IBUPROFEN 100- ibuprofen tablet, coated IBUPROFEN 250- ibuprofen tablet, coated JHK Inc dba American Safety & First Aid

IBUPROFEN

Drug Facts

Active ingredient (In Each Tablets)

Ibuprofen, USP 200 mg (NSAID) *Nonsteroidal Anti-Inflammatory Drug

Purpose

pain reliever/fever reducer

Use

temporarily relieves fever and minor aches and pains associated with:

- common cold
- backache
- headache
- toothache
- arthritis
- muscular aches
- menstrual cramps

Warnings

Allergy Alert

Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin. Symptoms may include:

- hives
- asthma (wheezing)
- facial swelling
- shock
- skin reddening
- rash
- blisters
- If an allergic reaction occurs, stop use and seek medical help right away.

Stomach bleeding warning

This product contains a nonsteroidal anti-inflammatory drug (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

Do not use

- if you ever had an allergic reaction to any other pain reliever/fever reducer
- right before or after heart surgery

Ask a doctor before use if

- stomach bleeding warning applies to you
- you have a history of stomach problems, such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, or kidney disease
- you are taking a diuretic

Ask a doctor or pharmacist before use if you are

- taking any other drug containing an NSAID (prescription or nonprescription)
- taking a blood thinning (anticoagulant) drug
- under a doctor's care for any serious condition
- taking any other drug

When using this product

- take with food or milk if stomach upset occurs
- long term continuous use may increase the risk of heart attack or stroke

Stop use and ask a doctor if you experience any of the following signs of stomach bleeding:

- feel faint
- vomit blood
- have bloody or black stools
- have stomach pain that does not get better
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 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 DURING THE LAST 3 MONTHS OF 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.

Directions

- do not take more than directed
- the smallest effective dose should be used
- do not take longer than 10 days, unless directed by a doctor (see Warnings)

Adults: 1 tablet every 4 to 6 hours while symptoms occur; if pain or fever does not respond to 1 tablet, 2 tablets may be used, but do not exceed 6 tablets in 24 hours unless directed by a doctor.

Children under 12 years: Ask a doctor.

Other Information

- do not use if packet is opened or torn
- Store at 20-25°C (68-77°F) avoid excessive heat above 40°C (104°F)

Inactive Ingredients

crosscarmellose sodium¹, FD& C Blue #2¹, FD& C red #40¹, FD& C Yellow #6¹, hypromellose¹, iron oxide red¹, lactose monohydrate¹, magnesium stearate¹, maltodextrin¹, medium chain triglycerides¹, microcrystalline cellulose¹, polydextrose¹, polyethylene glycol¹, polyvinyl alcohol¹, povidone (K-30)¹, silicon dioxide, sodium starch glycolate¹, starch, stearic acid¹, talc¹, titanium dioxide, triacetin¹,

1 contains one or more of these ingredients

Questions?

1-866-651-3660 Monday-Friday

8:00am-5:00pm EST

PRINCIPAL DISPLAY PANEL - 100 Tablet Packet Box

This product is not manufactured or distributed by Pfizer Consumer Healthcare Owner of the registered trademark Advil[®].

