HEAD AND SHOULDERS SUPREME DETOXIFYING PRE-WASH MASK- pyrithione zinc cream
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 ® Supreme Detoxifying Scalp Pre-Wash Mask

Drug Facts

Active ingredient

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 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, fragrance, argania spinosa kernel oil,

tocopheryl acetate, cocamidopropyl betaine, dimethicone, menthol, sodium benzoate, guar hydroxypropyltrimonium chloride, magnesium carbonate hydroxide, mentha piperita (peppermint) oil, mentha arvensis leaf oil, eucalyptus globulus leaf extract, methylchloroisothiazolinone, methylisothiazolinone.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 50 mL pouch Label

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

SUPREME

DETOXIFYING

SCALP PRE-WASH MASK

DEEPLY CLEANSES, REMOVES

BUILD UP & DETOXIFIES THE SCALP

WITH VITRAMIN E

& ARGAN OIL

1.6 FL OZ (50 mL)

HEAD AND SHOULDERS SUPREME DETOXIFYING PRE-WASH MASK

pyrithione zinc cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69423-407
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	
Ingredient Name	**Strength**
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	

METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
MAGNESIUM CARBONATE HYDROXIDE (UNII: YQO029V1L4)	
ZINC CARBONATE (UNII: EQR32Y7H0M)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
ARGAN OIL (UNII: 4V59G5UW9X)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
WATER (UNII: 059QF0KO0R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-407-50	50 mL in 1 POUCH; Type 0: Not a Combination Product	11/21/2019	09/01/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M032	11/21/2019	09/01/2024

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 7/2023 The Procter & Gamble Manufacturing Company