

**DHS SAL 4 OZ.- dhs sal shampoo**  
**Person and Covey**

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**DHS SAL**

**WARNINGS AND PRECAUTIONS**

For external use only. Ask a physician before use if you have Psoriasis or Seborrheic Dermatitis that covers a large area of the body. Do not get into eyes. If contact occurs, rinse eyes thoroughly with water. Stop use and ask a physician if condition worsens or does not improve after regular use.

**Dosage and Administration**

Wet hair thoroughly. Apply a liberal amount of shampoo and massage into a rich lather. Allow lather to remain on scalp for several minutes. Rinse hair well and repeat application. For best results use at least twice a week or as directed by a physician. Store away from direct sunlight.

**Indications and Use**

Controls crusty, flaky buildup, while relieving scalp itching and flaking, symptomatic of Psoriasis and Seborrheic Dermatitis.

If swallowed, get medical help or contact a Poison Control Center right away.

**PURPOSE**

Controls symptoms of Psoriasis, Seborrheic Dermatitis and Dandruff

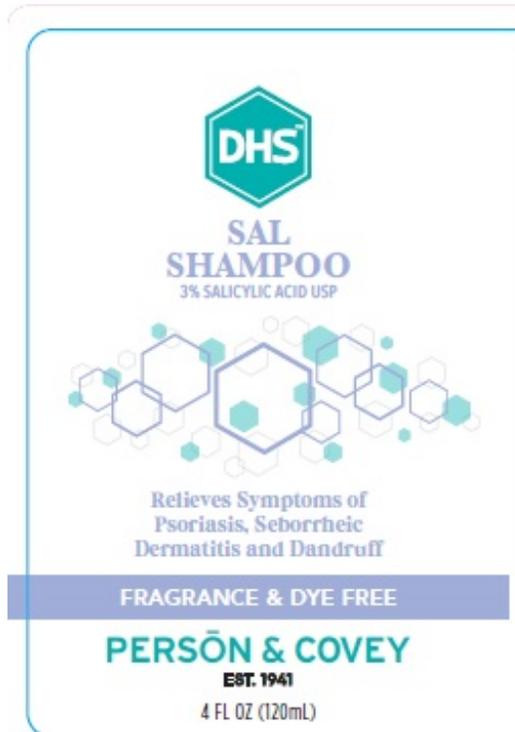
**INACTIVE INGREDIENTS**

Purified Water  
Sodium C14-16 Olefin Sulfonate  
Triethanolamine Lauryl Sulfate  
Cocamidopropyl Betaine  
Cocamidopropyl Hydroxysultaine  
PEG 8 Distearate

**Active Ingredient**

Salicylic Acid 35 mg/g

**PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**



**DHS SAL 4 OZ.**

dhs sal shampoo

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0096-0738
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>SALICYLIC ACID</b> (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	0.035 g in 1 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>SODIUM C14-16 OLEFIN SULFONATE</b> (UNII: 09W3D3YF5U)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>TRIETHANOLAMINE LAURYL SULFATE</b> (UNII: E8458C1KAA)	
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)	
<b>COCAMIDOPROPYL HYDROXYSULTAINE</b> (UNII: 62V75NI93W)	
<b>PEG-8 DISTEARATE</b> (UNII: 7JNC8VN07M)	

**Packaging**

		<b>Marketing Start</b>	<b>Marketing End</b>
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0096-0738-04	123.84 g in 1 BOTTLE; Type 0: Not a Combination Product	06/01/1995	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	06/01/1995	

**Labeler** - Person and Covey (008482473)

### Establishment

Name	Address	ID/FEI	Business Operations
Person and Covey		008482473	manufacture(0096-0738)

Revised: 11/2025

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