# **EQUATE PAIN RELIEVER- acetaminophen tablet WALMART INC.**

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#### Wal-Mart Pain Reliever 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

### **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
- 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 has ever had an allergic reaction to this product or any of its ingredients

#### Ask a doctor before use if the user

- has liver disease
- is a child with pain of arthritis

#### 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 (for adults) or 5 days (for children)
- 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> <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> <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	ask a doctor

## **Inactive ingredients**

croscarmellose sodium\*, povidone, pregelatinized starch, stearic acid \*may contain this ingredient

#### **Questions or comments?**

1-888-287-1915

## **Principal Display Panel**

equate™

Compare to Regular Strength Tylenol® Tablets active ingredient

Regular Strength

Pain Reliever

Acetaminophen 325 mg

Pain Reliever/Fever Reducer

**Actual Size** 

325 mg

100 TABLETS



## **EQUATE PAIN RELIEVER**

acetaminophen tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-299
Route of Administration	ORAL		
Active Ingredient/Active Moiety			

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg	

Inactive Ingredients		
Ingredient Name	Strength	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)		
STEARIC ACID (UNII: 4ELV7Z65AP)		

Product Characteristics			
Color	WHITE	Score	no score
Shape	ROUND (beveled edge)	Size	10mm
Flavor		Imprint Code	325MG;L403
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:79903-299- 10	1 in 1 CARTON	11/21/2024		
1		100 in 1 BOTTLE; 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	11/21/2024	

## Labeler - WALMART INC. (051957769)

Revised: 11/2024 WALMART INC.