PINWORM MEDICINE- pyrantal pamoate suspension
Reese Pharmaceutical Company

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.

Drug Facts

Active ingredient (in each mL)

Pyrantel Pamoate 144mg/mL (the equivalent of 50mg pyrantel base)

Purpose

Anthelimintic

Uses for the treatment of pinworns

Warnings Do not exceed recommended dosage

Ask a doctor before use if

- you're pregnant
- have liver disease

When using this product abdominal cramps, nausea, vomiting, diarrhea, headache, or dizziness sometimes occur after taking this drug. If any of these symptoms persist, consult a doctor.

Keep out of reach of children. In case of overdose get medical help or contact a poison control center right away.

Directions

- **shake well before use**
- **read bottle label and package insert carefully before taking this medication**
- treat the entire household unless otherwise advised
- do not repeat treatment unless directed by a doctor
- this product can be taken any time of day, with or without meals. It may be taken alone or with milk or fruit juice. Use of a laxative is not necessary prior to, during or after medication
- it signs of pinworms persist after treatment, consult a doctor

Weight	**Dosage** (taken as a single dose)
Less than 25 lb. or under 2 yrs.old	Do not use unless directed by a doctor
25-37 lb.	1/2 teaspoonful
38-62 lb.	1 teaspoonful
63-87 lb.	1 1/2 teaspoonfuls
88-112 lb.	2 teaspoonfuls
113-137 lb.	2 1/2 teaspoonfuls
138-162 lb.	3 teaspoonfuls
163-187 lb.	3 1/2 teaspoonfuls
188 lb. and over	4 teaspoonfuls

Inactive ingredients

banana flavor, citric acid, gylcerin, lecithin, magnesium aluminum silicate, methylcellulose, povidone, simethicone, sodium benzoate, sodium citrate, sodium saccharin, sorbitol, and water

PINWORM MEDICINE

pyrantal pamoate suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10956-618
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PYRANTEL PAMOATE (UNII: 81BK194Z5M) (PYRANTEL - UNII:4QIH0N49E7)	PYRANTEL PAMOATE	144 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	

LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
METHYLCELLULOSE (100 CPS) (UNII: 4GFU244C4J)	
POVIDONE (UNII: FZ989GH94E)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	BANANA	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10956-618-01	30 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part357B	01/01/2007	

Labeler - Reese Pharmaceutical Company (004172052)

Registrant - Elge Inc (610655136)

Establishment

Name	Address	ID/FEI	Business Operations
Elge Inc		610655136	manufacture

Revised: 3/2011 Reese Pharmaceutical Company