

**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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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.

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.

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.

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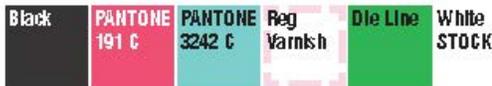
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Package Label - Principal Display Panel

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
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NDC 12539-779-01

**DECONGESTANT
ANTIHISTAMINE
COUGH SUPPRESSANT**

Neotuss Plus

Cherry Flavor
**Alcohol, Dye, Codeine,
and Sugar**
Free

Content: 1 FL. OZ. (30 ml)

Manufactured by:
 A.G. Marin Pharmaceuticals
 P.O. Box 174, Miami, FL 33144
 Phone 305-593-5333 • www.ag-marin.com

Drug Facts		Drug Facts (continued)
<p>Active Ingredients: In one tsp. (5mL)</p> <p>Dextromethorphan HBr 30 mg Chlorpheniramine Maleate 4 mg Phenylephrine HCl 7.5 mg</p> <p>Uses: Temporarily relieves cough due to minor throat and bronchial irritations associated with the common cold or inhaled irritants. Helps thin bronchial passage ways 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.</p> <p>WARNING: 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.</p> <p>Do not take this product if you have a persistent or chronic cough, such as occurs with smoking, asthma, emphysema or when a cough is accompanied by excessive secretions except under the advice and supervision of a physician.</p> <p>Do not take other sedatives, tranquilizers or alcohol while taking this medication.</p> <p>Do not use if you are taking monoamine oxidase inhibitors or other sympathomimetics.</p> <p>If you are pregnant or breastfeeding, ask a health professional before use.</p>	<p>Purpose</p> <p>Antitussive Antihistamine Decongestant</p>	<p>SHAKE WELL BEFORE USE</p> <p>DO NOT USE THIS PRODUCT IF SEAL IS TORN, BROKEN OR MISSING.</p> <p>KEEP OUT OF REACH OF CHILDREN.</p> <p>In case of accidental overdose, seek the advice of a health professional or contact a Poison Control Center in your locality.</p> <p>Directions: 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.</p> <p>Other Information: Pharmacist: Please note and dispense in tight light resistance containers as defined in the USP. Store between 15°-30°C (59°-86°F).</p> <p>Inactive Ingredients: Artificial and natural flavors, citric acid, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium carboxymethylcellulose, sorbitol, and sucrose.</p>

Lot #/Exp. Date:
Rev 12/19



NeoTuss Plus 1oz NDC 12539-779-01
 LT07866 (2188).ai (12/06/2019, 11:25 am)
 DORAL PHARMAMEDICS, INC. • 136308

NEOTUSS PLUS			
dextromethorphan hbr, chlorpheniramine maleate, phenylephrine hcl liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:12539-779
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	4 mg in 5 mL
	DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	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	Strength	
	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: Z8IX2SC1OH)	1 mg in 5 mL	
	SODIUM CARBOXYMETHYL STARCH (UNII: H8AV0SQX4D)	30 mg in 5 mL	

SUCRALOSE (UNII: 96K6UQ3ZD4)	7.5 mg in 5 mL
WATER (UNII: 059QF0KO0R)	2331 mg in 5 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:12539-779-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/28/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	02/28/2025	

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: 12/2025

DORAL PHARMAMEDICS INC dba A.G. Marin Pharmaceuticals