

**PAIN RELIEF REGULAR STRENGTH- acetaminophen tablet**  
**CHAIN DRUG MARKETING ASSOCIATION INC**

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**1001-QCH-2024-0612**

***Drug Facts***

**Active ingredient (in each tablet)**

Acetaminophen 325 mg

**Purpose**

Pain reliever/fever reducer

**Uses**

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

**Warnings**

**Liver warning**

This product contains acetaminophen. 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 the user is allergic to acetaminophen or any of the inactive ingredients in this product

**Ask a doctor before use if the user has** liver disease.

**Ask a doctor or pharmacist before use if the user is** 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.

**Directions**

- **do not take more than directed (see overdose warning)**

adults and children 12 years and over	<ul style="list-style-type: none"><li>▪ take 2 tablets every 4 to 6 hours while symptoms last</li><li>▪ do not take more than 10 tablets in 24 hours</li><li>▪ do not use for more than 10 days unless directed by a doctor</li></ul>
children 6-11 years	<ul style="list-style-type: none"><li>▪ take 1 tablet every 4 to 6 hours while symptoms last</li><li>▪ do not take more than 5 tablets in 24 hours</li><li>▪ do not use for more than 5 days unless directed by a doctor</li></ul>
children under 6 years	<ul style="list-style-type: none"><li>▪ ask a doctor</li></ul>

**Other information**

- store between 20-25°C (68-77°F)
- retain carton for complete product information

**Inactive ingredients**

povidone, pregelatinized starch, sodium starch glycolate, stearic acid

**PRINCIPAL DISPLAY PANEL**

Quality Choice®

NDC 83324-076-01

†Compare to Active Ingredient in TYLENOL® Regular Strength Tablets

Regular Strength

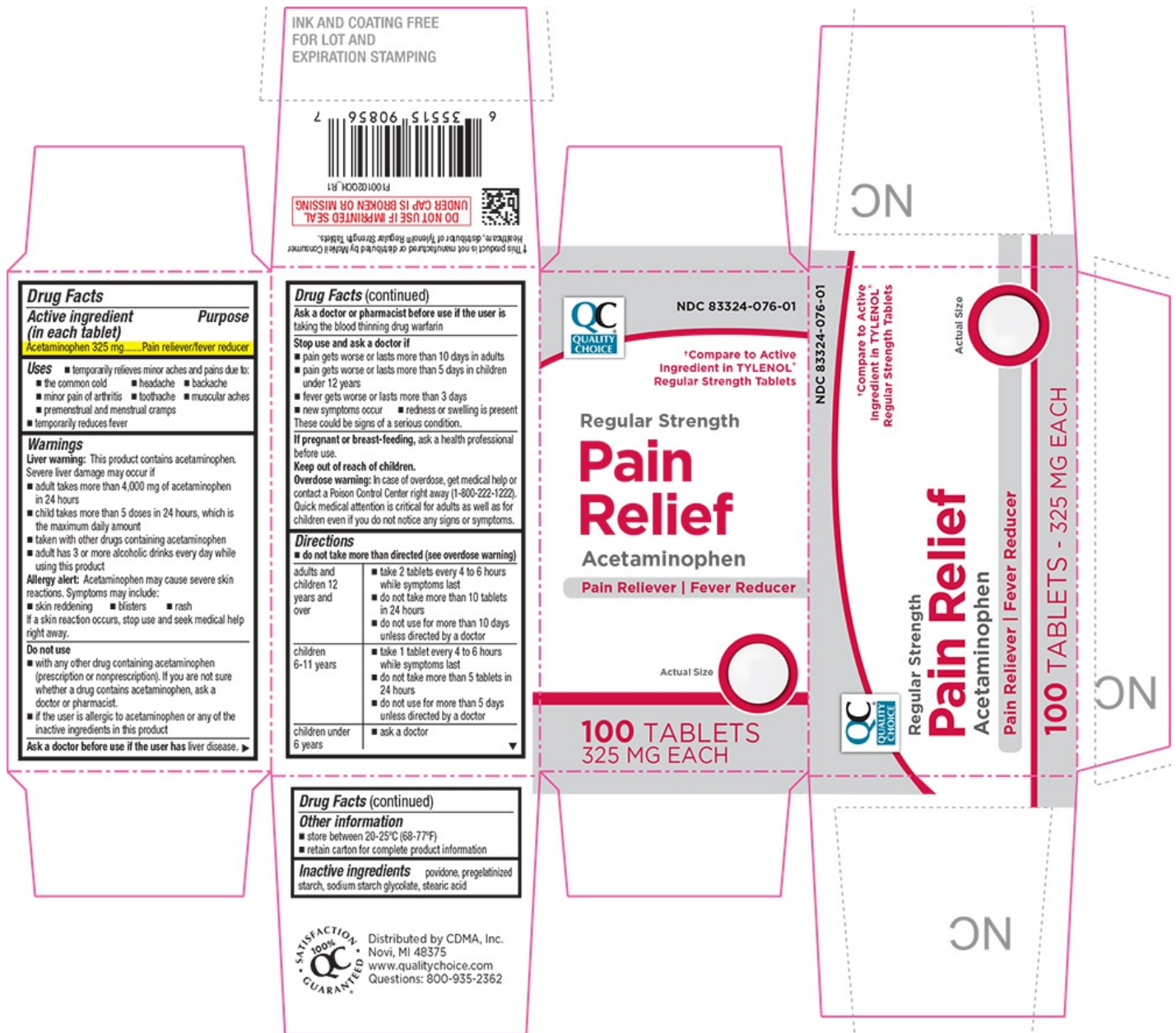
Pain Relief

Acetaminophen

Pain Reliever | Fever Reducer

Actual Size

100 TABLETS - 325 MG EACH



## PAIN RELIEF REGULAR STRENGTH

acetaminophen tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83324-076
<b>Route of Administration</b>	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
<b>POVIDONE</b> (UNII: FZ989GH94E)	
<b>STARCH, PREGELATINIZED CORN</b> (UNII: O8232NY3SJ)	
<b>SODIUM STARCH GLYCOLATE TYPE A CORN</b> (UNII: AG9B65PV6B)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND	<b>Size</b>	10mm
<b>Flavor</b>		<b>Imprint Code</b>	M2A3;57344
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-076-01	1 in 1 CARTON	04/05/2024	
1		100 in 1 BOTTLE, PLASTIC; 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	04/05/2024	

**Labeler** - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

Revised: 6/2024

CHAIN DRUG MARKETING ASSOCIATION INC