CONRX ALLERGY SINUS MULTI-SYMPTOMS- acetaminophen, chlorpheniramine maleate, and phenylephrine hydrochloride tablet Eagle Distributors, 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.

ConRx Allergy Sinus Multi-Symptoms

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325mg Chlorpheniramine maleate 2mg Phenylephrine HCl 5mg

Purpose

Pain reliever/ fever reducer Antihistamine Nasal decongestant

Uses

Uses For the temporary relieves these symptoms of hay fever or other upper respiratory allergies, headache, sinus congestion and pressure, nasal congestion, runny nose and sneezing, minor aches and pains, itching of the nose or throat, itchy watery eyes, help decongest sinus openings and passages.

Warnings

- **Liver warning**: This product contains acetaminophen. Severe liver damage may occur if you take: more than 8 tablets in 24 hours, which is the maximum daily amount for this product
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

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.
- If you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

- liver disease
- heart disease
- high blood pressure
- thyroid disease

- diabetes
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis,
- glaucoma

Ask a doctor or pharmacist before use if you are taking

• the blood thinning drug warfarin

When using this product

- do not exceed recommended dosage
- excitability may occur, especially in children
- drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- pain or nasal congestion 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

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

Keep out of reach of children.

Overdose warning

In case of overdose, get medical help or contact a Poison 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.

Directions

Do not taking more than directed (see overdose warning)

	 Take 1-2 tablets every 4 hours Do not taking more than 8 tablets in 24 hour
Children under 12 years of age	Consult a doctor

Other Information

- Do not use if pouch is torn or damaged.
- Store between 15-30°C (59-86°F). Avoid excessive heat and humidity.
- See side panel for lot number and expiration date.

Inactive ingredients

Colloidal silicon dioxide, Sodium Lauryl Sulphate, Edetaet disodium, Dioctyl Sodium Sulphosuccinate, Polyvinylpyrollidone, Gelatin, Corn starch', Magnesium Stearate, Microcrystalline Cellulose, Polyethylene glycol 6000, Sodium Benzoate, Sodium Starch Glycolate, Stearic acid, Purified Talc, Titanium Dioxide, Yellow iron oxide.

PRINCIPAL DISPLAY PANEL - 50 Pouch Carton

NDC:68737-225-09

Compare to the Active Ingredients in Tylenol® Allergy Multi-Symptom*

ConRx
Allergy MULTI - SYMPTOM
Sinus
Sneezing - Runny Nose - Congestion

■ **Acetaminophen** Headache

Phenylephrine HCl
 Sinus Pressure/Nasal Congestion

• Chlorpheniramine Maleate Watery Eyes/Runny Nose

Compare to the Active Ingredients in

Tylenol® Allergy Multi-Symptom



- Acetaminophen
 Phenylephrine HCl
 Chilorpheniramine Maleate

Headache Sinus Pressure/Nasal Congestion Watery Eyes/Runny Nose

DO NOT USE WITH OTHER MEDICINES CONTAINING **ACETAMINOPHEN**

*This product is not manufactured or distributed by: McNeil Consumer Speciality Pharmaceuticals Division of McNeil PPC, Inc. Eagle Distributors, Inc., does not own the Tylenol® Allergy Multi-Symptom



Product manufctured for: Eagle Distributors, Inc. Los Angeles, CA 90011

Compare to the Active Ingredients in

Tylenol® Allergy
Multi-Symptom®

Sneezing - Runny Nose - Congestion

- Acetaminophen Phenylephrine HCI Chlorpheniramine Maleate

Headache Sinus Pressure/Nasal Congestion Watery Eyes/Runny Nose



PUSH IN TAB AND PULL OUT

Compare to the Active Ingredients in

50 Pouches of 2 Caplets Each

Compare to the Active Ingredients in Tylenol Alle Multi-Symptom Sneezing - Runny Nose - Congestion ■ Acetaminophen ■ Phenylephrine HCl ■ Chlorpheniramine Maleate Headache Sinus Pressure/Nasal Congestion Watery Eyes/Runny Nose



