

**HEAD AND SHOULDERS CLINICAL STRENGTH DANDRUFF DEFENSE SENSITIVE  
SCALP- selenium sulfide lotion/shampoo  
The Procter & Gamble Manufacturing Company**

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**Head and Shoulders** ®

**Clinical Strength Dandruff Defense Sensitive Scalp Shampoo**

***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, sodiumchloride, cocamidopropyl betaine,

sodium citrate, sodium xylenesulfonate, dimethicone, fragrance, citric acid, sodium benzoate, tetrasodium EDTA, hydroxypropyl methylcellulose, aloe barbadensis leaf juice

**Questions (or comments)?**

**1-800-723-9569**

Dist. by PROCTER & GAMBLE,  
CINCINNATI, OH 45202

**PRINCIPAL DISPLAY PANEL - BOTTLE LABEL**

head &

shoulders ®

selenium sulfide dandruff &

seborrheic dermatitis shampoo

Clinical Strength

Dandruff Defense

Sensitive

Scalp

13.5 FL OZ (400 mL)



SELENIUM SULFIDE DANDRUFF & SEBORRHEIC DERMATITIS SHAMPOO

CLINICAL STRENGTH DANDRUFF DEFENSE

SENSITIVE SCALP

RELIEVES SCALP IRRITATION\* INFUSED WITH ALOE VERA



13.5 FL OZ (400 mL)

head & shoulders

CLINICAL STRENGTH SELENIUM SULFIDE DANDRUFF & SEBORRHEIC DERMATITIS SHAMPOO

DANDRUFF DEFENSE SENSITIVE SCALP

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

#1 DERMATOLOGIST RECOMMENDED

Drug Facts

Active ingredient	Purpose
Selenium sulfide 1% . . . . .	Anti-dandruff, Anti-seborrheic dermatitis

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

Warnings

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- When using this product
  - avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

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Questions (or comments)? 1-800-723-9569



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

www.headandshoulders.com

\*associated with dandruff \*100% dandruff protection satisfaction guarantee: if you are not satisfied, return this product with a proof of purchase for a refund via prepaid card.

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



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HEAD AND SHOULDERS CLINICAL STRENGTH DANDRUFF DEFENSE SENSITIVE SCALP

selenium sulfide lotion/shampoo

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:69423-642

**Route of Administration** TOPICAL

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

Ingredient Name	Strength
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)	
<b>GLYCOL DISTEARATE</b> (UNII: 13W7MDN21W)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>SODIUM XYLENESULFONATE</b> (UNII: G4LZF950UR)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-642-40	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/22/2022	09/01/2026

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	04/22/2022	09/01/2026

**Labeler** - The Procter & Gamble Manufacturing Company (004238200)

Revised: 12/2025

The Procter & Gamble Manufacturing Company