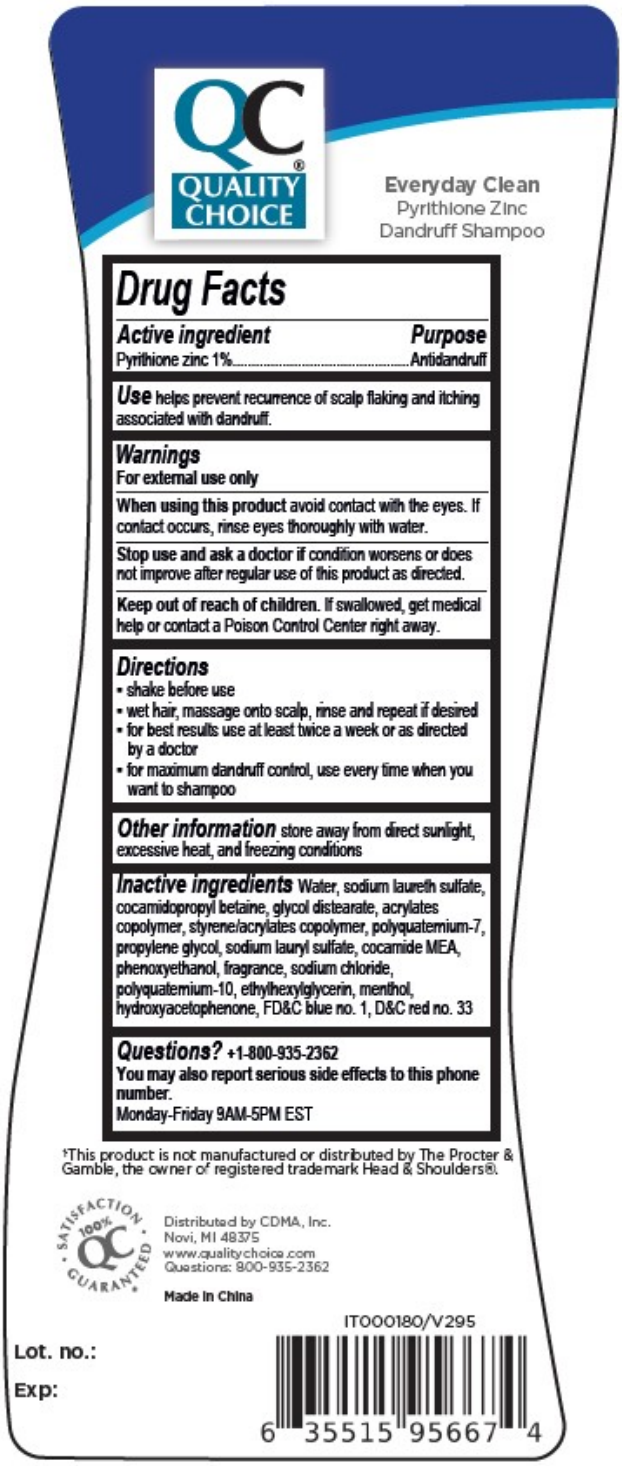
store away from direct sunlight, excessive heat, and freezing conditions

Inactive ingredients

Water, sodium laureth sulfate, cocamidopropyl betaine, glycol distearate, acrylates copolymer, styrene/acrylates copolymer, polyquaternium-7, propylene glycol, sodium lauryl sulfate, cocamide MEA, phenoxyethanol, fragrance, sodium chloride, polyquaternium-10, 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



QUALITY CHOICE DANDRUFF EVERYDAY CLEAN

pyrithione zinc lotion/shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-320
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

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

Product Characteristics

Color	blue	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-320-13	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	

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

Registrant - Medical Group Care LLC (085137089)

Revised: 1/2025 CHAIN DRUG MARKETING ASSOCIATION, INC.