UP AND UP ACETAMINOPHEN- acetaminophen tablet, film coated, extended release Target Corporation

Target Corporation Acetaminophen 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	 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	• ask a doctor

Other information

• store at 20-25°C (68-77°F). Avoid excessive heat 40°C (104°F).

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

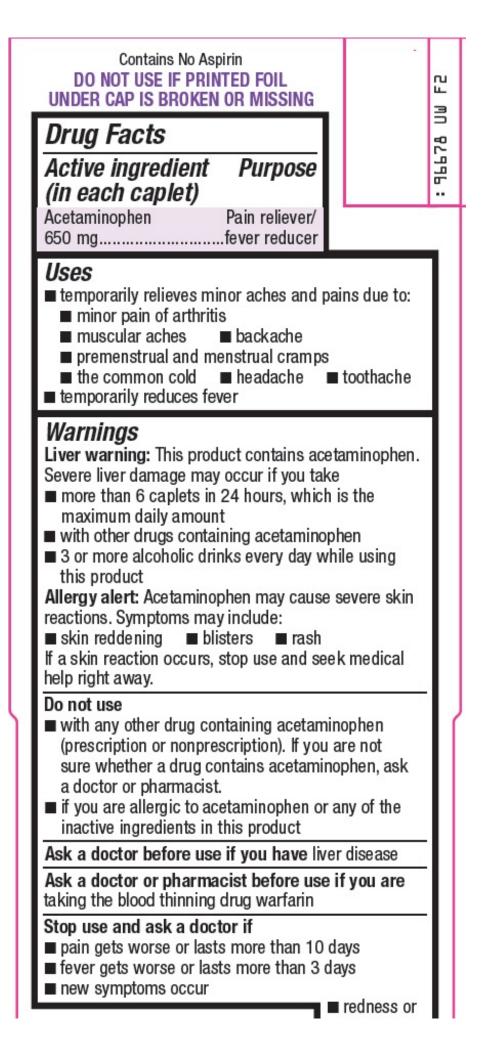
Questions? Call

1-888-547-7400

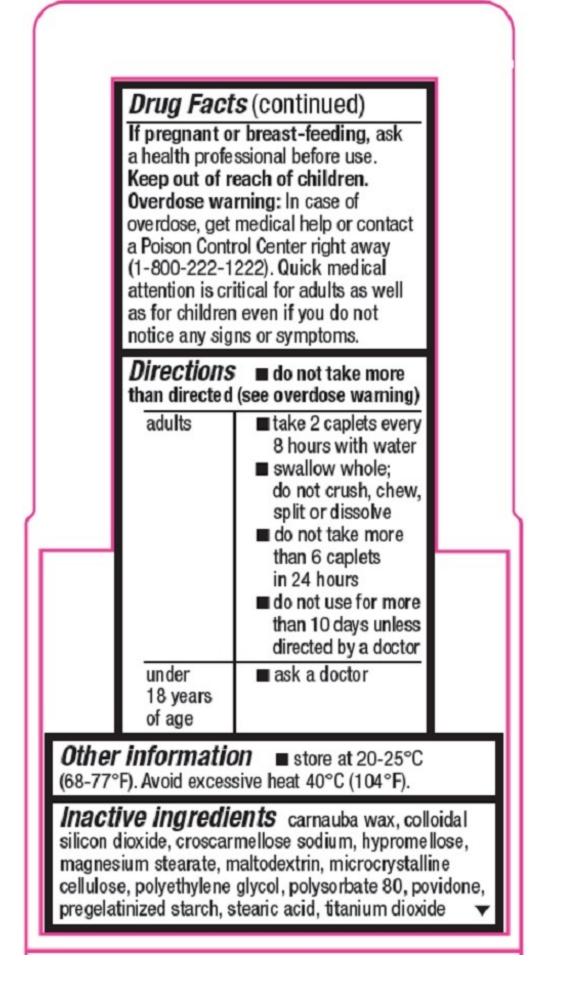
Principal Display Panel

see new warning Compare to active ingredient in Tylenol® 8HR Arthritis Pain arthritis pain acetaminophen extended-release tablets, 650 mg pain reliever / fever reducer for the temporary relief of minor arthritis pain DO NOT USE WITH OTHER MEDICINES CONTAINING ACETAMINOPHEN THIS PACKAGE FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN 100 CAPLETS* 100 CAPLETS* 650 mg EACH *(CAPSULE-SHAPED TABLETS)









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Product Informat	ion							
Product Type		HUMAN OTC DRUG Item Code (Source			ource)	NDC:11673-966		
Route of Administra	tion	ORAL						
Active Ingredient	/Active Mo	iety						
Ingredient Name Basis						f Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) ACETAMINOPH							650 mg	
Inactive Ingredie	nts							
Ingredient Name							Strength	
CARNAUBA WAX (UN								
SILICON DIO XIDE (U								
CROSCARMELLOSE								
HYPROMELLOSES (U								
MAGNESIUM STEARA								
MALTO DEXTRIN (UN								
MICRO CRYSTALLIN								
POLYETHYLENE GL								
POLYSORBATE 80 (U		э8Н)						
POVIDONE (UNII: FZ9								
STEARIC ACID (UNII: TITANIUM DIO XIDE (ID)						
TTANIUM DIO AIDE (UNII. 13F1X3 V2	Jr)						
Product Characte	ristics							
Color WHIT			Score	re		no score		
Shape	hape CAPSULE		Size	Size			19 mm	
Flavor			Imprint Co	de		L544		
Contains								
Packaging								
# Item Code		Package Description			eting Start Dat	e Marketii	ng End Dat	
	100 in 1 BOTT	LE; Type 0: Not a Con	nbination Proc	luct 03/15/2	20 18			
1 NDC:11673-966-78								
Marketing Info	ormation							
		ion Number or Mono	ograph Cita	tion Marl	ceting Start Date	e Marketii	ıg End Date	

Revised: 12/2019

Target Corporation