ROMPE PECHO SF FLU- dextromethorphan, brompheniramine maleate, phenylephrine hydrochloride liquid
Efficient Laboratories Inc

Drug Facts

Active ingredients (in each 5 mls tsp.) **Purpose**

Dextromethorphan HBr 10 mg ..... Cough Suppressant

Brompheniramine Maleate 4 mg ..... Antihistaminic

Phenylephrine HCl 10 mg ..... Nasal Decongestant

Purpose

Antihistamine

Cough Suppressant

Nasal Decongestant

USES

- temporarily relieves runny nose, sneezing, itching of the nose or throat and itchy watery eyes due to hay fever or other upper respiratory allergies
- temporarily relieves cough and nasal congestion as may occur with the common cold

Warnings

Do not exceed recommended dosage

Do not use • If you are 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 you are taking a prescription drug that contains an MAOI; ask your doctor or pharmacist before taking this product.

Ask a doctor before use if you have • heart disease • glaucoma • high blood pressure • thyroid disease • diabetes • difficulty in urination due to an enlarged prostate gland • a persistent or chronic cough such as occurs with smoking, asthma or emphysema • a cough that is accompanied by excessive phlegm (mucus) • a breathing problem such as emphysema or chronic bronchitis

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

When using this product • may cause drowsiness • avoid alcoholic drinks • alcohol, sedatives, and tranquilizers may increase drowsiness • be careful 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, cough tends to recur, or symptoms are accompanied by a fever, rash or persistent headache. A persistent cough may be a sign of a serious condition. • if nervousness, dizziness, or sleeplessness occur

If pregnant or breast-feeding, 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:

- shake well before use
- do not exceed more than 6 doses in any 24-hour period or as directed by a doctor

Age	Dose
adults and children 12 years of age and over	5 mL (1 tsp) every 4 hours
children 6 to under 12 years of age	2.5 mL (1/2 tsp) every 4 hours
children under 6 years of age	consult a doctor

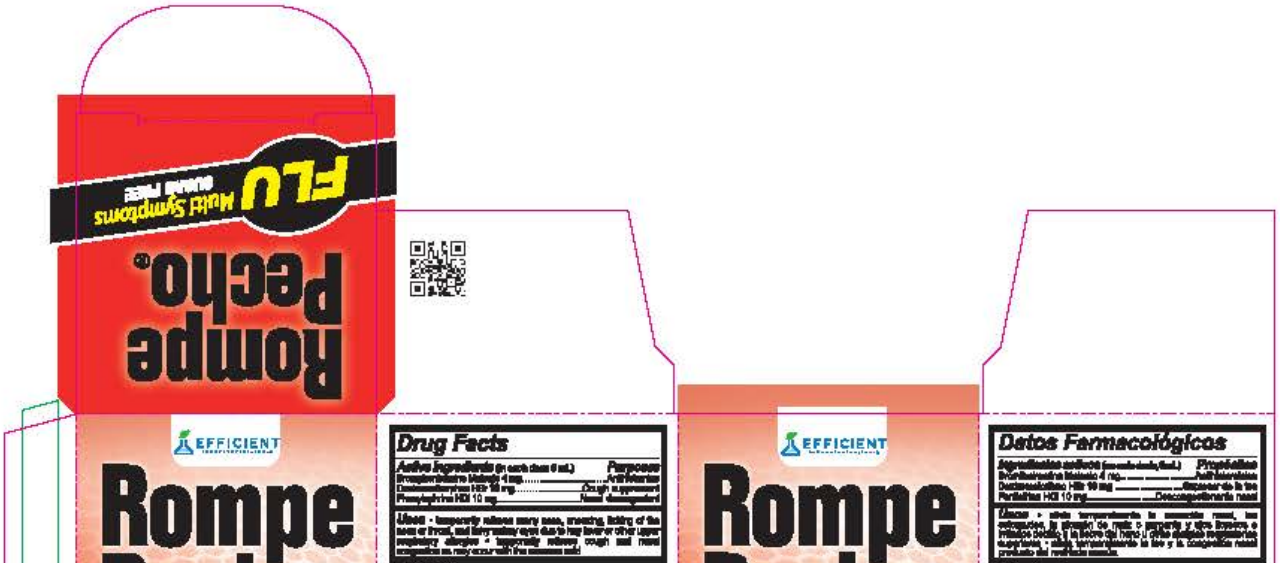
Inactive Ingredients:

Blue Cohosh root extract, Citric Acid, Echinacea root extract, Eucalyptus Oil, Ginkgo Biloba leaf extract, Glycerine, Golden Seal Root extract,
Honey Flavor, Horehound, Licorice Root extract, Menthol, Methylparaben, Mullein Leaf extract, Myrrh gum extract, Potassium Citrate,
Potassium Sorbate, Propylene Glycol, Propylparaben, Slippery Elm Bark extract, Sodium Chloride, Sucralose, Water, Wild Cherry Bark extract, Zinc Sulfate.

Questions or Comments?

305-805-3456 Monday - Friday 9AM to 5PM EST

www.efficientlabs.com



ROMPE PECHO SF FLU

dextromethorphan, brompheniramine maleate, phenylephrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58593-225

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL
BROMPHENIRAMINE MALEATE (UNII: IXA7C9ZN03) (BROMPHENIRAMINE - UNII:H57G17P2FN)	BROMPHENIRAMINE MALEATE	10 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
SUCRALOSE (UNII: 96K6UQ3ZD4)	
CAULOPHYLLUM THALICTROIDES ROOT (UNII: JTJ6HH6YEH)	
ECHINACEA, UNSPECIFIED (UNII: 4N9P6CC1DX)	
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	
GINKGO (UNII: 19FUJ2C58T)	
GOLDENSEAL (UNII: ZW3Z11D0JV)	
HONEY (UNII: Y9H1V576FH)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
MENTHOL (UNII: L7T10EIP3A)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
VERBASCUM DENSIFLORUM LEAF (UNII: 9936O846LI)	
MYRRH (UNII: JC71GJ1F3L)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
ULMUS RUBRA BARK (UNII: 91QY4PXU8Q)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PRUNUS SEROTINA BARK (UNII: 5D48E975HA)	
ZINC SULFATE (UNII: 89DS0H96TB)	
HOREHOUND (UNII: K08036XEJV)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POLYGLYCERIN-10 (UNII: P9060O936A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58593-225-06	178 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2007	10/31/2026

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M012	10/01/2007	10/31/2026

Labeler - Efficient Laboratories Inc (969044932)

Registrant - Efficient Laboratories Inc (969044932)

Revised: 3/2025 Efficient Laboratories Inc