

G-P-TUSS DXP - dextbrompheniramine, dextromethorphan, and phenylephrine syrup **Kramer Novis**

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 ingredients (in each 5mL tsp)	Purpose
Dextbrompheniramine Maleate, 2 mg	Antihistamine
Dextromethorphan Hydrobromide, 20 mg	Cough suppressant
Phenylephrine HCl, 10 mg	Nasal Decongestant

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
 - sneezing
 - itchy nose or throat
 - runny nose
 - itchy, watery eyes
 - nasal congestion
- temporarily controls cough due to minor throat and bronchial irritation associated with inhaled irritants
- temporarily restores freer breathing through nose

Warnings

Do not use

- in a child who is 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 child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.

Ask a doctor before use if the child has

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or occurs with smoking, asthma, chronic bronchitis or emphysema.

Ask a doctor or pharmacist before use if the child is taking sedatives or tranquilizers

When using this product

- do not exceed recommended dosage

- excitability may occur, especially in children
- may cause marked drowsiness
- sedatives and tranquilizers may increase the drowsiness effect

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- new symptoms occur
- symptoms do not improve within 7 days or are accompanied by fever
- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

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

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away.

Directions

- do not take more than 6 doses in any 24-hour period
- **EVERY 4 HOURS**
- Adults and Children 12 years of age and older: 5 mL (1 tsp)
- Children 6 to under 12 years of age: (2.5 mL (1/2 tsp))
- Children under 6 years of age: consult physician

Other information

- Tamper evident feature: Do not use if inner seal is torn, broken or missing
- Store at controlled room temperature 15-30°C (59-86°F)
- Avoid excessive heat or humidity

Inactive ingredients

Ammonium glycyrrhizinate, D&C red #33, flavor, glycerin, hydroxyethylcellulose, methylparaben, polysorbate, propylparaben, purified water, and sucralose

Manufactured in the USA for Kramer Novis.
San Juan, PR 00917
Tel: (787) 767-2072 / www.kramernovis.com

PRINCIPAL DISPLAY PANEL - G-P-TUSS DXP

NDC 52083-602-16

G-P-Tuss[®] DXP

Drug Facts

Active Ingredients (in each 5 mL tsp) *Purpose*
 Dextrompheniramine Maleate, 2 mg.....Antihistamine
 Dextromethorphan HBr, 20 mg.....Cough Suppressant
 Phenylephrine HCl, 10 mg.....Nasal Decongestant

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: • sneezing • itchy nose or throat • runny nose • itchy, watery eyes • nasal congestion
- temporarily controls cough due to minor throat and bronchial irritation associated with inhaled irritants
- temporarily restores freer breathing through nose

Warnings

Do not use in a child who is 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 child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.

Ask a doctor before use if the child has

- heart disease • high blood pressure • thyroid disease • diabetes • glaucoma • cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis or emphysema.

Ask a doctor or pharmacist before use if the child is taking sedatives or tranquilizers

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Lot #

Exp. Date

**ANTI-HISTAMINE
 COUGH SUPPRESSANT
 NASAL DECONGESTANT**

Contains the same active ingredients as Panatuss[®] DXP *

ALCOHOL FREE

RASPBERRY FLAVOR

16 Fl.oz. (473 mL)



Drug Facts (continued)

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur • new symptoms occur
- symptoms do not improve within 7 days or are accompanied by fever
- coughs lasts more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. These could be signs of serious conditions.

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

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away.

Directions do not take more than 6 doses in any 24-hour period.

**EVERY
 4 HOURS**

Adults and Children 12 years of age and over	5 mL (1 tsp)
Children 6 to under 12 years of age	2.5 mL (½ tsp)
Children under 6 years of age	consult physician

Other information

- Tamper evident feature: Do not use if inner seal is torn, broken or missing.
- Store at controlled room temperature 15-30°C (59-86°F).
- Avoid excessive heat or humidity.

Inactive Ingredients

Ammonium glycyrrhizinate, D&C red #33, flavor, glycerin, hydroxyethylcellulose, methylparaben, polysorbate, propylparaben, purified water and sucralose.

* Panatuss[®] DXP is a registered Trademark of Seyer Pharmatec. This product is not manufactured, distributed or marketed by Seyer Pharmatec.

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NDC 52083-602-16

G-P-TUSS DXP

**ANTI-HISTAMINE
 COUGH SUPPRESSANT
 NASAL DECONGESTANT**

**ALCOHOL FREE
 RASPBERRY FLAVOR**

16 Fl. oz. (473 mL)

Kramer Novis

Pharmaceuticals within reach of patients

G-P-TUSS DXP

dextrompheniramine, dextromethorphan, and phenylephrine syrup

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52083-602
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXBROMPHENIRAMINE MALEATE (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII:75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	2 mg in 5 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
AMMONIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYETHYL CELLULOSE (100 MPA.S AT 2%) (UNII: R33S7TK2EP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	RASPBERRY (Raspberry Flavor)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52083-602-16	473 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	01/24/2013	

Labeler - Kramer Novis (090158395)

Revised: 1/2013

Kramer Novis