PAIN RELIEF- acetaminophen tablet, coated DISCOUNT DRUG MART

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

1004-DDM-2021-1124

Drug Facts

Active ingredient (in each caplet)

Acetaminophen 500 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 you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

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 and children 12 years and over	 take 2 caplets every 6 hours while symptoms last do not take more than 6 caplets in 24 hours, unless directed by a doctor do not use for more than 10 days unless directed by a doctor
children under 12 years	ask a doctor

Other information

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

Inactive ingredients

hypromellose, mineral oil, polyethylene glycol, polysorbate 80, povidone, pregelatinized starch, sodium starch glycolate, stearic acid, titanium dioxide

PRINCIPAL DISPLAY PANEL

DISCOUNT drug mart FOOD FAIR

EXTRA STRENGTH

PAIN RELIEF

NON-ASPIRIN

ACETAMINOPHEN

Pain Reliever / Fever Reducer

†Compare to the active ingredient in EXTRA STRENGTH TYLENOL®

50 CAPLETS

500 mg Each



PAIN RELIEF

acetaminophen tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53943-304
Route of Administration	ORAL		

Active Ingredient/Active Moiety

l	Ingredient Name	Basis of Strength	Strength
	ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	500 mg

Inactive Ingredients				
Ingredient Name	Strength			
HYPROMELLOSES (UNII: 3NXW29V3WO)				
MINERAL OIL (UNII: T5L8T28FGP)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POLYSORBATE 80 (UNII: 6OZP39ZG8H)				
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)				
STARCH, CORN (UNII: O8232NY3SJ)				
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				

Product Characteristics				
Color	white	Score	no score	
Shape	OVAL	Size	17mm	
Flavor		Imprint Code	M2A4;57344	
Contains				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:53943- 304-02	1 in 1 CARTON	01/05/2010	02/28/2025	
1		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
2	NDC:53943- 304-04	1 in 1 CARTON	03/04/2010	02/28/2025	
2		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			
3	NDC:53943- 304-06	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/05/2010		
4	NDC:53943- 304-09	2 in 1 CARTON	04/04/2013	02/28/2025	
4		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product			

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC monograph not final	part343	01/05/2010	

Labeler - DISCOUNT DRUG MART (047741335)

Revised: 11/2022 DISCOUNT DRUG MART