HEAD AND SHOULDERS CLINICAL ITCH RELIEF SCALP MIST- pyrithione zinc liquid
The Procter & Gamble Manufacturing Company

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.

Head and Shoulders ® Clinical Itch Relief Scalp Mist

Drug Facts

Active ingredient

Pyrithione zinc 0.1%

Purpose

Anti-dandruff

Use

for the relief of dandruff.

Warnings

For external use only.

When using this product
- avoid contact with 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
- apply to affected areas one to four times daily or as directed by a doctor.

Inactive ingredients

Water, niacinamide, PEG-40 hydrogenated castor oil, phenoxyethanol, caffeine, bis-PEG/PPG-16/16 PEG/PPG-16/16 dimethicone, glycerin, benzyl alcohol, fragrance, menthol, acrylates/C10-30 alkyl acrylate crosspolymer, panthenol, tetrahydroxypropyl ethylenediamine, ethylhexylglycerin, mentha piperita (peppermint) oil, mentha arvensis leaf oil.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 125 mL bottle label

head &
shoulders ®
pyrithione zinc **dandruff treatment**

CLINICAL

ITCH RELIEF

SCALP MIST

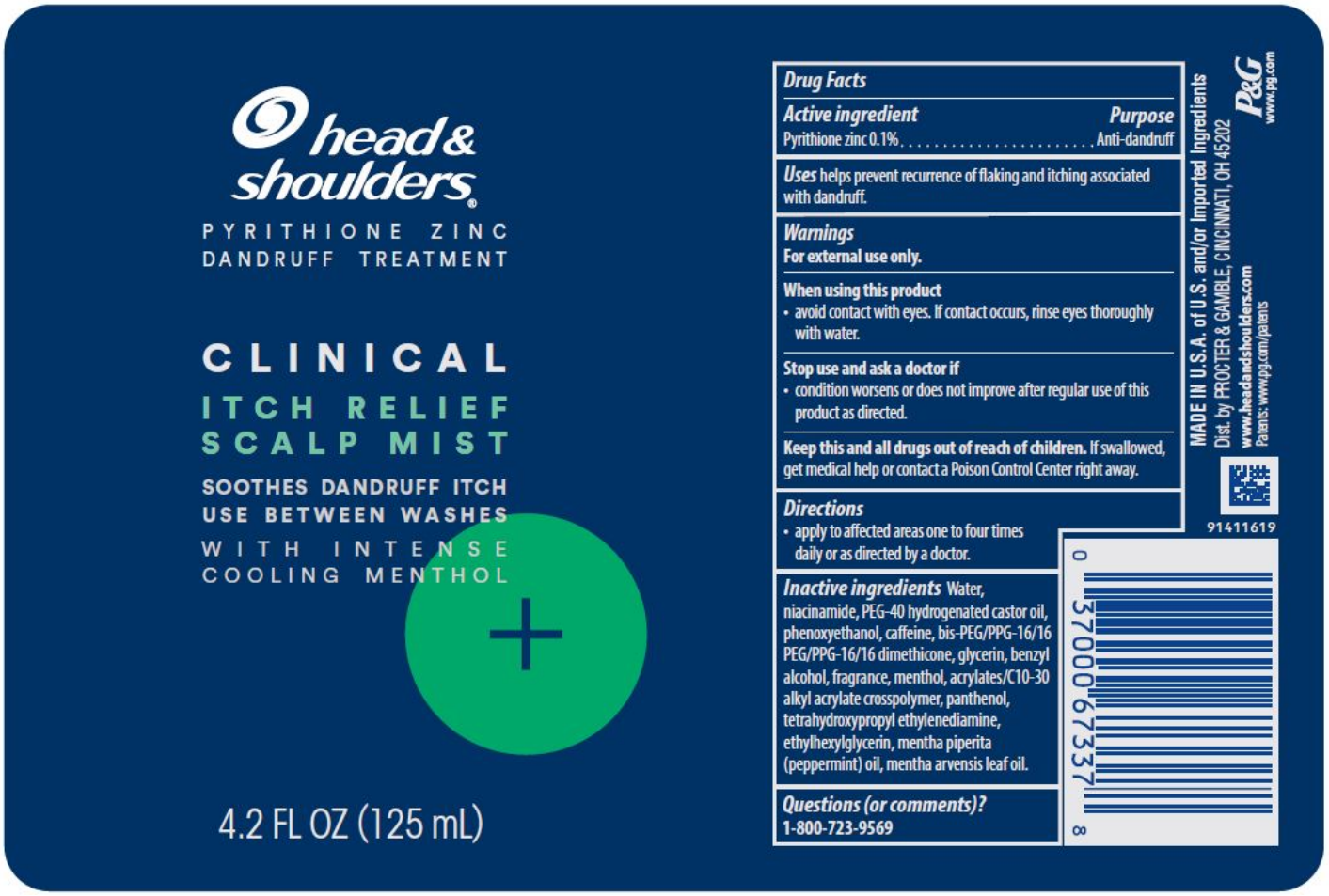
SOOTHES DANDRUFF ITCH

USE BETWEEN WASHES

WITH INTENSE

COOLING MENTHOL

4.2 FL OZ (125 mL)



HEAD AND SHOULDERS CLINICAL ITCH RELIEF SCALP MIST

pyrithione zinc liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-529
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
WATER (UNII: 059QF0KO0R)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PANTHENOL (UNII: WV9CM0O67Z)	
CAFFEINE (UNII: 3G6A5W338E)	
BIS-PEG/PPG-16/16 PEG/PPG-16/16 DIMETHICONE (UNII: 55A74AJ3KB)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETOL (UNII: Q4R969U9FR)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-529-12	125 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M032	11/17/2020	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 1/2023

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