Product #1102

Ibuprofen

Pain Reliever/Fever Reducer

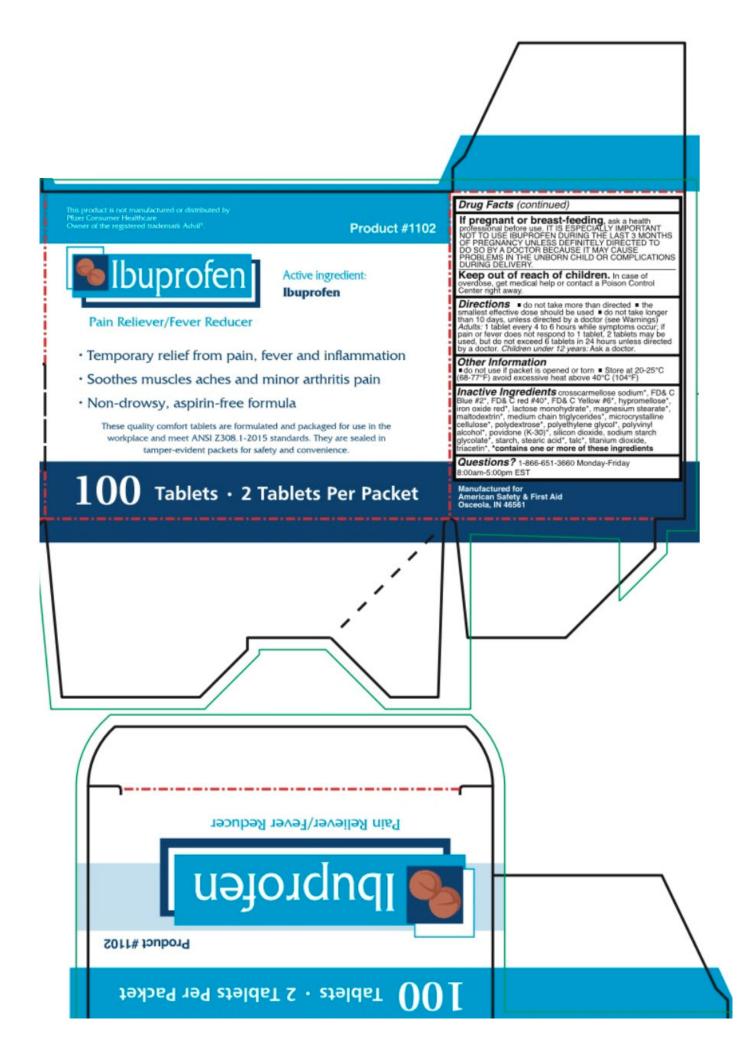
Active ingredient: Ibuprofen

- Temporary relief from pain, fever and inflammation
- Soothes muscles aches and minor arthritis pain
- Non-drowsy, aspirin-free formula

These quality comfort tablets are formulated and packaged for use in the

workplace and meet ANSI Z308.1-2015 standards. They are sealed in tamper-evident packets for safety and convenience.

100 Tablets • 2 Tablets Per Packet



Drug Facts		
	let)pain reliever/fever reducer	
Use temporarily relieves fever and m headache toothache arthritis	ninor aches and pains associated with: common cold backache muscular aches menstrual cramps	Ibuprofen
Symptoms may include: <pre>whites # asthr •blisters • If an allergic reaction occur Stomach bleeding warning: This pro- may cause severe stomach bleeding, i ulcers or bleeding problems • take a b containing prescription or nonprescript</pre>	severe allergic reaction, especially in people allergic to aspirin. na (wheezing) ●facial swelling ●shock ●skin reddening ●rash s, stop use and seek medical help right away. duct contains a nonsteroidal anti-inflammatory drug (NSAID), which he chance is higher if you: ■ are age 60 or older ●have had stomach lood thinning (anticoagulant) or steroid drug ●take other drugs ion NSAIDs (aspirin, ibuprofen, naproxen, or others) ● have 3 or more this product ● take more or for a longer time than directed	Pain Reliever/Fever Reduce
Do not use • if you ever had an a or after heart surgery	llergic reaction to any other pain reliever/fever reducer • right before	\vdash \bigcirc
Ask a doctor before use if stomach problems, such as heartburn kidney disease vou are taking a diure	stomach bleeding warning applies to you syou have a history of you have high blood pressure, heart disease, liver cirrhosis, or tic	PULL TO OPEN
Ask a doctor or pharmacist NSAID (prescription or nonprescription care for any serious condition = taking	t before use if you are • taking any other drug containing an) • taking a blood thinning (anticoagulant) drug • under a doctor's any other drug	Compare Active Ingredients To: Adv
When using this product	take with food or milk if stomach upset occurs ase the risk of heart attack or stroke	
feel faint • vomit blood • have bloody	 You experience any of the following signs of stomach bleeding: or black stools have stomach pain that does not get better pain fever gets worse or lasts more than 3 days redness or swelling is sw symptoms appear 	Do not use if individual packet is open
V2_3/24/20 7	47200 61102 4	

PRINCIPAL DISPLAY PANEL - 250 Tablet Packet Box

Product #1104

Ibuprofen

Pain Reliever/Fever Reducer

Active ingredient:

Ibuprofen

- Temporary relief from pain, fever and inflammation
- Soothes muscles aches and minor arthritis pain
- Non-drowsy, aspirin-free formula

These quality comfort tablets are formulated and packaged for use in the workplace and meet ANSI Z308.1-2015 standards. They are sealed in tamper-evident packets for safety and convenience.

