

REGULAR STRENGTH PAIN RELIEF- acetaminophen tablet
Preferred Pharmaceuticals Inc.

gc 102

Active Ingredients

Acetaminophen 325 mg

Purpose

Pain Reliever/Fever Reducer

Uses

- temporarily relieves minor aches and pains
- temporarily reduces fever

Warnings

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

- more than 12 tablets (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.

Ask a doctor before use if you have liver disease

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

warfarin.

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- new symptom occur
- redness or swelling is present

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. In case of overdose, get medical help or contact a Poison Control Center right away. 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
- adults and children 12 years and over: take 2 tablets every 4-6 hours, as needed; not more than 10 tablets in 24 hours. Do not take for more than 10 days unless directed by a doctor.
- children under 12 years: ask a doctor

Other Information

- TAMPER EVIDENT: Do not use if imprinted seal under cap is missing or broken.
- store at 20°C-25°C (68°F-77°F)

Repackaged by Preferred Pharmaceuticals, Inc.

Inactive Ingredients

povidone, sodium starch glycolate, starch stearic acid.

Questions or comments?

1-800-540-3765

Package Label

Acetaminophen Tablets 325mg

Compare to Tylenol

Each tablet contains Acetaminophen 325 mg
... pain reliever/fever reducer

Pkg Size: Exp Date:
Lot#: Batch#: Ins:
Mfg: Geri-Care; Brooklyn, New York
Prod#:

Warning
Do not use with any other drug containing acetaminophen (prescription or nonprescription). Acetaminophen may cause liver damage. Keep out of reach of children. May cause severe skin reactions. Keep out of reach of children. Tablet is round white and imprinted with GC 101.

Directions English
Use as directed by your doctor
Take _____ Tablet(s) _____ time(s) a day

Instrucciones Español:
Uso según lo dirigido por su doctor
Cada _____ Tableta(s) _____ vece(s) al día

CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed

Acetaminophen Tablets 325mg
Qty: Ins:
Lot#: Bat#:
Prod# (NDC):

Acetaminophen Tablets 325mg
Qty: Ins:
Lot#: Bat#:
Prod# (NDC):

Acetaminophen Tablets 325mg
Qty: Ins:
Insurance NDC:
Lot#: Bat#:

Acetaminophen Tablets 325mg
Qty: Ins:
Lot#: Bat#:
Prod# (NDC):

Log

Chart

Billing

Patient

REGULAR STRENGTH PAIN RELIEF				
acetaminophen tablet				
Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	
Route of Administration		ORAL	NDC:68788-7359(NDC:57896-102)	
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)			ACETAMINOPHEN	325 mg
Inactive Ingredients				
Ingredient Name				Strength
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
STARCH, CORN (UNII: O8232NY3SJ)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
Product Characteristics				
Color		white	Score	no score
Shape		ROUND	Size	10mm
Flavor			Imprint Code	GC101
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-7359-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	08/21/2019	

2	NDC:68788-7359-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	08/21/2019	
3	NDC:68788-7359-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	08/21/2019	
4	NDC:68788-7359-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	08/21/2019	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		343	08/21/2019	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment			
Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-7359)