CVS PINWORM TREATMENT - pyrantel pamoate suspension CVS PHARMACY 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.

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



CVS PINWORM TREATMENT

pyrantel pamoate suspension

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-618
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength

Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MONO HYDRATE (UNII: 2968 PHW8 QP)		
GLYCERIN (UNII: PDC6 A3C0 OX)		
LECITHIN, SO YBEAN (UNII: 1DI56 QDM62)		
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)		
METHYLCELLULOSE (100 CPS) (UNII: 4GFU244C4J)		
PO VIDO NE (UNII: FZ989 GH9 4E)		
DIMETHICO NE (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			

1	Packaging			
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-618- 02	2 in 1 CARTON	12/15/2016	
1		30 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	part357B	12/15/2016	

Labeler - CVS PHARMACY INC (062312574)

Registrant - Reese Pharmaceutical Co (004172052)

Establishment			
Name	Address	ID/FEI	Business Operations
Nexgen Pharma		079424083	manufacture(69842-618), label(69842-618)

Revised: 12/2016 CVS PHARMACY INC