PERT PLUS FOR MEN DAILY DANDRUFF- pyrithione zinc shampoo, suspension
Idelle Labs, Ltd

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.

Pert Plus For Men

Pert Plus for Men Daily Dandruff

Pert Plus for Men Daily Dandruff 2 in 1 Shampoo Plus Conditioner
controls dandruff with pyrithione zinc

with strengthening ingredients
MM1

Active Ingredient

Zinc Pyrithione 1.0%

Purpose:

Anti-Dandruff

Uses

Controls the symptoms of dandruff.

Warning:

For external use only

When using this product
avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor

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

Keep this and all drugs out of reach of children.

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

Directions:
For best results, use at least twice a week or as directed by a doctor. Gently massage into wet hair, lather and rinse. Repeat if desired.

Inactive Ingredients:

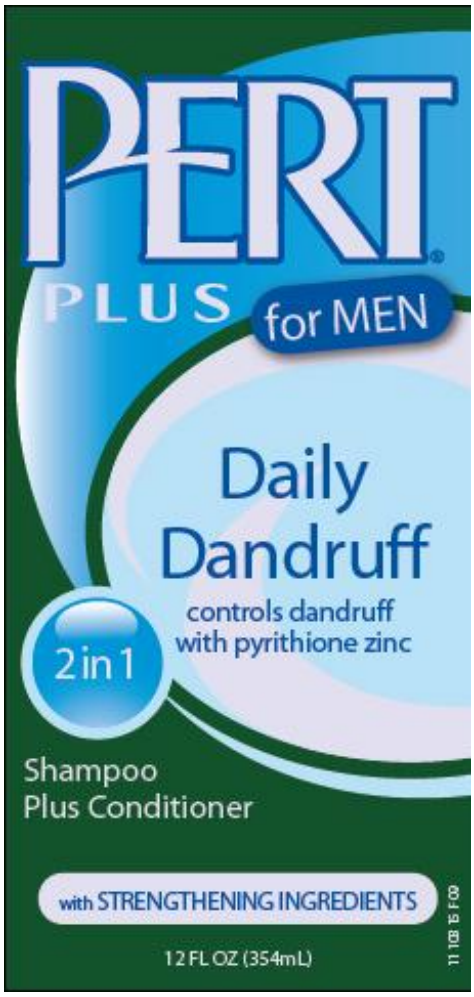
Water, ammonium laureth sulfate, ammonium lauryl sulfate, glycol distearate, dimethicone, hydrolyzed

vegetable protein PG-propyl silanetriol, cetyl alcohol, cocamide MEA, sodium chloride, fragrance, guar hydroxypropyltrimonium chloride, hydrogenated polydecene, sodium citrate, sodium benzoate, polyquaternium 10, PEG-7M, trimethylolpropane tricaprylate/tricaprate, citric acid, benzyl alcohol, methylchloroisothiazolinone, methylisothiazolinone, ammonium xylene sulfonate, D & C Yellow No. 10, FD & C Blue No. 1.

Questions?

1-800-487-7273 or visit us at www.pertplus.com

Made in the Canada and distributed by Idelle Labs, Ltd., El Paso, TX 79912



label

PERT PLUS FOR MEN DAILY DANDRUFF

pyrithione zinc shampoo, suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41595-5523
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	1 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
AMMONIUM LAURETH-5 SULFATE (UNII: 43ZIH89I48)	
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS)	
POLYETHYLENE GLYCOL 7000 (UNII: Q0JET65GEL)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:41595-5523-7	354 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part358H	04/01/2010	

Labeler - Idelle Labs, Ltd (128822926)

Registrant - Idelle Labs, Ltd (128822926)

Establishment

Name	Address	ID/FEI	Business Operations
Body Blue 2006 Inc.		243094112	MANUFACTURE

Revised: 12/2010 Idelle Labs, Ltd