GLOVERS DANDRUFF CONTROL MED., FLORAL- sulfur suspension
J. Strickland & Co.

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.

Glover's Dandruff Control Med., Floral

Active Ingredients:

Sulfur, 2.5%

Purpose

Antidandruff

Uses:

Controls scalp itching and flaking due to dandruff

Warnings:

For External Use Only.

When using this product

- do not get into 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.

Keep out of reach of children

If swallowed, get medical help or call a poison control center at once.

Flammable

Keep away from heat and open flame

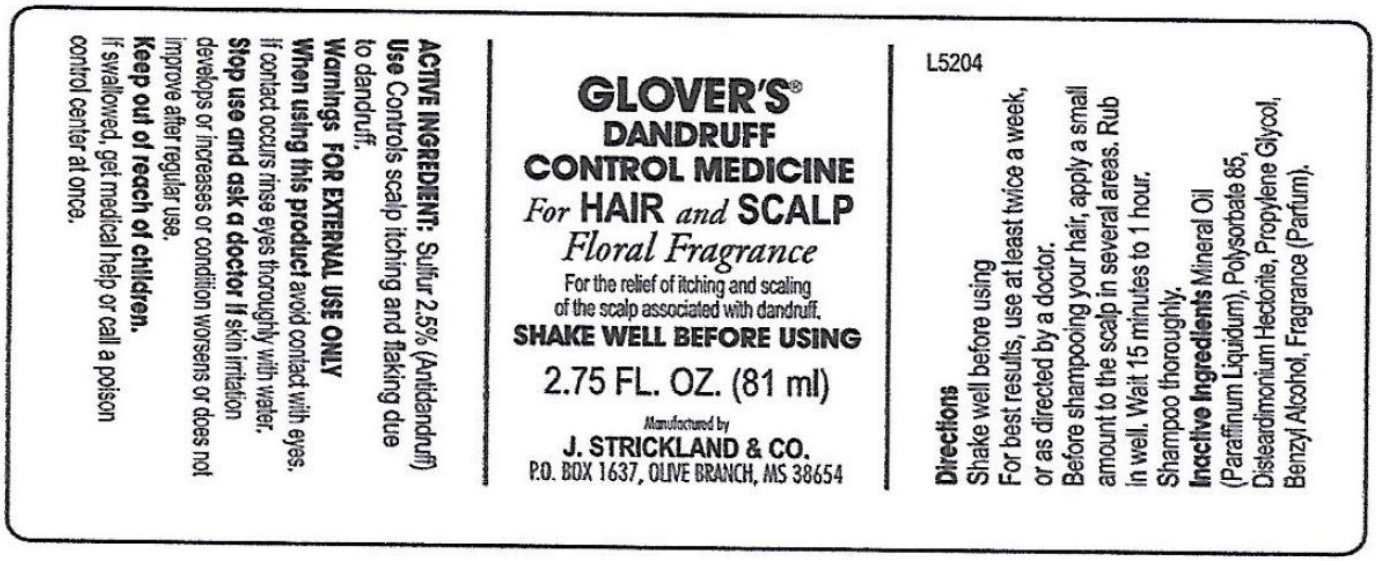
Directions:

- Shake well before using.
- For best results, use at leats twice a week, or as directed by a doctor.
- Before shampooimg your hair, apply a small amount to the scalp in several areas. Rub in well. Wait 15 minutes to 1 hour Shampoo thoroughly

