PENTREXCILINA CHILDREN TOTAL RELIEF MULTI-SYMPTOM- acetaminophen, dextromethorphan hydrobromide, guaifenesin usp, phenylephrine hydrochloride syrup
OPMX LLC

Pentrexcilina Children TOTAL RELIEF Multi-Symptom

Active ingredients

in each 10mL 2 teaspoonful

Acetaminophen 250mg

Dextromethorphan HBr 13.33mg

Guaifenesin USP 200mg

Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal descongestant

Uses

temporarily relieves these common cold and flu symptoms:

- sinus congestion and pressure
- minor aches and pains
- nasal congestion
- cough due to minor throat and bronchial irritation
- sore throat
- headache
- temporarily reduces fever
- temporarily promtoes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

Warnings

Liver warning

This product contains acetaminophen.

Severe liver damage may occur if you take:

- more than 6 doses in 24 hours, which is the maximum daily amount

- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily while using this product

Allergy alert

Acetaminophen may cause severe skin reactions. Symptoms may include:
- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

Sore throat warning

If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

Do not use
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist
- for children under 12 years of age
- 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
- liver disease
- heart disease
- diabetes
- thyroid disease
- high blood pressure
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus)

Ask a doctor or pharmacist before use if you are
- taking the blood thinning drug warfarin
- taking sodium restricted diet

When using this product

do not exceed recommended dose (see overdose warning).

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- pain, nasal congestion, or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back, or occurs with fever, rash, or headache that lasts. 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

Keep out of reach of children

Overdose warning

Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical even if you do not notice any sings or symptms.

Directions

- **do not take more than directed (see Overdose warning)**
- do not take more than 6 doses in any 24 hour period
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by a doctor
- mL=mililiter
- tsp=teaspoonful

Age	Dose
Children 6 to under 12 years of age	10 mL or 2 tsp provided every 4 hours
Children under 6 years of age	do not use

Other information

- **each 10 mL contains:**
- sodium 10mg
- store between 20-25°C (68-77°F)
- do not refrigerate
- keep carton for complete Drug Facts

Inactive ingredients

Aloe vera, citric acid, disodium EDTA, FD&C red no. 40, hydroxyethyl cellulose, natural & artificial strawberry flavor, propylene glycol USP, purified water, sodium benzoate, sorbitol 70% USP, sucralose.

Questions?

Call 619-600-5632 MN to FRI, 9 a.m. to 6 p.m. PTZ

MAXIMUM STRENGHT

NDC 69729-789-06

PENTREXCILINA

TOTAL RELIEF

ACETAMINOPHEN

DEXTROMETORPHAN HBr

PHENYLEPHRINE HCl

GUAIFENESIN

CHILDREN

Flu

Cold &

Cough

- FEVER
- BODYACHE
- HEADACHE
- SORE THROAT

6fl oz (177ml)

VERSIÓN EN ESPAÑOL EN EL INTERIOR DE LA CAJA

Exclusively distributed by:

OPMX

Chula Vista, CA 91910

Manufactured in FDA Registered in the USA

PENTREXCILINA CHILDREN TOTAL RELIEF MULTI-SYMPTOM

acetaminophen, dextromethorphan hydrobromide, guaifenesin usp, phenylephrine hydrochloride syrup

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69729-789
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 10 mL

ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	250 mg in 10 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	13.33 mg in 10 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SORBITOL SOLUTION 70% (UNII: 8KW3E207O2)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	STRAWBERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69729-789-06	1 in 1 CARTON	01/15/2025	
1		177 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 Drug	M012	01/15/2025	

Labeler - OPMX LLC (029918743)

Establishment

Name	Address	ID/FEI	Business Operations
GADAL Laboratories, Inc		841305639	manufacture(69729-789)