NEOTUSS PLUS- dextromethorphan hbr, chlorpheniramine maleate, phenylephrine hcl liquid DORAL PHARMAMEDICS INC dba A.G. Marin Pharmaceuticals

NeoTuss Plus Cherry Flavor

Dextromethorphan HBr 30 mg / 5 mL

Chlorpheniramine Maleate 4 mg / 5 mL

Phenylephrine HCl 7.5 mg / 5 mL

Artificial and natural flavors, citric acid, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium carboxymethylcellulose, sorbitol, and sucralose.

Antitussive, Antihistamine, and Decongestant.

Temporarily relieves cough due to minor throat and bronchial irritations associated with the common cold or inhaled irritants. Helps thin bronchial passageways of bothersome mucus.

Temporarily relieves nasal congestion due to the common cold, sinusitis, hay fever or other respiratory allergies.

Temporarily relieves sneezing, itching of the nose or throat, and itchy watery eyes due to hay fever or other respiratory allergies.

Adults and children over 12 years of age: One teaspoonful (5 mL) every 6-8 hours; do not exceed 4 teaspoonfuls in 24 hours period.

Children 6-12 years of age: 1/2 teaspoonful (2.5 mL) every 6-8 hours; do not exceed 2 teaspoonfuls in 24 hours period. Children under 6 years of age: Ask a doctor.

In case of accidental overdose seek the advice of a health professional or contact a Poison Control Center immediately.

Shake well before use.

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Do not use this product if the seal is torn, broken or missing.

Do not take this product if you have a persistent or chronic cough, such as occurs with smoking, asthma, emphysema or when cough is accompanied by excessive secretions except under the advice and supervision of a physician.

Do not take other sedatives, tranquilizers or alcohol while taking this medication.

Do not use if you are taking monoamine oxidase inhibitors or other sympathomimetics.

If you are pregnant or breast-feeding, ask a health professional before using this product.

Sympathomimetic amines should be used with caution in patients with hypertension, diabetes mellitus, heart disease, peripheral vascular disease, increased intraocular pressure, hyperthyroidism or prostatic hypertrophy. May cause drowsiness. Sedatives and tranquilizers may increase the drowsiness effect.

Pharmacist: Preserve and dispense in tight-light resistance containers as defined in the USP.

Store between 15°-30°C (59°-86°F).

Do not accept this product if the safety seal is broken or missing.

Keep a box for information.

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Package Label - Principal Display Panel

1 FL. OZ (30 mL) NDC: 12539-779-01

DIE 2188 • 1.5" x 3.5" DIE CUT 58T • 2 across, 0.125" space • 2 around, 3.625" repeat art by dan

