CURELIEF- diphenhydramine hcl solution
Wittman Pharma, Inc.

CuRELIEF

Directions: Take every 4-6 hours, or as directed by doctor. * Do NOT exceed the recommended dose

Age	Dose
Adults and Children 12 years of age and over	take 10 to 20 mL (2 to 4 tsp), not to exceed 300 mg in 24 hours
Children 6 to under 12 years of age	take 5 to 10 mL (1 to 2 tsp) not to exceed 150 mg in 24 hours
Children under 6 years of age	Consult a doctor

Warnings:

Do not use: ■ with any other product containing diphenhydramine, even one used on skin ■ to make a child sleepy

Ask a doctor before use if you have: ■ a breathing problem such as emphysema or chronic bronchitis ■ glaucoma ■ trouble urinating due to the enlarged prostate gland

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.

When using this product: ■ excitability might occur, especially in children ■ marked drowsiness may occur ■ avoid alcoholic drinks ■ alcohol, sedatives, and tranquilizers may increase drowsiness effect ■ be careful when driving a motor vehicle or operating machinery

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Inactive Ingredients: Citric Acid, Flavor, Methylparaben, Monoammonium

Glycyrrhizinate, Potassium Citrate, Propylene Glycol, Propylparaben, Purified Water, Sorbitol, Sucralose

Uses: Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: ■ runny nose ■ sneezing ■ itchy, watery eyes ■ itching of the nose or throat ■ Temporarily relieves these symptoms due to the common cold: ■ runny nose ■ sneezing

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Purpose: Antihistamine

Active Ingredient (in each 5 mL tsp)

Diphenhydramine HCL, 12.5 mg

Drug Facts

CURELIEF

diphenhydramine hcl solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83335-103

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	12.5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
AMMONIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SORBITOL (UNII: 506T60A25R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83335-103-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2024	
2	NDC:83335-103-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/01/2024	

Labeler - Wittman Pharma, Inc. (830980947)

Establishment

Name	Address	ID/FEI	Business Operations

Wittman Pharma, Inc.		830980947	manufacture(83335-103) , analysis(83335-103) , label(83335-103)

Revised: 4/2024 Wittman Pharma, Inc.