

ROMPE PECHO DM- dextromethorphan hbr, guaifenesin liquid
Efficient Laboratories Inc

Rompe Pecho

Active Ingredients:(in each teaspoonful 5 ml.)

Purpose

Dextromethorphan Hydrobromide 10 mg Cough Suppressant
Guaifenesin 100 mg Expectorant

Purpose

Cough Suppressant

Expectorant

Uses:

- temporarily relieves cough as may occur with the common cold
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus.

Warnings

Do not exceed recommended dosage.

Ask a doctor before use if you have

- a cough that is accompanied by excessive phlegm (mucus)
- a persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema.

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 on MAOI, ask a doctor or pharmacist before taking this product.
- **Stop use and ask a doctor**
- If cough persists more than 1 week, tends to recur or is accompanied by a fever, rash or persistent headache. A persistent cough may be a sign of a serious condition.

If pregnant or breast-feeding,ask a health professional before use.

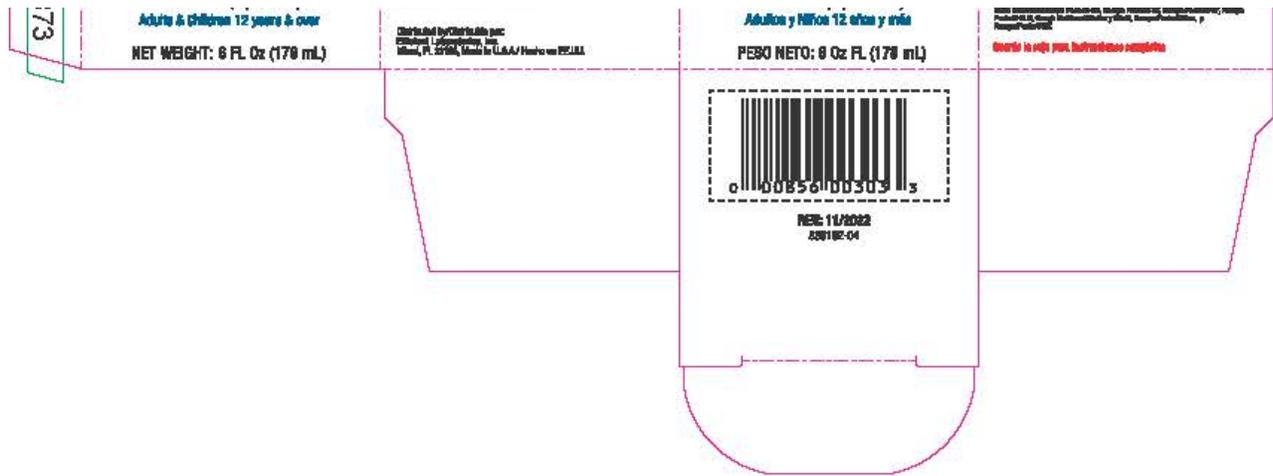
Keep out of reach of children.

In case of accidental 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



ROMPE PECHO DM

dextromethorphan hbr, guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58593-275
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
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
SUCRALOSE (UNII: 96K6UQ3ZD4)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	
CAULOPHYLLUM THALICTROIDES ROOT (UNII: JTJ6HH6YEH)	

ECHINACEA, UNSPECIFIED (UNII: 4N9P6CC1DX)
EUCALYPTUS GLOBULUS LEAF (UNII: S546YWLW6E6)
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)
ASCLEPIAS TUBEROSA ROOT (UNII: 1ZL82U5D18)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
PROPYLPARABEN (UNII: Z8IX2SC1OH)
WATER (UNII: 059QF0KOOR)
ULMUS RUBRA BARK (UNII: 91QY4PXU8Q)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
SPEARMINT (UNII: J7I2T6IV1N)
PRUNUS SEROTINA BARK (UNII: 5D48E975HA)
ZINC SULFATE (UNII: 89DS0H96TB)
GLYCERIN (UNII: PDC6A3C0OX)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58593-275-06	178 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2001	
2	NDC:58593-275-10	10 mL in 1 POUCH; Type 0: Not a Combination Product	10/01/2001	05/16/2023
3	NDC:58593-275-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2015	05/16/2023
4	NDC:58593-275-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2015	05/16/2023

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/01/2001	

Labeler - Efficient Laboratories Inc (969044932)

Registrant - Efficient Laboratories Inc (969044932)