# SULFUR 8 SCALP THERAPY MEDICATED DANDRUFF CONTROL SCALP OIL SERUM- sulfur suspension J. Strickland and Co. 

## Sulfur 8 Scalp Therapy Medicated Dandruff Control Scalp Oil Serum

## Drug Facts

## Active Ingredient

Sulfur, 2.5\%

## Purpose

Antidandruff

## Use

Controls scalp itching an flaking due to 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 consult a doctor if

- if skin irritation develops or increases.
- condition worsens or does not improve after regular use as directed.


## Keep out of reach of children.

If swallowed, get medical help or call a Poison Control Center at once.
Flammability is increased by proudct build-up. Keep hair away from sparks, flame, extreme heat or lit tobacco. Hair is Flammable.

## Directions

- Shake well before using.
- For best results, use at least twice a week, or as directed by a doctor.
- Before shampooing your hair, apply a small amount to the scalp in several areas. Rub in well. Wait 15 minutes. Shampoo thoroughly.


## Inactive Ingredients

Mineral Oil (Paraffinum Liquidum), Polysorbate 85, Disteardimonium Hectorite, Propylene Glycol, Benzyl Alcohol, Fragrance (Parfum)

Package Labeling:



## SULFUR 8 SCALP THERAPY MEDICATED DANDRUFF CONTROL SCALP OIL SERUM

sulfur suspension

| Product Information |  |  |  |
| :---: | :---: | :---: | :---: |
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:12022-034 |
| Route of Administration | TOPICAL |  |  |
| Active Ingredient/Active Moiety |  |  |  |
| Ingredient Name |  | Basis of Strength | Strength |
| SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70) |  | SULFUR | 25 mg in 1 mL |
| Inactive Ingredients |  |  |  |
| Ingredient Name |  |  | Strength |
| MINERAL OIL (UNII: T5L8T28FGP) |  |  |  |
| POLYSORBATE 85 (UNII: A7F3N56197) |  |  |  |
| DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L) |  |  |  |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) |  |  |  |
| BENZYL ALCOHOL (UNII: LKG8494WBH) |  |  |  |


| \# Item Code | Package Description | ivarketing start Date | iviarketing mnu Date |
| :---: | :---: | :---: | :---: |
| $\begin{aligned} & 1 \begin{array}{l} \text { NDC:12022-034- } \\ 00 \end{array} \end{aligned}$ | 1 in 1 BOX | 01/01/2020 |  |
| 1 | 81 mL in 1 BOTTLE; 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 | 01/01/2020 |  |

Labeler - J. Strickland and Co. (007023112)

