PHARBETOL- acetaminophen 325 mg tablet Ulai Health LLC

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.

Drug Facts

Active ingredient (in each tablet)

Acetaminophen 325 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. The maximum daily dose of this product is 10 tablets (3,250 mg) in 24 hours for adults or 5 tablets (1,625 mg) in 24 hours for children. 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, which is the maximum daily amount
- 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 a 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 in adults
- pain gets worse or lasts more than 5 days in children under 12 years
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur

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).

Directions

• do not take more than directed (see overdose warning)

adults and children 12 years and over	 take 2 tablets every 4 to 6 hours while symptoms last do not take more than 10 tablets in 24 hours, unless directed by a doctor do not use for more than 10 days unless directed by a doctor
children 6 to under 12 years	 take 1 tablet every 4 to 6 hours while symptoms last do not take more than 5 tablets in 24 hours do not use for more than 5 days unless directed by a doctor
children under 6 years	ask a doctor

Other information

• store at 20-25 ⁰C (68-77 ⁰F)

Inactive ingredients

povidone, pregelatinized corn starch, sodium starch glycolate, stearic acid

Questions or comments?

(866) 562-2756 (Mon - Fri 8 AM to 4 PM EST)

PHARBEST

NDC 73057-353-11

Manufactured in the USA

Regular Strength *COMPARE TO the active ingredient in TYLENOL $^{\ensuremath{\mathbb{R}}}$ Regular Strength

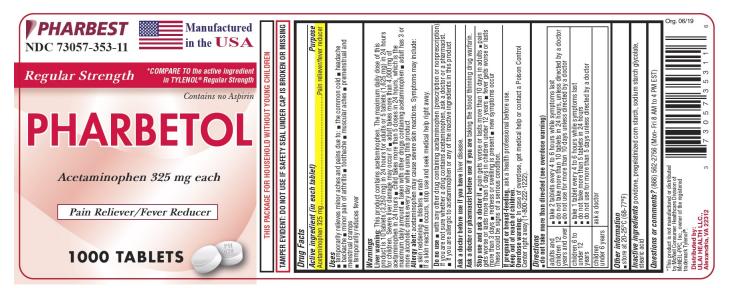
Contains no Aspirin

PHARBETOL

Acetaminophen 325 mg each

Pain Reliever/Fever Reducer

1000 TABLETS



PHARBETOL acetaminophen 325 mg tablet											
Product Information											
Product Type	HUMAN OTC DRUG	ltem Code (Source)	NDC:73057-353								
Route of Administration	ORAL										

A	ctive Ingred	lient/Act	ive Moietv	,						
-							Basis of S	Strength	Strength	
AC			-		HEN - UNII:36209IT	-L9D)	ACETAMINOPHEN		325 mg	
		- (01							010g	
In	active Ingr	edients								
			Ingr	edient	Name			5	Strength	
PC	VIDONE, UNSF	PECIFIED (U	INII: FZ989GH9	94E)						
ST	ARCH, CORN (JNII: 082321	NY3SJ)							
SO	DIUM STARCH	GLYCOLAT	Е ТҮРЕ А СО	RN (UNII	: AG9B65PV6B)					
ST	EARIC ACID (UI	NII: 4ELV7Z6	55AP)							
D -	roduct Char									
		racteristi								
Color white				Score			no score			
			ROUND		Size			10mm		
Flavor				Ir	Imprint Code			PH020		
Co	ontains									
Pa	ackaging									
						Mark	eting Start	Marke	eting End	
#	Item Code		Package I	Descrip	otion		Date		Date	
1	NDC:73057- 353-11	1000 in 1 BOTTLE, PLASTIC; Type Combination Product			0: Not a	07/08/2019				
2	NDC:73057-	100 in 1 BOTTLE, PLASTIC; Type): Not a	07/08/2	n19			
-	353-08	Combination Product				07,00,2	515			
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Labeler - Ulai Health LLC (081181535)

Revised: 1/2022

Ulai Health LLC