

GREEN GUARD ADVANCED SINUS RELIEF- acetaminophen, phenylephrine hydrochloride tablet
Unifirst First Aid Corporation

819R Green Guard Advanced Sinus Relief

Drug Facts

Active ingredients (in each tablet)

Acetaminophen 500 mg

Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer

Nasal decongestant

Uses

temporarily relieves these common cold/flu symptoms:

- nasal congestion
- headache
- minor aches and pains
- stuffy nose
- sinus congestion and pressure

temporarily reduces fever

Warnings

Liver warning: This product contains acetaminophen. 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.
- for more than 10 days unless directed by a doctor

Ask a doctor before use if you have

- liver disease
- thyroid disease
- diabetes
- high blood pressure
- heart disease
- trouble urinating due to an enlarged prostate gland

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.

Stop use and ask a doctor if

- you get nervous, dizzy, or sleepless
- 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 appear

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 use more than directed**

Adults and children: (12 years and over) Take 2 tablets every 6 hours. Do not take more than 8 tablets in 24 hours.

Children under 12 years: Ask a doctor

Other information

- store at room temperature 59°-86°F (15°-30°C)
- tamper evident sealed packets
- do not use any opened or torn packets
- avoid excessive heat and humidity

Inactive ingredients

corn starch, crospovidone, hypromellose, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, stearic acid.

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CROSPVIDONE (UNII: 68401960MK)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

Product Characteristics

Color	white (white)	Score	no score
Shape	ROUND (ROUND)	Size	12mm
Flavor		Imprint Code	AZ;261
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-001-33	50 in 1 BOX	12/30/2008	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-001-48	125 in 1 BOX	12/30/2008	
2	NDC:47682-001-99	2 in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:47682-001-99	2 in 1 PACKET; Type 0: Not a Combination Product	12/30/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/30/2008	

Labeler - Unifirst First Aid Corporation (832947092)