

**HEAD AND SHOULDERS 2IN1 CLINICAL STRENGTH DANDRUFF DEFENSE
INTENSIVE ITCH RELIEF- selenium sulfide lotion/shampoo
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 ®

2in1 Clinical Strength Dandruff Defense Intensive Itch Relief

Drug Facts

Active ingredient

Selenium Sulfide 1%

Purpose

Anti-dandruff, anti-seborrheic dermatitis

Uses

helps prevent recurrence of flaking, itching, irritation, scaling and redness associated with dandruff and seborrheic dermatitis.

Warnings

For external use only.

Ask a doctor before use if you have a condition that covers a large area of the body.
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.
- caution: if used on bleached, tinted, grey, or permed hair, rinse for 5 minutes.

Inactive ingredients

Water, sodium lauryl sulfate, glycol distearate, sodium chloride, cocamidopropyl betaine, sodium citrate, dimethicone, sodium xylenesulfonate, menthol, fragrance, citric acid, sodium benzoate, tetrasodium EDTA, hydroxypropyl methylcellulose, mentha piperita (peppermint) oil, mentha arvensis leaf oil, red 4.

Questions (or comments)?

1-800-723-9569

Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 400 mL BOTTLE LABEL

head &

shoulders ®

selenium sulfide dandruff &

seborrheic dermatitis shampoo

2in1

Clinical Strength

Dandruff Defense

Intensive

Itch Relief

Extra strength formula

with Intense Cooling Menthol

13.5 FL OZ (400 mL)



SELENIUM SULFIDE DANDRUFF
& SEBORRHEIC DERMATITIS
SHAMPOO + CONDITIONER

— 2 IN 1 —
**CLINICAL
STRENGTH
DANDRUFF
DEFENSE**

**INTENSIVE
ITCH RELIEF**
EXTRA STRENGTH FORMULA
WITH INTENSE COOLING MENTHOL



13.5 FL OZ (400 mL)

head & shoulders

CLINICAL STRENGTH
SELENIUM SULFIDE DANDRUFF
& SEBORRHEIC DERMATITIS
SHAMPOO + CONDITIONER

**DANDRUFF DEFENSE
INTENSIVE ITCH RELIEF**

- Extra strength formula relieves itch associated with dandruff
- Clinically proven. Up to 100% dandruff protection*

#1 DERMATOLOGIST RECOMMENDED

Drug Facts

Active Ingredient	Purpose
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90243395

MADE IN U.S.A. of U.S. and/or
Imported Ingredients
Dist. by PROCTER & GAMBLE,
CINCINNATI, OH 45202

www.headandshoulders.com

*Satisfaction Guarantee: If you
are not satisfied return this product
with a proof of purchase for a refund
via prepaid card. Call 1-800-843-3543
for further details.

*visible flakes; with regular use
Patents: www.pg.com/patents

P&G
www.pg.com

HEAD AND SHOULDERS 2IN1 CLINICAL STRENGTH DANDRUFF DEFENSE INTENSIVE ITCH RELIEF

selenium sulfide lotion/shampoo

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-527
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
SELENIUM SULFIDE (UNII: Z 69D9E381Q) (SELENIUM SULFIDE - UNII:Z 69D9E381Q)			SELENIUM SULFIDE	1 g in 100 mL
Inactive Ingredients				
Ingredient Name				Strength
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)				
PEPPERMINT OIL (UNII: AV092KU4JH)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
WATER (UNII: 059QF0KO0R)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
GLYCOL DISTEARATE (UNII: 13W7MDN21W)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SODIUM XYLENESULFONATE (UNII: G4LZF950UR)				
DIMETHICONE (UNII: 92RU3N3Y1O)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
FD&C RED NO. 4 (UNII: X3W0AM1JLX)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-527-40	400 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)