# IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE- ibuprofen and pseudoephedrine hydrochloride tablet, sugar coated Sunmark

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#### Ibuprofen and Pseudoephedrine Hydrochloride

#### **Drug** Facts

Purposes
Pain reliever/fever reducer
Nasal decongestant

\* nonsteroidal anti-inflammatory drug

#### Uses

temporarily relieves these symptoms associated with the common cold or flu:

- headache
- fever
- sinus pressure
- nasal congestion
- minor body aches and pains

#### Warnings

#### 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.

#### 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

- in children under 12 years of age
- if you have ever had an allergic reaction to any other pain reliever/fever reducer
- right before or after heart surgery
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

#### Ask a doctor before use if

- stomach bleeding warning applies to you
- you have problems or serious side effects from taking pain relievers or fever reducers
- you have a history of stomach problems, such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma, thyroid disease, diabetes, have trouble urinating due to an enlarged prostate gland, 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 any other product that contains pseudoephedrine or any other nasal decongestant
- 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:
  - feel faint
  - vomit blood
  - have bloody or black stools
  - have stomach pain that does not get better
- you have symptoms of heart problems or stroke:
  - chest pain
  - trouble breathing
  - weakness in one part or side of body
  - slurred speech
  - leg swelling
- fever gets worse or lasts more than 3 days
- nasal congestion lasts for more than 7 days
- symptoms continue or get worse
- redness or swelling is present in the painful area
- you get nervous, dizzy, or sleepless
- 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. (1-800-222-1222)

#### Directions

- do not take more than directed
- the smallest effective dose should be used
- adults and children 12 years of age and over:
  - take 1 caplet every 4 to 6 hours while symptoms persist. If symptoms do not respond to 1 caplet, 2 caplets may be used.
  - do not use more than 6 caplets in any 24-hour period unless directed by a doctor
- children under 12 years of age: do not use

#### Other information

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

#### Inactive ingredients

acacia, calcium carbonate, carnauba wax, confectioner's sugar, corn starch, croscarmellose sodium, crospovidone, FD&C Blue no. 2 Aluminum Lake, FD&C Red no. 40 Aluminum Lake, FD&C Yellow no. 6 Aluminum Lake, gelatin, guar gum, hydrogenated vegetable oil, hydroxypropyl cellulose, iron oxide black, kaolin, polyethylene glycol, powdered cellulose, povidone, pregelatinized starch, propylene glycol, shellac, silicon dioxide, sodium benzoate, sucrose, talc, titanium dioxide, white wax

Questions?

call 1-800-406-7984

Distributed By McKesson One Post Street, San Francisco, CA 94104

#### PRINCIPAL DISPLAY PANEL - 200 mg/30 mg Caplet Blister Pack Carton

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COMPARE TO ADVIL<sup>®</sup> COLD & SINUS ACTIVE INGREDIENTS<sup>†</sup> NDC 49348-097-47

See New Warnings Information

cold & sinus relief

Ibuprofen and Pseudoephedrine HCl Tablets, USP

ibuprofen, USP 200 mg • pain reliever/fever reducer (NSAID)\* pseudoephedrine HCl, USP 30 mg • nasal decongestant \*nonsteroidal anti-inflammatory drug

Relieves sinus pressure, nasal congestion and fever

Non-Drowsy

20 Coated Caplets \*\* (\*\*oval-shaped tablets)



#### 0000051 ISS 720012900

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Drug Facts (continued) Inscribe ingredients exais, calcium carbonate, carrauba wax, confectioner's sugar, com starch, croscamellose sodium, crospovidone, FD&C Blue no. 2 Aluminum Lake, FD&C Red no. 40 Aluminum Lake, FD&C Vellow no. 6 Aluminum Lake, getatin, guar gum, hydrogenated vegetable oli, hydroxypropyi celludose, iron oxide black, kaolin, polyeinylene gyoot, powdered celludose, povidone, pregetatinized starch, propylene gycot, anellos and toxide, sodium benzoate, such finanum doxide, white wax polyeinylene gyoot, powdered celludose, povidone, pregetatinized starch, propylene gyoot, anellose, such finanum doxide, white wax

<b>IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE</b> ibuprofen and pseudoephedrine hydrochloride tablet, sugar coated							
Product Information							
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:49		NDC:49348	9348-097		
		nem Coue (3)	ource)	1100.4554	1340-097		
Route of Administration	ORAL						
Active Ingredient/Active Moi	etv						
	edient Name		Basis of St	rength	Strength		
IBUPROFEN (UNII: WK2XYI10QM) (IBU			IBUPROFEN	in eing ein	200 mg		
PSEUDO EPHEDRINE HYDRO CHLORIDE (UNII: 6 V9 V2RYJ8N) (PSEUDO EPHEDRINE - UNII:7CUC9 DDI9F) PSEUDO EPHEDRINE HYDRO CHLORIDE					30 mg		
Inactive Ingredients							
	Ingredient Name				Strength		
ACACIA (UNII: 5C5403N26O)							
CALCIUM CARBONATE (UNII: H0 G9 3	79FGK)						
CARNAUBA WAX (UNII: R12CBM0EIZ)							
STARCH, CORN (UNII: 08232NY3SJ)							
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)							
CROSPOVIDONE (15 MPA.S AT 5%)	(UNII: 68401960MK)						
FD&C BLUE NO.2 (UNII: L06K8R7DC	<u>I</u> K)						
FD&C RED NO. 40 (UNII: WZB9127XOA)							
FD&C YELLOW NO.6 (UNII: H77VEI9	3A8)						
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)							
GUAR GUM (UNII: E8911637KE)							
GLYCERYL TRISTEARATE (UNII: P60							
HYDROXYPROPYL CELLULOSE (12	, ,	P)					
FERROSOFERRIC OXIDE (UNII: XM0	M87F357)						
KAOLIN (UNII: 24H4NWX5CO)							
POLYETHYLENE GLYCOL, UNSPEC	,						
POWDERED CELLULOSE (UNII: SMD	,						
POVIDONE, UNSPECIFIED (UNII: FZ9							
PROPYLENE GLYCOL (UNII: 6DC9Q	167V3)						
<b>SHELLAC</b> (UNII: 46 N107B710)							
SILICON DIO XIDE (UNII: ETJ7Z6 XBU							
SODIUM BENZOATE (UNII: OJ245FE5	EU)						
SUCROSE (UNII: C151H8 M554)							
TALC (UNII: 7SEV7J4R1U)							
TITANIUM DIO XIDE (UNII: 15FIX9V2J	Р)						

WHITE WAX (UNII: 7G1J5DA97F)								
Product Characteristics								
Color brown Score no score	ore							
ShapeOVAL (Caplets)Size14mm	1							
FlavorImprint Code423	423							
Contains								
Packaging								
# Item Code Package Description Marketing Start Date Mark	eting End Date							
NDC:49348-097- 47     20 in 1 BLISTER PACK; Type 0: Not a Combination Product     10/13/2001								
Marketing Information								
Marketing Category Application Number or Monograph Citation Marketing Start Date Mark	eting End Date							
ANDA ANDA074567 10/13/2001								

### Labeler - Sunmark (177667227)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

## Establishment

Name	Address	ID/FEI	<b>Business Operations</b>
Ohm Laboratories Inc.		184769029	manufacture(49348-097)

Revised: 5/2018

Sunmark