250 Tablets • 2 Tablets Per Packet





IBUPROFEN 100 ibuprofen tablet, coated			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73598-1102
Route of Administration	ORAL		

	Ingredient Na	ame	Basis of St	rength Strength
BUPROFEN (UNII: \	MK2XYI10QM) (IBUPROFEN	N - UNII:WK2XYI10QM)	IBUPROFEN	200 mg
nactive Ingre	dients			
	Ingre	edient Name		Strength
CROSCARMELLOSI	E SODIUM (UNII: M280L1	LHH48)		
D&C BLUE NO. 2	(UNII: L06K8R7DQK)			
FD&C RED NO. 40	(UNII: WZB9127XOA)			
FD&C YELLOW NO	.6 (UNII: H77VEI93A8)			
HYPROMELLOSE, U	JNSPECIFIED (UNII: 3NX	W29V3WO)		
FERRIC OXIDE RED	(UNII: 1K09F3G675)			
LACTOSE MONOH	(DRATE (UNII: EWQ57Q8	I5X)		
MAGNESIUM STEA	RATE (UNII: 70097M6I30)			
MALTODEXTRIN (U	NII: 7CVR7L4A2D)			
MEDIUM-CHAIN TR	IGLYCERIDES (UNII: C9	H2L21V7U)		
MICROCRYSTALLIN	IE CELLULOSE (UNII: OF	P1R32D61U)		
POLYDEXTROSE (U	INII: VH2XOU12IE)			
POLYETHYLENE GI	YCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POLYVINYL ALCOH	OL, UNSPECIFIED (UNI	l: 532B59J990)		
POVIDONE K30 (UI	NII: U725QWY32X)			
	UNII: ETJ7Z6XBU4)			
SODIUM STARCH C	GLYCOLATE TYPE A (UN	III: H8AV0SQX4D)		
STARCH, CORN (UN	III: 08232NY3SJ)			
STEARIC ACID (UNI	I: 4ELV7Z65AP)			
TALC (UNII: 7SEV7]4	.R1U)			
TITANIUM DIOXIDE				
TRIACETIN (UNII: XI				
	·			
Product Chara	cteristics			
Color	BROWN	Score		2 pieces
Shape	ROUND	Size		10mm
Flavor		Imprint Cod	•	G2
Contains		imprint Cou	C	02
Contains				
Packaging				
			Marketing Start	Marketing End
# Item Code	Package D	escription	Date	Date
1 NDC:73598- 1102-1	50 in 1 BOX		02/02/2000	
1	2 in 1 PACKET; Type 0: Product	Not a Combination		

