

**EQUATE MEDICATED DANDRUFF- selenium sulfide liquid**  
**WAL-MART STORES INC**

*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.*

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**DRUG FACTS**

**ACTIVE INGREDIENT**

SELENIUM SULFIDE 1%

**PURPOSE**

ANTI-DANDRUFF

**USES**

FOR RELIEF OF FLAKING AND ITCHING ASSOCIATED WITH DANDRUFF AND SEBORRHEIC DERMATITIS AND TO HELP PREVENT THE CHANCE OF RECCURENCE

**WARNINGS**

FOR EXTERNAL USE ONLY

**ASK A DOCTOR BEFORE USING IF YOU HAVE**

SEBORRHEIC DERMATITIS IN AREAS OTHER THAN THE SCALP

**WHEN USING THIS PRODUCT**

- AVOID CONTACT WITH EYES
- IF PRODUCT GETS INTO EYES, RINSE THOROUGHLY WITH WATER
- FOR USE ON COLOR TREATED OR PERMED HAIR, RINSE THOROUGHLY

*STOP USE AND ASK A DOCTOR IF*

CONDITION WORSENS OR DOES NOT IMPROVE AFTER REGULAR USE OF THIS PRODUCT AS DIRECTED

**KEEP OUT OF REACH OF CHILDREN**

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

**DIRECTIONS**

- SHAKE WELL, APPLY SHAMPOO, RINSE THOROUGHLY
- FOR BEST RESULTS USE AT LEAST TWICE A WEEK OR AS DIRECTED BY A DOCTOR

**OTHER INFORMATION**

STORE AT ROOM TEMPERATURE

**INACTIVE INGREDIENTS**

WATER (AQUA), SODIUM LAURETH SULFATE, TEA-LAURYL SULFATE, COCAMIDOPROPYL BETAINE, ACRYLATES COPOLYMER, CITRIC ACID, FRAGRANCE, AMMONIUM CHLORIDE, DMDM HYDANTOIN, MENTHOL, SODIUM HYDROXIDE, MAGNESIUM ALUMINUM SILICATE, HYDROXYPROPYL METHYLCELLULOSE, BLUE 1 (CI 42090), RED 33 (CI 17200)

**QUESTIONS?**

CALL: 1-888-287-1915

**LABEL COPY**



**EQUATE MEDICATED DANDRUFF**

selenium sulfide liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:49035-719
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SELENIUM SULFIDE (UNII: Z69D9E381Q) (SELENIUM SULFIDE - UNII:Z69D9E381Q)	SELENIUM SULFIDE	10 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
TEA-LAURYL SULFATE (UNII: E8458C1KAA)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
AMMONIUM CHLORIDE (UNII: 01Q9PC255D)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
MENTHOL (UNII: L7T10EIP3A)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-719-11	325 mL in 1 BOTTLE, PLASTIC		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	08/17/2014	

**Labeler** - WAL-MART STORES INC (051957769)

**Registrant** - APOLLO HEALTH AND BEAUTY CARE (201901209)

### Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(49035-719)