DELTUSS DP NASAL DECONGESTANT ANTIHISTAMINE CHERRY FLAVORdexchlorpheniramine maleate, pseudoephedrine hydrochloride liquid Deliz Pharmaceutical Corp

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.

DELTUSS DP Nasal Decongestant Antihistamine CHERRY Flavor

Drug Facts

Active ingredients (in each 5 mL teaspoonful)

Dexchlorpheniramine maleate 1 mg Pseudoephedrine HCl 30 mg

Purpose

Antihistamine

Nasal Decongestant

Uses

- temporarily relieves nasal congestion due to the common cold, hay fever or other respiratory allergies
- temporarily relieves runny nose, sneezing, itching of the nose or throat, and itchy, watery eyes due to hay fever (allergic rhinitis)
- 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 giving this product.

Ask a doctor before use if

you have

- heart disease
- high blood pressure

- thyroid disease
- diabetes
- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlarged prostrate gland

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

When using this product

- do not exceed recommended dosage
- marked drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase the drowsiness effect
- avoid alcoholic beverages
- use caution when driving a motor vehicle or operating machinery
- excitability may occur especially in children

Stop use and ask a doctor if

- symptoms do not improve within 7 days or are accompanied by fever
- nervousness, dizziness, or sleeplessness occur

If pregnant or breastfeeding

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 right away.

Directions

- use only with enclosed measuring cup
- do not use enclosed measuring cup for any other drug product

	2 teaspoonfuls (tsp) every 6 hours not to exceed 8 teaspoonfuls in 24 hours, or as directed by a doctor.
Children 6 to under 12 years of age:	1 teaspoonful (tsp) every 6 hours not to exceed 4 teaspoonfuls in 24 hours, or as directed by a doctor.
Children 2 to under 6 years of age:	Consult a doctor.

Other information

• Store at room temperature 15°C-30°C (59°F-86°F)

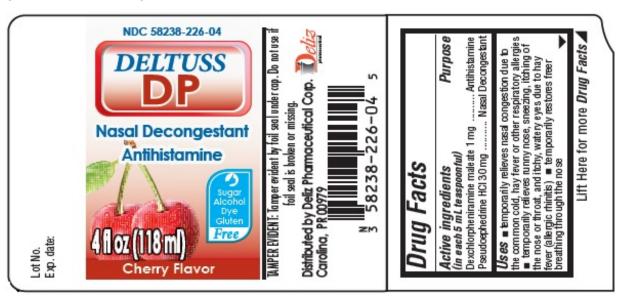
Inactive ingredients

cherry flavor, citric acid, glycerin, propylene glycol, purified water, sodium citrate, sodium saccharin, sorbitol

Questions?

Call 1-787-701-3312. You may also report serious side effects to this phone number.

DELTUSS DP Nasal Decongestant Antihistamine CHERRY Flavor 4oz/118ml (58238-226-04)



Drug Facts (continued) Warnings Do not use if you are now taking a prescription monoamine exidase 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 giving this product. Ask a doctor before use if you have me heart disease ■high blood pressure ■thyroid disease ■diabetes ■a breathing problem such as emphysema or chronic bronchitis glaucoma difficulty in urination due to enlarged prostrate gland Ask a doctor or pharmacist before use if you are taking sedatives of tranquilizers. When using this product = do not exceed recommended dosage = marked drowsiness may occur ■alcohol, sedatives, and tranquilizers may increase the drowsiness effect avoid alcoholic beverages use caution when driving a motor vehicle or operating machinery excitability may occur especially in children Stop use and ask a doctor if symptoms do not improve within 7 days or are accompanied by fever ■ nervousness, dizziness, or sleeplessness occur If pregnant or breastfeeding ask a health professional before use Keep out of reach of children. In case of overdose, get medical help or contact Poison Control Center right away. Directions ■use only with enclosed measuring cup do not use enclosed measuring cup for any other drug Adults and children 2 teaspoonfuls (tsp) every 6 hours not to exceed 8 teaspoonfuls in 24 hours, or as directed by a doctor. 12 years of age and older: Children 6 to under 1 teaspoonful (tsp) every 6 hours

not to exceed 4 teaspoonfuls in 24

hours, or as directed by a doctor.

Consult a doctor.

12 years of age:

Children 2 to under

6 years of age:

Drug Facts (continued)

Other information

■ store at room temperature 15°C-30°C (59°F-86°F)

Inactive ingredients

cherry flavor, citric acid, glycerin, propylene glycol, purified water, sodium citrate, sodium saccharin, sorbitol

Questions? Call 1-787-701-3312. You may also report serious side effects to this phone number.

DELTUSS DP NASAL DECONGESTANT ANTIHISTAMINE CHERRY FLAVOR

dexchlorpheniramine maleate, pseudoephedrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58238-226
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
DEXCHLORPHENIRAMINE MALEATE (UNII: B10YD955QW) (DEXCHLORPHENIRAMINE - UNII:3Q9Q0B929N)	DEXCHLORPHENIRAMINE MALEATE	1 mg in 5 mL		
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	30 mg in 5 mL		

Inactive Ingredients			
Ingredient Name	Strength		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
GLYCERIN (UNII: PDC6A3C0OX)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
SODIUM CITRATE (UNII: 1Q73Q2JULR)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
SORBITOL (UNII: 506T60A25R)			

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	CHERRY (Cherry Flavor)	Imprint Code	
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:58238-226- 04	1 in 1 PACKAGE	03/20/2015		
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part341	03/20/2015		

Labeler - Deliz Pharmaceutical Corp (826391138)

Establishment					
Name	Address	ID/FEI	Business Operations		
Woodfield Pharmaceutical, LLC		079398730	manufacture(58238-226)		

Revised: 11/2021 Deliz Pharmaceutical Corp