DIMETAPP- brompheniramine maleate, pseudoephedrine hydrochloride elixir GlaxoSmithKline Consumer Healthcare Holdings (US) LLC

Dimetapp®

Active ingredients (in each Purpose 5 mL tsp)

Brompheniramine maleate, USP Antihistamine

l mg

Pseudoephedrine HCl, USP 15 Nasal decongestant

mg

Uses

- temporarily relieves nasal and sinus congestion due to the common cold, hay fever or other upper respiratory allergies, or associated with sinusitis
- temporarily relieves these symptoms due to hay fever (allergic rhinitis):
 - o runny nose
 - o sneezing
 - o itchy, watery eyes
 - o itching of the nose or throat
- temporarily restores freer breathing through the nose

Warnings

Do not use if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- glaucoma
- a breathing problem such as emphysema or chronic bronchitis

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

When using this product

- do not use more than directed
- drowsiness may occur

- avoid alcoholic beverages
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor if

- you get nervous, dizzy, or sleepless
- symptoms do not get better within 7 days or are accompanied by fever

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

Keep out of reach of children. In case of overdose, get medical help.

Other information

- store at 20-25°C (68-77°F)
- not a USP elixir
- dosage cup provided
- Pleasant tasting children find its great grape taste easy to take.
- Adults like it, too especially those who have difficulty swallowing tablets or capsules.

Directions

do not take more than 4 doses in any 24-hour period

age	dose
adults and children 12 years	4 tsp every 4 hours
and over	
children 6 to under 12 years	2 tsp every 4 hours
children under 6 years	Do not use

Inactive ingredients

artificial flavor, citric acid, FD&C blue no. 1, FD&C red no. 40, glycerin, high fructose corn syrup, propylene glycol, saccharin sodium, sodium benzoate, sorbitol, purified water

PRINCIPAL DISPLAY PANEL - 118 mL Bottle Carton

Dimetapp® **Elixir**NASAL DECONGESTANT
ANTIHISTAMINE

GREAT GRAPE TASTE FOR CHILDREN & ADULTS RELIEVES COLD AND ALLERGY SYMPTOMS

NASAL CONGESTION

RUNNING NOSE ITCHY, WATERY EYES SNEEZING 4 FL. OZ. (118 mL)



DIMETAPP

brompheniramine maleate, pseudoephedrine hydrochloride elixir

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0031-2231
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BROMPHENIRAMINE MALEATE (UNII: IXA7C9ZN03) (BROMPHENIRAMINE - UNII: H57G17P2FN)	BROMPHENIRAMINE MALEATE	1 mg in 5 mL	
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	15 mg in 5 mL	

Inactive Ingredients			
Ingredient Name	Strength		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SORBITOL (UNII: 506T60A25R)			
WATER (UNII: 059QF0KO0R)			

Product Characteristics			
Color	PURPLE	Score	
Shape		Size	
Flavor	GRAPE	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0031-2231- 89	1 in 1 CARTON	07/09/2015	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0031-2231- 86	1 in 1 CARTON	07/09/2015	
2		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:0031-2231- 88	1 in 1 CARTON	07/09/2015	
3		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information					
Marketing	Application Number or Monograph	Marketing Start	Marketing End		
Category	Citation	Date	Date		

Export only 07/09/2015

Labeler - GlaxoSmithKline Consumer Healthcare Holdings (US) LLC (079944263)

Revised: 1/2024 GlaxoSmithKline Consumer Healthcare Holdings (US) LLC