EQUATE PAIN RELIEVER ACETAMINOPHEN FOR ADULTS 250 MGacetaminophen bar, chewable Walmart Inc

Equate Pain Reliever Acetaminophen for Adults 250 mg

Active ingredient (in each chewable bar)

Acetaminophen 250mg

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 you take

- more than 4,000mg of acetaminophen in 24 hours, which is the maximum daily amount
- 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.

• 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
- a sodium-restricted diet

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 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 give more than directed (see overdose warning)

Age (yrs.) Adults and children 12	 Dose (chewable bar) take 4 chewable bars every 6 hours while symptoms last do not take more than 12 chewable bars in 24 hours, unless directed by a doctor
years and over	 do not use for more than 10 days unless directed by a doctor
Children under 12 years	ask a doctor

Other information

- each chewable bar contains: potassium 5mg, sodium 16mg.
- Store in a cool dry place between 20-25°C (68-77°F).
- **Child Resistant Container;**do not use if printed seal under cap is broken or missing.

Inactive ingredients:

ammonium glycyrrhizate, FD & C yellow #6, flavors, geleol mono and diglycerides, glucose syrup, hydroxypropyl betadex, maltitol solution, maltodextrin, menthol, neotame, polyethylene glycol 400, povidone, propylene glycol, purified water, seaweed extract (carrageenan), sodium chloride, starch, sucralose, sucrose, trisodium citrate dihydrate.

Questions or comments?

Call **1-888-287-1915**

U.S. Patent 11,273,123

Principal Display Panel - Equate Pain Reliever Acetaminophen for Adults 250 mg-Orange Vanilla Flavor

NDC 79903-247-40

equate[™]

Compare to Tylenol ® active ingredient**

Pain Reliever Acetaminophen for Adults 250 mg

Pain Reliever/Fever Reducer

Contains No Aspirin

Soft Chew Bars

Orange Vanilla Flavor

250 mg EACH 40 CHEWABLE BARS

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.



If you are all ergic to acetaminophen or any of the inactive ingredents Overdose wa ming: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222), Quick medical ■ pain gets worse or lasts more than 10 days ■ fever gets worse or lasts more than 3 days ■ new symptoms occur ■ redness or swelling is present attention is critical for adults as well as for children even if you do not Ask a doctor or pharmacist before use if you are taking the blood If pregnant or breast-feeding, ask a health professional before use with any other drug containing acetaminophen (prescription or than directed (see overdose warning) nonprescription). If you are not sure whether a drug contains Ask a doctor before use if you have I liver disease acetaminophen, ask a doctor or pharmacist. These could be signs of a serious condition. Drug Facts (continued) Keep out of reach of children Stop use and ask a doctor if notice any signs or symptoms. a sodium-restricted diet Adults and children 12 thinning drug warfarin do not give more Do not use **Directions** years and over in this product. Age (yrs.)

■ do not take more than 12 chewable bars in 24 hours, unless directed by a doctor ■ do not use for more than 10 days unless directed by a doctor #6, flavors, geleof mono and diglycerides, glucose syrup, hydroxypropyl betadex, mattitol solution, maltodextrin, menthol, neotame, polyethylene Inactive ingredients: ammonium glycyrthizate, FD&C yellow glycol 400, povidone, propylene glycol, purified water, seaweed extract Child Resistant Container, do not use if printed seal under cap is take 4 chewable bars every 6 hours while carrageenan), sodium chloride, starch, sucralose, sucrose, trisodium each chewable bar contains; potassium 5mg, sodium 16mg. store in a cool dry place between 20-25°C (68-77°F). Dose (chewable bar) symptoms last ask a doctor Other information Children under 12 years broken or missing. citrate dihydrate

Questions or comments? U.S. Patent 11,273,123 3111-888-287-1915

EQUATE PAIN RELIEVER ACETAMINOPHEN FOR ADULTS 250 MG

acetaminophen bar, chewable

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-247
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	250 mg	

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POVIDONE K30 (UNII: U725QWY32X)	
CORN SYRUP (UNII: 9G5L16BK6N)	
HYDROXYPROPYL BETADEX (UNII: 11960HX6EK)	
NEOTAME (UNII: VJ597D52EX)	
AMMONIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)	
GLYCERYL MONO AND DIPALMITOSTEARATE (UNII: KC98RO82HJ)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CARRAGEENAN (UNII: 5C69YCD2YJ)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SUCROSE (UNII: C151H8M554)	
MALTITOL (UNII: D65DG142WK)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
WATER (UNII: 059QF0KO0R)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics			
Color	orange	Score	no score
Shape	RECTANGLE	Size	25mm
Flavor	ORANGE (Orange Vanilla Flavor)	Imprint Code	0
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-247- 40	40 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2024	

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M013	06/01/2024	

Labeler - Walmart Inc (051957769)

Establishment			
Name	Address	ID/FEI	Business Operations
USpharma Ltd		080664601	manufacture(79903-247)

Revised: 3/2024 Walmart Inc