ACETAMINOPHEN, CHLORPHENIRAMINE MALEATE, PHENYLEPHRINE HCLacetaminophen, chlorpheniramine maleate, phenylephrine hcl tablet OPMX LLC

Resfriol D

(Acetaminophen 500mg, Chlorpheniramine Maleate 4mg, Phenylephrine HCl 10mg Tablet)

Acetaminophen 500mg

Chlorpheniramine Maleate 4mg

Phenylephrine HCl 10mg

Pain reliever/fever reducer

Antihistamine

Nasal decongestant

magnesium stearate, povidone, silicon dioxide, sodium benzoate, sodium starch glycolate, starch, talc

- Do not take more than directed see overdose warning
- swallow whole; do not crush, chew or dissolve
- do not exceed 6 tablets per 24 hours

Adults and children 12 years of age &	1 tablet with water every 4-6 hours as
over	needed
Children 4 to 12 years of age	Ask a doctor
Children under 4 years of age	Do not use

Temporarily relieves these symptoms of hay fever or other respiratory allergies:

- Headache
- Nasal congestion
- Sinus congestion & pressure
- Runny nose and sneezing
- Minor aches & pain

Temporarily relieves these additional symptoms of hay fever:

- itching of the nose or throat
- itchy, watery eyes
- helps clear nasal passages
- helps decongest sinus opening and passages

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 6 tablets in 24 hours, which is the maximum daily amount for this product
- with other drugs contains acetaminophen
- 3 or more alcoholic drinks every day 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 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
- 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
- High blood pressure
- Thyroid disease
- Diabetes
- Trouble urinating due to enlarged prostate gland
- A breathing problem such as emphysema or chronic bronchitis

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

When using this product, do not use more than directed.

Stop use and ask a doctor if:

You get nervous, dizzy, or sleepless Pain, nasal congestion, or cough gets worse or lasts more than 7 days Fever gets worse or last more than 3 days Redness or swelling is present New symptoms occur

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.

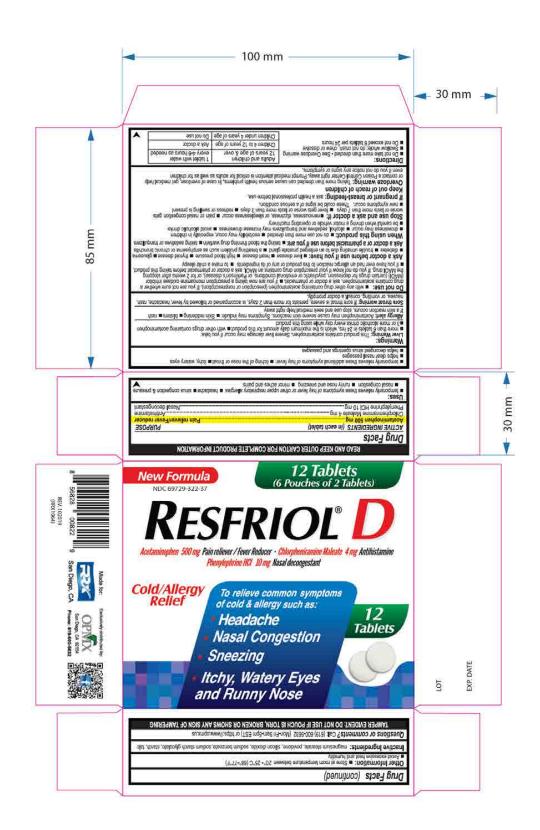
Keep out of reach of children.

OVERDOSE WARNING: Read directions carefully. In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

store at room temperature between 20-25°C (68-77°F)







HCL

acetaminophen, chlorpheniramine maleate, phenylephrine hcl tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69729-322	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	500 mg	
CHLORPHENIRAMINE MALEATE (UNII: V1Q0090J9Z) (CHLORPHENIRAMINE - UNII: 3U6I01965U)	CHLORPHENIRAMINE MALEATE	4 mg	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg	

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			
TALC (UNII: 7SEV7J4R1U)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
STARCH, CORN (UNII: O8232NY3SJ)			
POVIDONE K30 (UNII: U725QWY32X)			

Product Characteristics			
Color	white	Score	no score
Shape	ROUND (BICONVEX)	Size	13mm
Flavor		Imprint Code	S78
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69729-322- 27	2 in 1 POUCH; Type 0: Not a Combination Product	08/26/2019	
2	NDC:69729-322- 37	12 in 1 POUCH; Type 0: Not a Combination Product	08/26/2019	
3	NDC:69729-322- 38	144 in 1 POUCH; Type 0: Not a Combination Product	08/26/2019	

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date

OTC Monograph Drug	M012	08/26/2019	

Labeler - OPMX LLC (029918743)

Revised: 3/2022 OPMX LLC