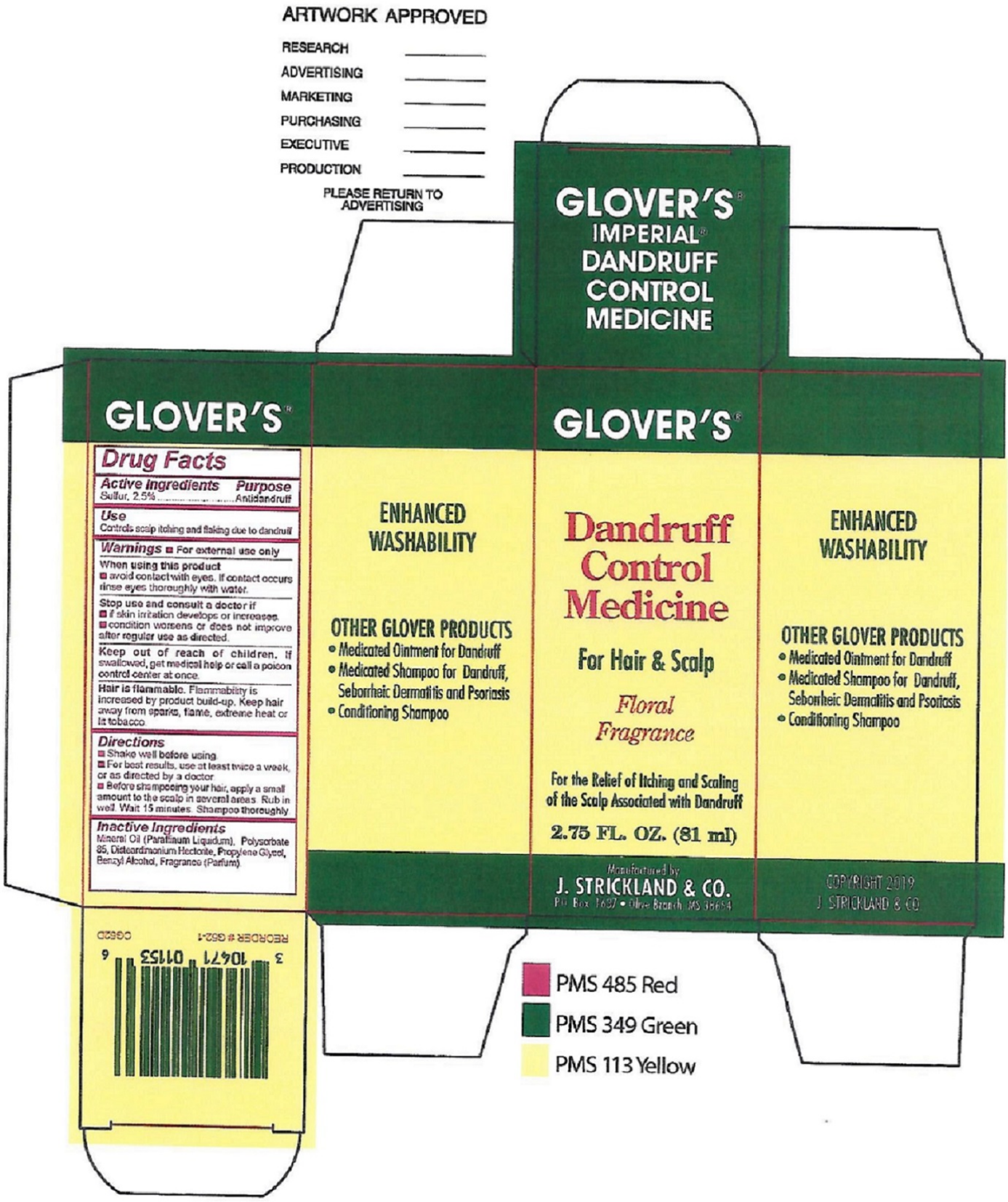
Inactive Ingredients:

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

Package Labeling Bottle



Package Labeling Carton



GLOVERS DANDRUFF CONTROL MED., FLORAL

sulfur suspension

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:12022-008
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12022-008-00	1 in 1 CARTON	11/05/2001	
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 final	part358H	11/05/2001	

Labeler - J. Strickland & Co. (007023112)

Registrant - J. Strickland & Co. (007023112)

Establishment

Name	Address	ID/FEI	Business Operations
J. Strickland & Co.		007023112	manufacture(12022-008) , pack(12022-008) , label(12022-008)

Revised: 11/2019 J. Strickland & Co.