PERCY MEDICINE- bismuth subsalicylate liquid
Merrick Medicine Company, Inc

PERCY MEDICINE

Drug Facts Active Ingredient

Active Ingredients

Bismuth Subsalicylate

Purpose

Antidiarrheal

Warnings

Children and teenagers that have or are recovering from Chicken Pox, flu systems, or flue should NOT use this product. If nausea, vomiting, or fever occur, consult a doctor because these symptoms could be an early sign of Reye Syndrome, a rare but serious condition.

Allergy Alert: Contains Salicylates. Do Not take if you are - allergic to salicylates (including aspirin) - taking other salicylate products.

Do not use if you have - blood or black stool - an ulcer - bleeding problem.

Ask a doctor before use if you are taking any drug for anti coagulation (thinning of the blood), diabetes,gout, arthritis. - When using this product a temporary but harmless darkening of the stool and/or tongue may occur.

Stop use and consult a doctor if - symptoms get worse - ringing in the ears or loss of hearing occurs - diarrhea lasts more that 2 days.

if pregnant or breast feeding *ask a professional before use*. In case of accidental overdose, get medical help or contact a Poison Control Center immediately.

Inactive Ingredients

INACTIVE INGREDIENTS: CALCIUM HYDROXIDE, CITRIC ACID, ETHYL ALCOHOL, GLYCERINE, GUM ARABIC, OIL OF CINNAMON, OIL OF ORANGE, POTASSIUM CARBONATE, RHUBARB FLUID EXTRACT, SUGAR, PURIFIED WATER

Directions

(shake well before using)

Adults and children 12 yrs and over : 2 teaspoonfuls every hour as needed , not to exceed 4 doses in 24 hours and not more than 2 days.

Children under 12: Consult a doctor.

Keep out of reach of children

Keep out of reach of children.

Uses:

Relieves diarrhea, reduces number of bowel movements.

Helps firm stool.

Relieves nausea, heartburn, fullness due to overindulgence of food and drink.

Package Label

PERCY MEDICINE
FOR DIARRHEA
ALCOHOL 5%
90 ml/ 3 fl oz

PERCY MEDICINE

bismuth subsalicylate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0322-2222
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BISMUTH SUBSALICYLATE (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ)	BISMUTH SUBSALICYLATE	1050 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CALCIUM HYDROXIDE (UNII: PF5DZW74VN)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
ACACIA (UNII: 5C5403N26O)	
ORANGE OIL (UNII: AKN3KSD11B)	
CINNAMON OIL (UNII: E5GY4I6YCZ)	
POTASSIUM CARBONATE (UNII: BQN1B9B9HA)	
RHUBARB (UNII: G280W4MW6E)	
SUCROSE (UNII: C151H8M554)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0322-2222-00	1 in 1 BOX	09/09/2019	
1	NDC:0322-2222-03	90 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0322-2222-03	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/09/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M008	09/09/2019	

Labeler - Merrick Medicine Company, Inc (007331838)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(0322-2222) , manufacture(0322-2222) , pack(0322-2222) , label(0322-2222)

Revised: 12/2023 Merrick Medicine Company, Inc