ACETAMINOPHEN, IBUPROFEN- acetaminophen, ibuprofen tablet, film coated GLENMARK THERAPEUTICS INC., USA

Drug Facts

Active ingredients (in each caplet)

Acetaminophen 250 mg

- Ibuprofen 125 mg (NSAID*)
- *nonsteroidal anti-inflammatory drug

Purposes

Pain reliever

Pain reliever

Uses

- temporarily relieves minor aches and pains due to:
 - o headache
 - o toothache
 - o backache
 - o menstrual cramps
 - o muscular aches
 - o minor pain of arthritis

Warnings

Acetaminophen liver damage warning:

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

- with other drugs containing acetaminophen
- more than 6 caplets in 24 hours, which is the maximum daily amount for this product
- 3 or more alcoholic drinks every day while using this product

Acetaminophen allergy alert: may cause severe skin reactions. Symptoms may include:

skin reddening
 blisters
 rash

If skin reaction occurs, stop use and seek medical help right away.

NSAID allergy alert: ibuprofen may cause a severe allergic reaction, especially in

people allergic to aspirin. Symptoms may include:

hives

• facial swelling

asthma (wheezing)

- shock
- skin reddening
 rash
- blisters

If an allergic reaction occurs, stop use and seek medical help right away.

NSAID stomach bleeding warning:

This product contains an NSAID, which may cause severe stomach bleeding. The chance is higher if you:

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs [aspirin, ibuprofen, naproxen, or others]
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

Heart attack and stroke warning:

NSAIDs, except aspirin, increase the risk of heart attack, heart failure, and stroke. These can be fatal. The risk is higher if you use more than directed or for longer than directed.

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 have ever had an allergic reaction to acetaminophen or any other pain reliever
- right before or after heart surgery

Ask a doctor before use if

- you have liver disease
- stomach bleeding warning applies to you
- you have problems or serious side effects from taking pain relievers
- you have a history of stomach problems, such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma, or had a stroke

• you are taking a diuretic

Ask a doctor or pharmacist before use if you are

- under a doctor's care for any serious condition
- taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
- taking any other drug

When using this product

• take with food or milk if stomach upset occurs

Stop use and ask a doctor if

- you experience any of the following signs of stomach bleeding:
 - o feel faint
 - o vomit blood
 - o have bloody or black stools
 - o have stomach pain that does not get better
- you have symptoms of heart problems or stroke:
 - o chest pain
 - o trouble breathing
 - o weakness in one part or side of body
 - o slurred speech
 - o leg swelling
- pain gets worse or lasts more than 10 days
- redness or swelling is present in the painful area
- any new symptoms appear

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen at 20 weeks or later in pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away. Prompt 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

adults and children 12 years and	 take 2 caplets every 8 hours
over	while symptoms persist
children under 12 years	

• do not take more than 6 caplets in 24 hours, unless directed by a doctor

Other information

- read all warnings and directions before use. Keep carton.
- store at 20°C to 25°C (68°F to 77°F)
- avoid excessive heat above 40°C (104°F)

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, ferrosoferric oxides, glyceryl dibehenate, hypromellose, iron oxide yellow, iron oxide red, polydextrose, povidone, polyethylene glycol, propylene glycol, shellac, pregelatinized starch and titanium dioxide

Questions or comments?

call weekdays 9 AM to 5 PM at 1 (888) 721-7115

Distributed by:

Glenmark Therapeutics Inc., USA

Mahwah, NJ 07430

March 2023

PRINCIPAL DISPLAY PANEL

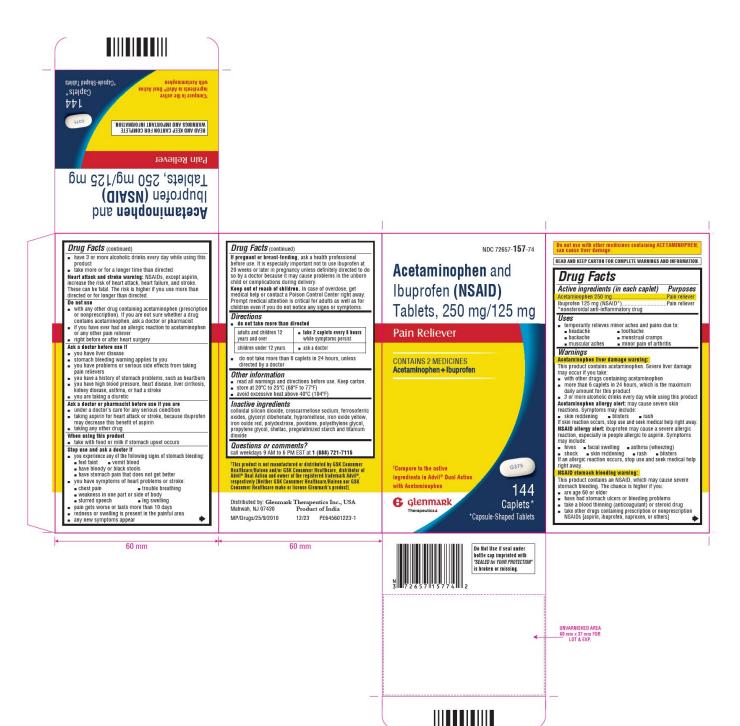
NDC 72657-157-74

Acetaminophen 250 mg and Ibuprofen (NSAID) 125 mg Tablets

Pain Reliever

144'sCaplets*

*Capsule-Shaped Tablets



ACETAMINOPHEN, IE					
acetaminophen, ibuprofen tal	piet, film coated				
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:72657-15		57-157	
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingre	edient Name		Basis of St	trength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (I	QM)	IBUPROFEN		125 mg	

250 mg

Inactive Ingredients				
Ingredient Name	Strength			
STARCH, CORN (UNII: 08232NY3SJ)				
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)				
POVIDONE K90 (UNII: RDH86HJV5Z)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
GLYCERYL DIBEHENATE (UNII: R8WTH25YS2)				
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ 8WG20P6)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
POLYDEXTROSE (UNII: VH2XOU12IE)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
FERRIC OXIDE RED (UNII: 1K09F3G675)				
FERRIC OXIDE YELLOW (UNII: EX43802MRT)				
SHELLAC (UNII: 46N107B710)				
FERROSOFERRIC OXIDE (UNII: XM0M87F357)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				

Product Characteristics

Color	YELLOW (light yellow to yellow)	Score	no score
Shape	CAPSULE	Size	15mm
Flavor		Imprint Code	G375
Contains			

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72657-157- 18	1 in 1 CARTON	05/07/2024	
1		18 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:72657-157- 36	1 in 1 CARTON	05/07/2024	
2		36 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:72657-157- 72	1 in 1 CARTON	05/07/2024	
3		72 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:72657-157- 74	1 in 1 CARTON	05/07/2024	
4		144 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:72657-157- 76	1 in 1 CARTON	05/07/2024	
5		216 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:72657-157- 20	1 in 1 CARTON	05/07/2024	

6		250 in 1 BOTTLE; Type 0: Not a Combination Product			
7	NDC:72657-157- 05	1 in 1 CARTON	05/07/2024		
7		500 in 1 BOTTLE; Type 0: Not a Combination Product			
Μ	Marketing Information				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
AN	IDA	ANDA218311	05/07/2024		

Labeler - GLENMARK THERAPEUTICS INC., USA (969085666)

Establishment			
Name	Address	ID/FEI	Business Operations
Glenmark Pharmaceuticals Limited		862603186	ANALYSIS(72657-157), MANUFACTURE(72657-157)

Revised: 5/2024

GLENMARK THERAPEUTICS INC., USA