

HEAD AND SHOULDERS CLINICAL DRY SCALP RESCUE KIT- pyrithione zinc
The Procter & Gamble Manufacturing Company

Head and Shoulders® Clinical Dry Scalp Rescue Kit

head & shoulders

CLINICAL STRENGTH SHAMPOO

DANDRUFF DEFENSE

DRY SCALP RESCUE

Drug Facts

Active ingredient

Selenium Sulfide 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

sodium lauryl sulfate, glycol distearate, sodiumchloride, cocamidopropyl betaine, sodiumcitrate, sodiumxylenesulfonate, fragrance, dimethicone, citric acid, sodium benzoate, tetrasodiumEDTA, hydroxypropylmethylcellulose, honey extract, red 4.

Questions (or comments)?

1-800-723-9569

head & shoulders

CLINICAL STRENGTH **CONDITIONER**

DANDRUFF DEFENSE

DRY SCALP RESCUE

Drug Facts

Active ingredient

Pyrithione zinc 0.5%

Purpose

Anti-dandruff

Uses

helps prevent recurrence of flaking and itching associated with dandruff.

Warnings

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Keep this and all drugs out of reach of children.

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Directions

- for best results use at least twice a week or as directed by a doctor.
- apply to wet hair after shampooing by gently massaging into hair and scalp, rinse well.

Inactive ingredients

Water, stearyl alcohol, cetyl alcohol, stearamidopropyl dimethylamine, glutamic acid, dimethicone, fragrance, phenoxyethanol, benzyl alcohol, sodium chloride, citric acid, honey extract, methylchloroisothiazolinone, methylisothiazolinone

Questions (or comments)?

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Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - Kit

head & shoulders ®

Shampoo & Conditioner

head & shoulders ®

selenium sulfide **dandruff & seborrheic dermatitis shampoo**

CLINICAL STRENGTH

DANDRUFF DEFENSE

DRY SCALP

RESCUE

head & shoulders ®

pyrithione zinc **dandruff conditioner**

CLINICAL STRENGTH

DANDRUFF DEFENSE

DRY SCALP

RESCUE

1 SELENIUM SULFIDE DANDRUFF & SEBORRHEIC DERMATITIS

SHAMPOO 13.5 FL OZ (400 mL)

1 PYRITHIONE ZINC DANDRUFF

CONDITIONER 9.1 FL OZ (270 mL)



HEAD AND SHOULDERS CLINICAL DRY SCALP RESCUE KIT

pyrithione zinc kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-673
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-673-01	1 in 1 PACKAGE, COMBINATION	03/15/2023	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE, PLASTIC	400 mL
Part 2	1 TUBE	270 mL

Part 1 of 2

HEAD AND SHOULDERS CLINICAL STRENGTH DANDRUFF DEFENSE

DRY SCALP RESCUE

selenium sulfide lotion/shampoo

Product Information

Item Code (Source) NDC:69423-523

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SELENIUM SULFIDE (UNII: Z69D9E381Q) (SELENIUM SULFIDE - UNII:Z69D9E381Q)	SELENIUM SULFIDE	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
WATER (UNII: 059QF0K00R)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
HONEY (UNII: Y9H1V576FH)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-523-40	400 mL in 1 BOTTLE, PLASTIC; 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	03/15/2023	

Part 2 of 2

HEAD AND SHOULDERS CLINICAL DANDRUFF DEFENSE DRY SCALP RESCUE

pyrithione zinc lotion

Product Information

Item Code (Source) NDC:69423-568

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	0.75 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
WATER (UNII: 059QF0KO0R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
GLUTAMIC ACID (UNII: 3KX376GY7L)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
HONEY (UNII: Y9H1V576FH)	
COCONUT (UNII: 3RT3536DHY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-568-27	270 mL in 1 TUBE; 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	01/11/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	03/15/2023	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 6/2024

The Procter & Gamble Manufacturing Company