ALLERGY MULTISYMPTOM- acetaminophen, chlorpheniramine maleate, and phenylephrine hydrochloride tablet, coated RETAIL BUSINESS SERVICES , LLC.

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.

1128B-CAR-2021-1022

Drug Facts

Active ingredients (in each caplet)	Purpose		
Acetaminophen 325 mg	Pain reliever		
Chlorpheniramine maleate 2 mg	Antihistamine		
Phenylephrine HCl 5 mg	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
- temporarily relieves these additional symptoms of hay fever:
 - itching of the nose or throat
 - itchy, watery eyes
- helps clear nasal passages
- helps decongest sinus openings and passages

Warnings

Liver warning

This product contains acetaminophen. The maximum daily dose of this product is 10 caplets (3,250 mg acetaminophen) in 24 hours. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing 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.

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 sedatives or tranquilizers
- 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

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 (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

do not take more than directed (see overdose warning)

adults and children 12 years and over	 take 2 caplets every 4 hours swallow whole; do not crush, chew, or dissolve do not take more than 10 caplets in 24 hours
children under 12 years	 ask a doctor

Other information

- store between 20°-25°C (68°-77°F) in a dry place
- retain carton for complete product information and warnings

Inactive ingredients

acesulfame potassium, colloidal silicon dioxide, corn starch, croscarmellose sodium, crospovidone, flavor, iron oxide yellow, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, pregelatinized starch, propylene glycol, stearic acid, talc, titanium dioxide

PRINCIPAL DISPLAY PANEL

NDC 72476-128-02

CAREONE®

FOR ADULTS

ALLERGY MULTI-SYMPTOM

Pain Reliever - Acetaminophen

Nasal Decongestant - Phenylephrine HCL

Antihistamine - Chlorpheniramine Maleate

FOR RELIEF OF:

- Headache
- Sinus Pressure

- Nasal Congestion
- Runny Nose
 Sneezing
- Itchy, Watery Eyes

COOL TASTE

Instant Cooling Sensation

May Cause Drowsiness

Actual Size

24 CAPLETS



ALLERGY MULTISYMPTOM

acetaminophen, chlorpheniramine maleate, and phenylephrine hydrochloride tablet, coated

Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72476-128			
Route of Administration	ORAL					

Active Ingred	ient/Active Moiety			
	Ingredient Name		Basis of Streng	gth Strength
ACETAMINOPHEN	IOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPHEN			325 mg
CHLORPHENIRAMINE MALEATE (UNII: V1Q0090J9Z) (CHLORPHENIRAMINE - CHLORPHENIRAMINE JNII: 3U6I01965U) CHLORPHENIRAMINE			2 mg	
PHENYLEPHRINE	ENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - PHENYLEPHRINE II: 1WS297W6MV) PHENYLEPHRINE HYDROCHLORIDE			5 mg
Inactive Ingre	edients			
	Strength			
ACESULFAME PO	TASSIUM (UNII: 230V73Q5G9)			
SILICON DIOXIDE	(UNII: ETJ7Z6XBU4)			
CROSCARMELLOS	E SODIUM (UNII: M28OL1HH48	3)		
FERRIC OXIDE YE	LLOW (UNII: EX438O2MRT)			
MAGNESIUM STE	ARATE (UNII: 70097M6I30)			
CROSPOVIDONE (UNII: 2S7830E561)			
CELLULOSE, MICI	ROCRYSTALLINE (UNII: OP1R3	2D61U)		
POLYETHYLENE G	ILYCOL, UNSPECIFIED (UNII:	3WJQ0SDW1A)		
POLYVINYL ALCO	HOL, UNSPECIFIED (UNII: 532	B59J990)		
POVIDONE, UNSP	ECIFIED (UNII: FZ989GH94E)			
STARCH, CORN (U	INII: 08232NY3SJ)			
PROPYLENE GLYC	OL (UNII: 6DC9Q167V3)			
STEARIC ACID (UN	III: 4ELV7Z65AP)			
TALC (UNII: 7SEV7J	4R1U)			
TITANIUM DIOXID	E (UNII: 15FIX9V2JP)			
Product Char	acteristics			
Color	white (Off-White)	Score	no	score
Shape	OVAL	Size	Size 17	
Flavor	MINT	Imprint Cod	Imprint Code A	
Contains				
Packaging				
# Item Code	Package Desc	ription Ma	rketing Start M Date	arketing End Date
1 NDC:72476- 128-02	2 in 1 CARTON	10/21	/2020	
1	12 in 1 BLISTER PACK; Type 0 Product	Not a Combination		
Marketing	Information			
Marketing Category	Application Number Citatio		arketing Start M Date	larketing End Date
category				2010

Revised: 10/2021

RETAIL BUSINESS SERVICES , LLC.