MEDI FIRST SINUS PAIN AND PRESSURE- acetaminophen, phenylephrine hydrochloride tablet
MEDI FIRST PLUS SINUS PAIN AND PRESSURE- acetaminophen, phenylephrine hydrochloride tablet
OTIS CLAPP MYGREX- acetaminophen, phenylephrine hydrochloride tablet
Unifirst First Aid Corporation

819R Sinus Pain and Pressure UniFirst MF/MFP OC

Drug Facts

Active ingredients (in each tablet)

Acetaminophen 500mg

Phenylephrine HCl 5mg

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.

Over dose 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 59º-86ºF (15º-30ºC)

■ 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

Questions or comments? 1-800-634-7680

Medi-First Plus Sinus Pain and Pressure Label

100 Tablets
(505 x 2's)
Medi First ® Plus
Sinus Pain and Pressure
Aches & Fever • Acetaminophen 500mg

Nasal Decongestant • Phenylephrine HCl 5 mg

Pull to Open

This Package is For Households Without Young Children.

Sinus Pain Relief

Tamper Evident

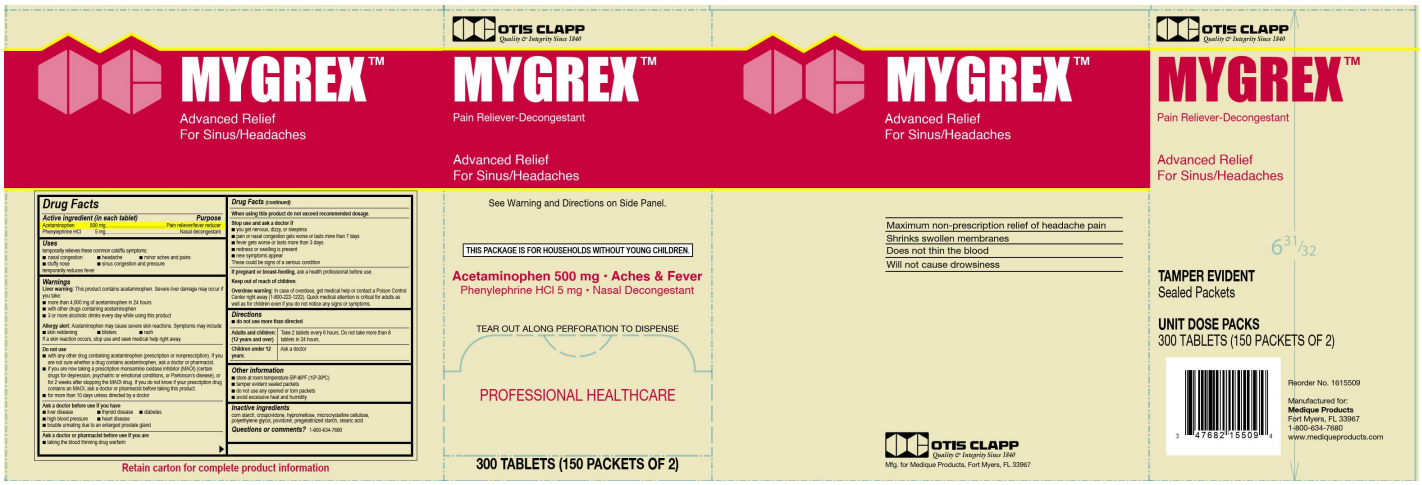
Unit Dose Packet

Otis Clapp Mygrex Label

Otis Clapp
Quality and Integrity Since 1840
MYGREX ™
Pain Reliever-Decongestant
Advanced Relief

For Sinus/Headaches
See Warnings and Directions on Side Panel

This package is for Households without Young Children.
Acetaminophen 500 mg • Aches & Fever
Phenylephrine HCl 5 mg • Nasal Decongestant
Tear Out Along Perforation To Dispense
PROFESSIONAL HEALTHCARE
300 Tablets (150 PACKETS OF 2)



Medi-First Sinus Pain and Pressure Label

Medi-First ®
Sinus Pain and Pressure

Aches & Fever • Acetaminophen 500 mg

Nasal Decongestant • Phenylephrine HCl 5 mg

Pull to Open

This Package is For Households Without Young Children.

Tamper Evident Unit Dose Packets

100 Tablets

(50 x 2)

MEDI FIRST SINUS PAIN AND PRESSURE

acetaminophen, phenylephrine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47682-819
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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
CROSPOVIDONE (UNII: 68401960MK)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

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:47682-819-33	50 in 1 BOX	12/30/2008	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-819-48	125 in 1 BOX	12/30/2008	
2		2 in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:47682-819-13	250 in 1 BOX	12/30/2008	
3	NDC:47682-819-99	2 in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:47682-819-99	2 in 1 PACKET; Type 0: Not a Combination Product	12/30/2008	
5	NDC:47682-819-50	25 in 1 BOX	04/16/2019	

5		2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

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

MEDI FIRST PLUS SINUS PAIN AND PRESSURE

acetaminophen, phenylephrine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47682-919
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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
CROSPOVIDONE (UNII: 68401960MK)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

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:47682-919-33	50 in 1 BOX	12/30/2008	

1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-919-48	125 in 1 BOX	12/30/2008	
2		2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

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

OTIS CLAPP MYGREX

acetaminophen, phenylephrine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47682-509
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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
CROSPOVIDONE (UNII: 68401960MK)	
POVIDONE (UNII: FZ989GH94E)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

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:47682-509-03	150 in 1 BOX	12/30/2008	
1	NDC:47682-509-99	2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-509-99	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)

Revised: 3/2023

Unifirst First Aid Corporation