

WECARE ANTI-DANDRUFF- antidandruff shampoo liquid
Dynarex Corporation

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.

1398 Antidandruff Shampoo NDC 67777-139-80

Active ingredient

Pyrithione Zinc 0.5%

Purpose

Anti-Dandruff

Use

Helps eliminate irritation and flaking associated with dandruff

Warnings

For External Use Only

When using this product

- Avoid contact with eyes
- If contact occurs, rinse eyes with plenty of water

Stop use and ask a doctor if

condition does not improve or worsens after regular use of this product as directed.

Keep out of reach of children

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

Directions

- wet hair
- lather
- massage onto scalp
- rinse well
- repeat if desired
- for best results, use regularly

Inactive ingredients

Acrylates Copolymer, Amyl Cinnamal, Benzyl Alcohol, Benzyl Salicylate, CI 42090, Citric Acid, Cocamide MEA, Cocamidopropyl Betaine, Coco-Glucoside, Dimethiconol, Glycerol Oleate, Glycol Distearate, Methylchloroisothiazolinone, Methylisothiazolinone, Perfume, Polyquaternium-10, Sodium Benzoate, Sodium Chloride, Sodium Hydroxide, Sodium Laureth Sulfate, Styrene/Acrylamide Copolymer, TEA-Dodecylbenzenesulfonate, Water

Label



DANDRUFF SHAMPOO

Relieves Itching and Flaking

12 fl. oz.
(355 mL)

Reorder No. 1398

Drug Facts

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Questions?
1-888-DYNAREX Monday-Friday, 9AM-5PM EST



Manufactured for: Dynarex Corporation
10 Glenshaw Street • Orangeburg, NY 10962
USA • www.dynarex.com

Made in Türkiye R230223
NDC# 67777-139-81 Mfg Lic No-13065



SYMBOL GLOSSARY
For an explanation of symbols used in Dynarex packaging, visit dynarex.com/symbols.php



Dandruff Shampoo

WECARE ANTI-DANDRUFF

antidandruff shampoo liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67777-139
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	5 g in 1000 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
.ALPHA.-AMYLCINNAMALDEHYDE (UNII: WC51CA3418)	
WATER (UNII: 059QF0KO0R)	
STYRENE/ACRYLAMIDE COPOLYMER (MW 500000) (UNII: 5Z4DPO246A)	
DIMETHICONOL (2000 CST) (UNII: T74O12AN6Y)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
BENZYL SALICYLATE (UNII: WAO5MKN9TU)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
GLYCERYL MONOOLEATE (UNII: C4YAD5F5G6)	
POLYQUATERNIUM-10 (1000 MPA.S AT 2%) (UNII: GMR4PEN8PK)	
TRIETHANOLAMINE DODECYLBENZENESULFONATE (UNII: 8HM7ZD48HN)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCO GLUCOSIDE (UNII: ICS790225B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-139-80	12 in 1 CASE	10/11/2018	

1	NDC:67777-139-81	355 mL in 1 BOTTLE, PLASTIC; 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		10/11/2018	

Labeler - Dynarex Corporation (008124539)

Revised: 2/2023

Dynarex Corporation