

ARTHRITIS PAIN RELIEVER- acetaminophen tablet, film coated, extended release

Walgreen Company

Walgreen Co. Arthritis Pain Reliever Drug Facts

Active ingredient (in each caplet)

Acetaminophen 650 mg

Purpose

Pain reliever/fever reducer

Uses

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

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 6 caplets 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

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 take more than directed (see overdose warning)**

adults	<ul style="list-style-type: none">• take 2 caplets every 8 hours with water• swallow whole; do not crush, chew, split or dissolve• do not take more than 6 caplets in 24 hours• do not use for more than 10 days unless directed by a doctor
under 18 years of age	<ul style="list-style-type: none">• ask a doctor

Other information

- store at 20-25°C (68-77°F)
- meets the requirements of USP *Dissolution Test 4*

Inactive ingredients

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polysorbate 80, povidone, pregelatinized starch, stearic acid, titanium dioxide

Questions or comments?

1-800-719-9260

Principal Display Panel

Walgreens

Compare to the active ingredient in Tylenol® 8HR Arthritis Pain

Arthritis Pain Reliever

ACETAMINOPHEN EXTENDED-RELEASE TABLETS, 650 mg / PAIN RELIEVER / FEVER REDUCER

Arthritis

8 Hour

- For the temporary relief of minor arthritis pain

100 CAPLETS*

650 mg EACH

(*capsule-shaped tablets)

ACTUAL SIZE

3



ITEM 915876 W10107-0624-F

U.S. Patent 7,897,172
Contains No Aspirin**DO NOT USE IF PRINTED FOIL
UNDER CAP IS BROKEN OR MISSING**Gluten Free
DISTRIBUTED BY:
WALGREEN CO.
DEERFIELD, IL 60015
**100% SATISFACTION
GUARANTEED**
walgreens.com
©2021 Walgreen Co.
NDC 0363-0544-78Walgreens Pharmacist Recommended.
Our pharmacists recommend the Walgreens brand. We invite
you to compare to national brands.
†This product is not manufactured or distributed by Johnson &
Johnson Consumer Inc., McNeil Consumer Healthcare Division,
distributor of Tylenol® 8HR Arthritis Pain.

EXP. LOT

: 54478 94 F9

how2recycle.com
REV-0324**Walgreens**Compare to the active ingredient
in Tylenol® 8HR Arthritis Pain†**Arthritis
Pain Reliever**ACETAMINOPHEN EXTENDED-RELEASE
TABLETS, 650 mg / PAIN RELIEVER / FEVER REDUCER

Arthritis 8 Hour

• For the temporary relief of
minor arthritis pain**100** CAPLETS*
650 mg EACH

(*capsule-shaped tablets)



ACTUAL SIZE

Drug Facts

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PEEL BACK HERE ▶**ADHESIVE
AREA
NO COPY****Drug Facts (continued)****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
 - 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.**Drug Facts (continued)****Directions****do not take more than directed (see overdose warning)**

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acetaminophen tablet, film coated, extended release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0544
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg

Inactive Ingredients

Ingredient Name	Strength
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	WHITE	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	L544
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0544-71	1 in 1 CARTON	02/11/2011	01/31/2022
1		50 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0363-0544-47	1 in 1 CARTON	02/11/2011	04/23/2013
2		150 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:0363-0544-85	1 in 1 CARTON	02/04/2011	03/23/2017
3		250 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:0363-0544-78	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/31/2014	
5	NDC:0363-0544-62	1 in 1 CARTON	08/15/2014	11/30/2024
5		24 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075077	05/26/2000	

