

ACETAMINOPHEN- acetaminophen tablet

Major Pharmaceuticals

0220K- Major

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325 mg

Purpose

Pain reliever/ Fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- headache
- muscular aches
- backache
- minor pain of arthritis
- the common cold
- toothache
- premenstrual and menstrual cramps
- temporarily reduces fever

Liver warning: This product contains acetaminophen. The maximum daily dose of this product is 10 tablets (3,250 mg) in 24 hours for adults or 5 tablets (1,625 mg) in 24 hours for children. Severe liver damage may occur if

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours, which is the maximum daily amount
- taken with other drugs containing acetaminophen
- adult has 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 allergic to acetaminophen or any of the inactive ingredients in this product

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 in adults
- pain gets worse or lasts more than 5 days in children under 12 years
- fever gets worse or lasts more than 3 days
- new symptoms 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.

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. **This Unit Dose package is not child resistant and is Intended for Institutional Use Only.**

Directions

- **do not take more than directed (see overdose warning)**
- **adults and children 12 years and over:**
 - take 2 tablets every 4 to 6 hours while symptoms last
 - do not take more than 10 tablets in 24 hours, unless directed by a doctor
 - do not use for more than 10 days unless directed by a doctor
- **children 6 years to under 12 years**
 - take 1 tablet every 4 to 6 hours while symptoms last
- do not take more than 5 tablets in 24 hours
- do not use for more than 5 days unless directed by a doctor
- **children under 6 years:** ask a doctor

Other information

Store in a dry place at 15° - 30°C (59° - 86°F).

Povidone, pregelatinized starch, sodium starch glycolate, stearic acid.

Questions or comments?

1-800-231-4670

Keep this carton for complete product information

Tamper Evident: Do not use if sealed blister units are broken or damaged.

Distributed by: MAJOR® PHARMACEUTICALS
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NDC 0904-6773-61

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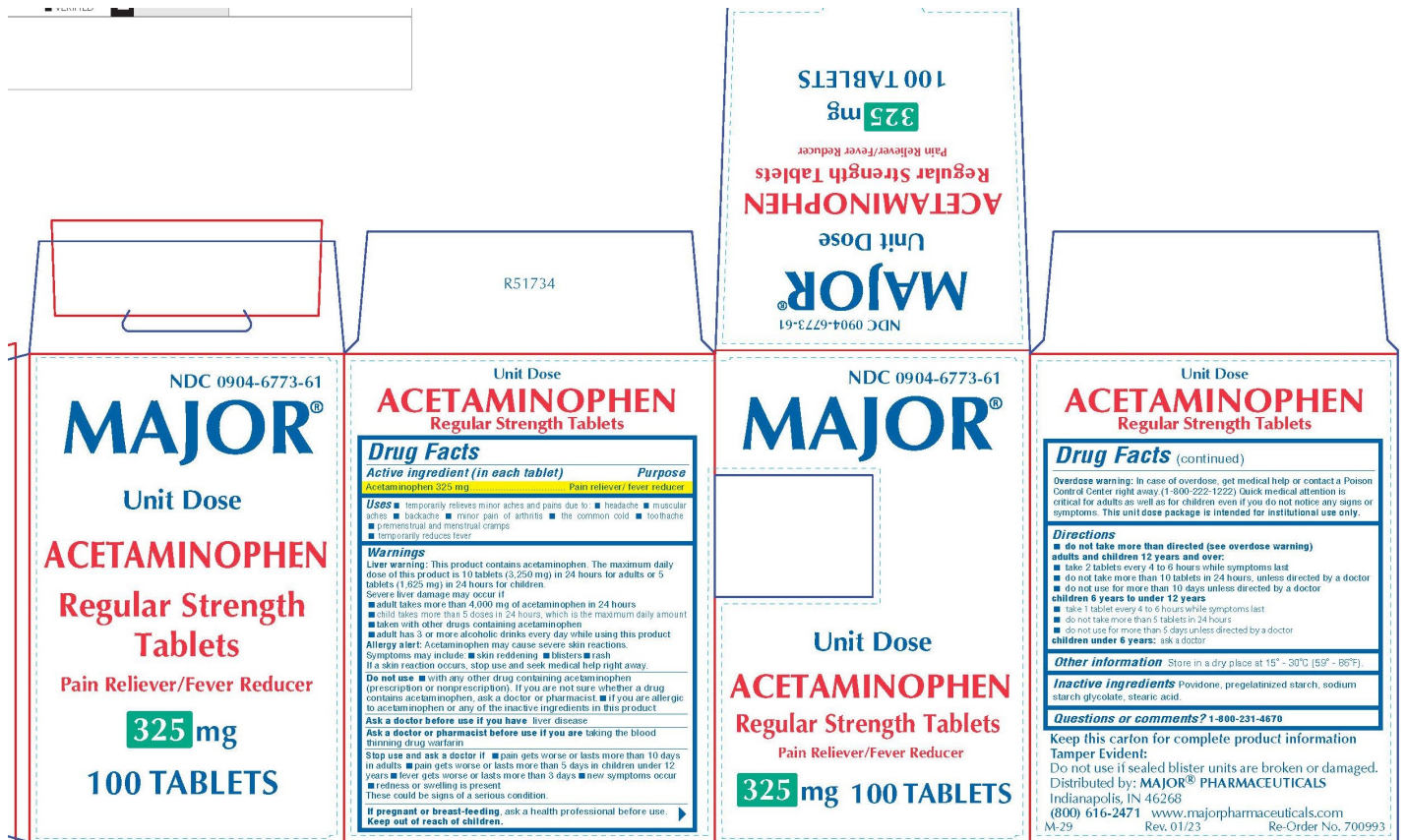
Unit Dose

Acetaminophen

Regular Strength Tablets

325 mg

100 Tablets



ACETAMINOPHEN acetaminophen tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-6773
Route of Administration	ORAL		
Active Ingredient/Active Moiety			

M-29 Rev. 01/23 Re-Order No. 700993

Ingredient Name		Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)		ACETAMINOPHEN	325 mg	
Inactive Ingredients				
Ingredient Name		Strength		
STEARIC ACID (UNII: 4ELV7Z65AP)				
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)				
STARCH, CORN (UNII: O8232NY3SJ)				
POVIDONE K30 (UNII: U725QWY32X)				
Product Characteristics				
Color	white	Score	2 pieces	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	CPC;220	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-6773-61	10 in 1 BOX, UNIT-DOSE	10/23/2018	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M013	10/23/2018		

Labeler - Major Pharmaceuticals (191427277)

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