Marketing Category	Applica	tion Number or Mono Citation	graph	Ma	rketing Start Date	Mar	keting End Date
ANDA	ANDA07209	6		02/02	2/2000		
BUPROFEN							
buprofen tablet, o	coated						
Product Inforr	mation						
	Πατιοπ						
Product Type		HUMAN OTC DRUG	Item Co	ode (S	Source)	NDC:735	598-1104
Route of Adminis	stration	ORAL					
Active Ingredie	ent/Active	Moiety					
	Ingre	dient Name			Basis of Str	ength	Strength
IBUPROFEN (UNII: V	VK2XYI10QM) (IBUPROFEN - UNII:WK2XYI1	0QM)		IBUPROFEN		200 mg
IBUPROFEN (UNII: V	VK2XYI10QM) (IBUPROFEN - UNII:WK2XYI1	0QM)		IBUPROFEN		200 mg
		IBUPROFEN - UNII:WK2XYI1	0QM)		IBUPROFEN		200 mg
		IBUPROFEN - UNII:WK2XYI1 Ingredient Name			IBUPROFEN		200 mg
IBUPROFEN (UNII: V Inactive Ingree CROSCARMELLOSE	dients	Ingredient Name			IBUPROFEN		
Inactive Ingree	dients E S ODIUM (UN	Ingredient Name			IBUPROFEN		
Inactive Ingree	dients E SODIUM (UN (UNII: L06K8R7	Ingredient Name III: M280L1HH48) ZDQK)			IBUPROFEN		
Inactive Ingred CROSCARMELLOSE FD&C BLUE NO. 2	dients SODIUM (UN (UNII: L06K8R7 (UNII: WZB912	Ingredient Name III: M280L1HH48) ZDQK) I7XOA)			IBUPROFEN		
Inactive Ingred CROSCARMELLOSE FD&C BLUE NO. 2 FD&C RED NO. 40 FD&C YELLOW NO	dients SODIUM (UN (UNII: L06K8R7 (UNII: WZB912 . 6 (UNII: H77)	Ingredient Name III: M280L1HH48) ZDQK) I7XOA)			IBUPROFEN		
Inactive Ingred CROSCARMELLOSE FD&C BLUE NO. 2 FD&C RED NO. 40 FD&C YELLOW NO	dients SODIUM (UN (UNII: L06K8R7 (UNII: WZ B912 . 6 (UNII: H77) INSPECIFIED	Ingredient Name III: M280L1HH48) ZDQK) ZTXOA) ZEI93A8) (UNII: 3NXW29V3WO)			IBUPROFEN		
Inactive Ingree CROSCARMELLOSE FD&C BLUE NO. 2 FD&C RED NO. 40 FD&C YELLOW NO HYPROMELLOSE, U	dients SODIUM (UN (UNII: L06K8R7 (UNII: WZ B912 . 6 (UNII: H77) INSPECIFIED (UNII: 1K09F3	Ingredient Name III: M28OL1HH48) 7DQK) 7XOA) /EI93A8) (UNII: 3NXW29V3WO) G675)			IBUPROFEN		
Inactive Ingree CROSCARMELLOSE FD&C BLUE NO. 2 FD&C RED NO. 40 FD&C YELLOW NO HYPROMELLOSE, U FERRIC OXIDE RED	dients SODIUM (UN (UNII: L06K8R7 (UNII: WZ B912 . 6 (UNII: H77) INSPECIFIED 0 (UNII: 1K09F3 (DRATE (UNII:	Ingredient Name III: M280L1HH48) ZDQK) ZTXOA) /EI93A8) (UNII: 3NXW29V3WO) G675) EWQ57Q8I5X)			IBUPROFEN		
Inactive Ingree CROSCARMELLOSE FD&C BLUE NO. 2 FD&C RED NO. 40 FD&C YELLOW NO HYPROMELLOSE, U FERRIC OXIDE RED LACTOSE MONOHY MAGNESIUM STEAF	dients SODIUM (UN (UNII: L06K8R7 (UNII: WZ B912 . 6 (UNII: H77V INSPECIFIED) (UNII: 1K09F3 (DRATE (UNII: 70	Ingredient Name III: M280L1HH48) ZDQK) ZTXOA) (EI93A8) (UNII: 3NXW29V3WO) G675) EWQ57Q8I5X) 0097M6I30)			IBUPROFEN		
Inactive Ingree CROSCARMELLOSE FD&C BLUE NO. 2 FD&C RED NO. 40 FD&C YELLOW NO HYPROMELLOSE, U FERRIC OXIDE RED LACTOSE MONOHY MAGNESIUM STEAF MALTODEXTRIN (UP	dients 5 SODIUM (UN (UNII: L06K8R7 (UNII: WZ B912 . 6 (UNII: H77) INSPECIFIED 0 (UNII: 1K09F3 (DRATE (UNII: 70 NII: 7CVR7L4A2	Ingredient Name III: M280L1HH48) ZDQK) ZTXOA) (EI93A8) (UNII: 3NXW29V3WO) G675) EWQ57Q8I5X) 0097M6I30)			IBUPROFEN		
Inactive Ingree CROSCARMELLOSE FD&C BLUE NO. 2 FD&C RED NO. 40 FD&C YELLOW NO HYPROMELLOSE, U FERRIC OXIDE RED LACTOSE MONOHY MAGNESIUM STEAF MALTODEXTRIN (UP MEDIUM-CHAIN TR	dients SODIUM (UN (UNII: L06K8R7 (UNII: WZ B912 . 6 (UNII: H77) INSPECIFIED (UNII: 1K09F3 /DRATE (UNII: 70 NII: 7CVR7L4A2 IGLYCERIDES IE CELLULOS	Ingredient Name III: M280L1HH48) 7DQK) 7XOA) 7EI93A8) (UNII: 3NXW29V3WO) G675) EWQ57Q8I5X) 0097M6I30) 2D) 5 (UNII: C9H2L21V7U) E (UNII: OP1R32D61U)			IBUPROFEN		

TRIACETIN (UNII: XHX3C3X673)

TALC (UNII: 7SEV7J4R1U)

POVIDONE K30 (UNII: U725QWY32X) SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

STARCH, CORN (UNII: 08232NY3SJ) **STEARIC ACID** (UNII: 4ELV7Z65AP)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)

SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)

Product Characteristics					
Color	BROWN	Score	2 pieces		
Shape	ROUND	Size	10mm		

Flavor		Imprint Cod	Imprint Code		
Contains					
Pa	ackaging				
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:73598- 1104-1	125 in 1 BOX	02/02/2000		
1		2 in 1 PACKET; Type 0: Not a Combination Product			
Μ	arketing	nformation			
M	arketing Marketing Category	nformation Application Number or Monograp Citation	h Marketing Start Date	Marketing End Date	

Labeler - JHK Inc dba American Safety & First Aid (867236309)

Revised: 4/2024

JHK Inc dba American Safety & First Aid