1	NDC 12539.779.01	Drug Facts		Drug Facts (continued)	
THIS	DECONGESTANT ANTIHISTAMINE COUCH SUPPRESSANT DECUGAS PLUS Cherry Raver Alcohol, Dye, Codeine, and Sugar Free Content: 1 FL. OZ. (30 ml) Manufactured for A. Marin Pharmaceutcals FD. Box 174, Miand H. 33144	Active Ingredients: In onetsp. (Sm.) Dextrome thorphan HBr 30 mg Chlorpheniramine Makate 4 mg Phenykephrine HCl 7.5 mg	Purpose Antituzive Antihistamine Decongestant	SHARE WELL BEFORE USE DONOT USETHES PRODUCTIFSEALIS TO RN, BROKEN OR MISSING. REEP OUT OF REACH OF CHILDREN.	
		Uses : Temporarily relieves cough due to minor throat and bronchial imitations associated with the common cold or inhaled initiants. Heips thin bin nchial passage ways of bothersome muons. Temporarily relieves nearal congestion due to the common cold, sinusitis, haylever or other regimanty allegies. Temporarily relieves sneezing, itshing of the nose or throat, and itshy watery eyes due to haylever or other regimatory allegies.		In case of accidental overdose, seek the advice of a health professional or contact a Poison Control Center immediately	
SSIDE				Directions: Adults and children over 12 years of age: One teapoonful r (5 mL) every(>8 hours; do note sceed 4 is agoonful is in 24 hours period Children 6-12 years of age: 1/2 is agoonful (2,5 mL) every 6-8 hours; do note sceed 2 waspoonful is in 24 hours period. Children under 6 years of age: 4 sk a doobr.	
Ri		WAINING: Sympatho reirce to arrines should be used with caution in patients with hypertension, dizbe to mellitus, heard iseas, peripheral vaccular dizease, increased intracoular persure, hyperthymidismor purstain hypertruphy. May cause downines. Sedairues and tranquilizers rray increase the downiness effect		Other Information: Pharmacist: Preserve and dispense in tight light resistance containers as defined in the USP. Size between 15°-30°C (59°-86°F).	
				Inactive Ingredients: Artificial and naturalflators, citric acid, glycerin, methylparaben, propylene glycol, propylparaben, purfiled water, sodium.	
FIRST		Do not ta le this product f you have apersistentor chronicocough, such as occurs with smoking, asthrag, emphysema or when cough is accorpaneidoby excessive scene ions except under the advice and supermision of a physician. Do not ta le tother sedatives, tranquilizers or alcohol while taking this redication. Do not use f you are taking ronocarrine coidses inhibitors or other sympahomimetics. If you as program to be ast feeding, aska health professional before use		calboxymethylcellulose, sochitol, and sucralose.	
				Lot #/Incp. Date: Rev 12/19	
	Phone 305-593-5333 + www.ag-marin.com				

BAY TECH/LABEL

NeoTuss Plus 1oz NDC 12539-779-01

LT07866 (2188).ai (12/06/2019, 11:25 am) DORAL PHARMAMEDICS, INC. • 136308

NEOTUSS PLUS						
dextromethorphan hbr, chlo	rpheniramine maleate	, phenylephrine	e hcl liquid			
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:12		NDC:125	39-779	
Route of Administration	ORAL					
Active Ingredient/Active	e Moiety					
Ingr	Basis of	Basis of Strength				
CHLORPHENIRAMINE MALEATE UNII: 3U6IO1965U)	CHLORPHENIRAMINE MALEATE		4 mg in 5 mL			
DEXTROMETHORPHAN HYDRO (DEXTROMETHORPHAN - UNII:735	DEXTROMETHORPHAN HYDROBROMIDE		30 mg in 5 mL			
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)			PHENYLEPHRINE HYDROCHLORIDE		7.5 mg in 5 mL	
Inactive Ingredients						
	Ingredient Name			Strer	ngth	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)					300 mg in 5 mL	
SORBITOL (UNII: 506T60A25R)		1750 mg in 5 mL				
GLYCERIN (UNII: PDC6A3C0OX)		500 mg in 5 mL				
CITRIC ACID (UNII: 2968PHW8QP)					10 mg in 5 mL	
METHYLPARABEN (UNII: A2I8C7HI9T)					9 mg in 5 mL	
PROPYLPARABEN (UNII: Z8IX2SC10H)					1 mg in 5 mL	
SODIUM CARBOXYMETHYL ST	ARCH (UNII: H8AV0SQX4D)			30 mg in 5 mL		

SUCRALOSE (UNII: 9	6K6UQ3ZD4)	7.5	mg in 5 mL
NATER (UNII: 059QF	233	2331 mg in 5 mL	
Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/28/2025	
Marketing I	nformation		
Marketing	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Category			

Labeler - DORAL PHARMAMEDICS INC dba A.G. Marin Pharmaceuticals (076007996)

Establishment								
Name	Address	ID/FEI	Business Operations					
DEXTRUM LABORATORIES INC.		007392322	manufacture(12539-779)					

Revised: 2/2025

DORAL PHARMAMEDICS INC dba A.G. Marin Pharmaceuticals