ACETAMINOPHEN- acetaminophen tablet
Rebel Distributors Corp

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.

Acetaminophen Drug Facts

Active ingredient

Acetaminophen 325mg

Acetaminophen 500mg

Keep Out of Reach of Children

Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose get medical help or contact a Poison Control Center immediately. Quick medical attention is critical for adults as well as children, even if you do not notice symptoms or signs.

Uses

Use for temporary relief of minor aches and pains due to:

° headache ° muscular aches ° backache ° arthritis ° the common cold ° toothache ° menstrual cramps ° temporarily reduces fever

Warnings

Alcohol Warning: If you consume 3 or more alcoholic beverages every day, ask your doctor whether you should take Acetaminophen or other pain relievers/fever reducers. Acetaminophen may cause liver damage.

Do not use with any other product containing Acetaminophen or if you have ever had an allergic reaction to pain relievers/fever reducers.

Stop use and ask a doctor if

° new symptoms occur ° redness or swelling is present ° pain gets worse or lasts more than 10 days ° fever gets worse or lasts more than 3 days.

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

Directions

Do not take more than directed

° **adults and children 12 years and older,** take 2 tablets every 4 to 6 hours as needed. Do not take more than 12 tablets in 24 hours. ° **children under 12,** consult a doctor before use.

Other Information

For your protection, this bottle has an imprinted Safety Seal under cap. Do not use if Safety Seal is broken or missing.

Store at room temperature.

Questions? To report an adverse drug effect, contact 1-800-795-9775.

Inactive ingredients

Crospovidone, Povidone, Pregelatinized Starch, Stearic Acid.

Distributed by:

Plus PharmaCommack

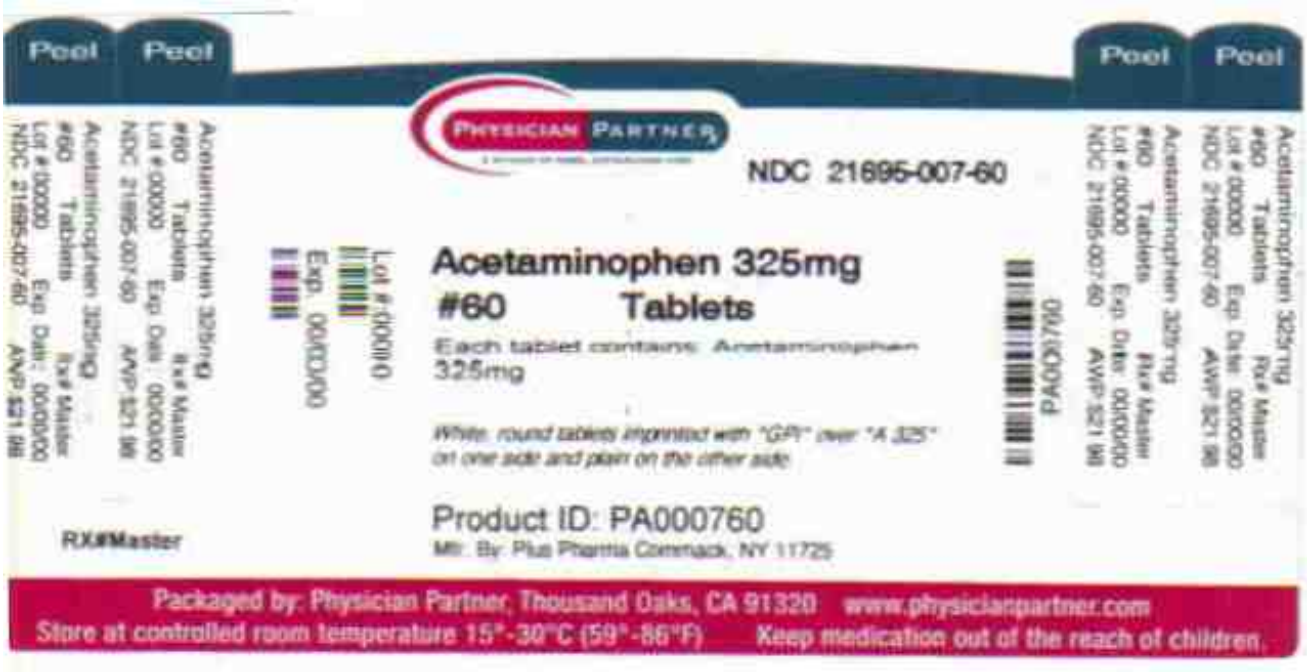
New York 11725

Repackaged by:

Rebel Distributors Corp

Thousand Oaks, CA 91320

Principal Display Panel



Principal Display Panel

ACETAMINOPHEN

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21695-007(NDC:51645-703)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CROSPOVIDONE (UNII: 68401960MK)	
POVIDONE (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	WHITE	**Score**	no score
Shape	ROUND	**Size**	12mm
Flavor		**Imprint Code**	GPI;A325
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:21695-007-30	30 in 1 BOTTLE		
2	NDC:21695-007-60	60 in 1 BOTTLE		
3	NDC:21695-007-00	100 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	01/01/2007	

ACETAMINOPHEN

acetaminophen tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21695-008(NDC:51645-706)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients

Ingredient Name	Strength
CROSPOVIDONE (UNII: 68401960MK)	
POVIDONE (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	WHITE	**Score**	2 pieces
Shape	ROUND	**Size**	14mm
Flavor		**Imprint Code**	GPI;A5
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21695-008-14	14 in 1 BOTTLE		
2	NDC:21695-008-15	15 in 1 BOTTLE		
3	NDC:21695-008-20	20 in 1 BOTTLE		
4	NDC:21695-008-30	30 in 1 BOTTLE		
5	NDC:21695-008-50	50 in 1 BOTTLE		
6	NDC:21695-008-60	60 in 1 BOTTLE		

7	NDC:21695-008-00	100 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	01/01/2007	

Labeler - Rebel Distributors Corp (118802834)

Establishment

Name	Address	ID/FEI	Business Operations
Rebel Distributors Corp		118802834	RELABEL, REPACK

Revised: 2/2011 Rebel Distributors Corp