RYMED- dexchlorpheniramine maleate and phenylephrine hydrochloride tablet, coated EDWARDS PHARMACEUTICALS, INC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

RYMED TABLETS

Drug Facts

Active Ingredients	Purpose	
(in each tablet)		
Dexchlorpheniramine Maleate 2 mg	Antihistamine	
Phenylephrine HCl 10 mg	Nasal Decongestant	

Uses

temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other upper respiratory allergies:

- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes
- nasal congestion
- reduces swelling of nasal passages

Warnings

Do not exceed recommended dosage.

Do not use this product

• 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

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma
- heart disease
- high blood pressure
- thyroid disease
- diabetes mellitus
- difficulty in urination due to enlargement of the prostate gland

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

When using this product

- excitability may occur, especially in children
- may cause drowsiness
- alcohol, sedatives and tranquilizers may increase the drowsiness effect
- avoid alcoholic beverages
- use caution when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- symptoms do not improve within 7 days or are accompanied by a fever
- new symptoms occur

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

Keep out of reach of children. In case of accidental overdose, seek professional help or contact a Poison Control Center immediately.

Directions

Do not exceed recommended dosage.

Adults and children 12 years of age and over:	1 tablet every 4 hours, not to exceed 6 tablets in 24 hours, or as directed by a doctor
Children 6 to under 12 years of age:	1/2 tablet every 4 hours, not to exceed 3 tablets in 24 hours, or as directed by a doctor
Children under 6 years of age:	Consult a doctor.

Inactive ingredients

Magnesium Stearate, Microcrystalline Cellulose, Sodium Starch Glycolate

Questions or Comments?

Call 1-800-543-9560

PRINCIPAL DISPLAY PANEL - 100 Tablet Bottle Label

E NDC 0485-0080-01 RYMED TABLETS

Antihistamine • Nasal Decongestant

Each tablet contains:

Dexchlorpheniramine Maleate 2 mg Phenylephrine HCl 10 mg This bottle is not to be dispensed to consumer. Dispense in a tight, light-resistant container with a child-resistant cap.

Store at 59°-86° (15°-30°C) [see USP Controlled Room Temperature]. Tamper evident by foil seal under cap. Do not use if foil seal is broken or missing.

Manufactured for:

EDWARDS

Pharmaceuticals, Inc.

Ripley, MS 38663

100 tablets

Drug Facts

Active Ingredients Purpose (in each tablet)

Dexchlorpheniramine
Maleate 2 mg Antihistamine
Phenylephrine HCl

10 mg Nasal Decongestant

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■ nasal congestion ■ reduces swelling of nasal passages





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Pharmaceuticals, Inc. Ripley, MS 38663 100 tablets



Lot: Exp. Date

Drug Facts (continued)

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Drug Facts (continued)

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Questions or Comments?

Rev. 11/11

RYMED

dexchlorpheniramine maleate and phenylephrine hydrochloride tablet, coated

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Product	Intorm	ation
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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0485-0080
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Route of Administration ORAL

Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength
DEXCHLORPHENIRAMINE MALEATE (UNII: B10YD955QW) (DEXCHLORPHENIRAMINE - UNII:3Q9Q0B929N)	DEXCHLORPHENIRAMINE MALEATE	2 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS 297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength	
MAGNESIUM STEARATE (UNII: 70097M6I30)		
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)		
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)		

Product Characteristics

Color	white	Score	2 pieces
Shape	OVAL	Size	12mm
Flavor		Imprint Code	R;1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:0485- 0080-01	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/15/2011	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC monograph final	part341	11/15/2011	

Labeler - EDWARDS PHARMACEUTICALS, INC. (195118880)

Revised: 8/2023 EDWARDS PHARMACEUTICALS, INC.