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- Chlorpheniramine Maleate Watery Eyes/Runny Nose Sinus Pressure/Nasal Congestion Неадасре
 - - Phenylephrine HCI
 - Acetaminophen



GNOI Allergy
Multi-Symptom:

Compare to the Active Ingredients in

Drug Facts

Active ingredients (in each tablet)

Purpose

Chlorpheniramine maleate 2 mg... Phenylephrine HCl 5 mg

Pain reliever ...Antihistamine .Nasal decongestant

USES temporarily relieves these symptoms of hay fever or other upper respiratory allergies:

- headache sinus congestion and pressure nasal congestion runny nose and sneezing minor aches and pains itching of the nose or throat itchy, watery eyes
- helps decongest sinus openings and passages

Warnings

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

- more than 8 tablets in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

- 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.
- if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

- liver disease heart disease high blood pressure thyroid disease diabetes trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis glaucoma

Ask a doctor or pharmacist before use if you are

■ taking the blood thinning drug warfarin
■ taking sedatives or tranquilizers

When using this product do not exceed recommended dosage

- excitability may occur, especially in children
 drowsiness may occur
 alcohol, sedatives and tranquilizers may increase drowsiness
 avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur pain or nasal congestion gets worse or lasts more than 7 days rever gets worse or lasts 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. Overdose warning: 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. Directions and do not take more than directed (see overdose warning) ■ take 2 tablets every 4 hours adults and children 12 years and over ■ do not take more than 8 tablets in 24 hours children under 12 years consult a doctor Other information avoid excessive heat and humidity store between 15-30°C (59-86°F) do not use if pouch is torn or open
see side panel for lot number and expiration date Inactive ingredients coloidal silicon dioxide, sodium lauryl sulphate, edetaet disodium, dioctyl sodium sulphosuccinate, polyvinylpyrolidone, gelatin, corn starch, magnesium stearate, microcrystalline cellulose, polyethylene glycol 6000, sodium benzoate, sodium starch glycolate, stearic acid, purified talc, titanium dioxide, yellow iron oxide.

Questions or comments? all 1-800-570-8650 (M-F 9 am to 5 pm PST)

Part No: 88881_2012

CONRX ALLERGY SINUS MULTI-SYMPTOMS

acetaminophen, chlorpheniramine maleate, and phenylephrine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68737-225
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Acetaminophen (UNII: 36209ITL9D) (Acetaminophen - UNII:36209ITL9D)	Acetaminophen	325 mg	
Chlorpheniramine maleate (UNII: V1Q0O9OJ9Z) (Chlorpheniramine - UNII:3U6IO1965U)	Chlorpheniramine maleate	2 mg	
Phenylephrine Hydrochloride (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV)	Phenylephrine Hydrochloride	5 mg	

Inactive Ingredients			
Ingredient Name	Strength		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)			
SODIUM LAURYL SULFATE (UNII: 368GB5141J)			
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)			
BIS(1-METHYLAMYL) SODIUM SULFOSUCCINATE (UNII: 772Y8KZU65)			
POVIDONE K90 (UNII: RDH86HJV5Z)			
GELATIN (UNII: 2G86QN327L)			
STARCH, CORN (UNII: O8232NY3SJ)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)			
POLYETHYLENE GLYCOL 6000 (UNII: 30 IQX730 WE)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)			
STEARIC ACID (UNII: 4ELV7Z65AP)			

TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

Product Characteristics			
Color	WHITE	Score	2 pieces
Shape	ROUND	Size	12mm
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68737-225-09	50 in 1 BOX		
1		2 in 1 POUCH		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	07/11/2012	

Labeler - Eagle Distributors,Inc. (929837425)

Revised: 7/2012 Eagle Distributors,Inc.