IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE- ibuprofen and pseudoephedrine hydrochloride tablet, sugar coated Ohm Laboratories Inc.

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: Ohm Laboratories Inc. New Brunswick, NJ 08901

PRINCIPAL DISPLAY PANEL - 200 mg/30 mg Tablet Blister Pack Carton

[†]Compare To the active ingredients of Advil[®] Cold & Sinus

NDC 51660-493-41

See New Warnings Information

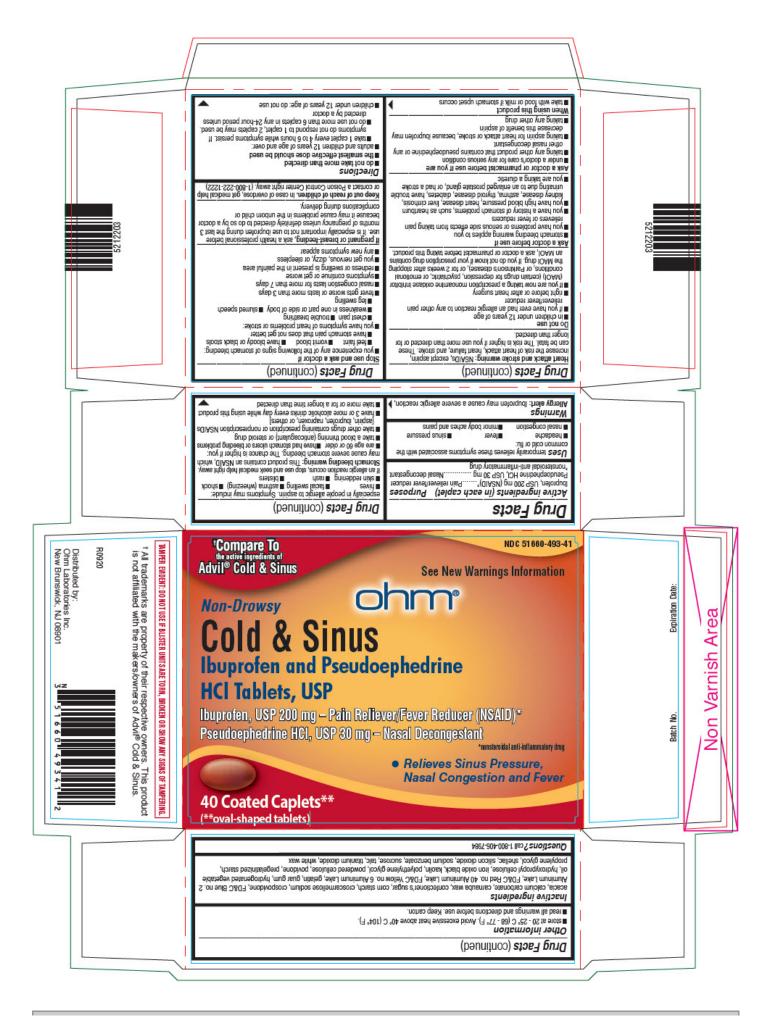
ohm®

Non-Drowsy Cold & Sinus 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 40 Coated Caplets** (**oval-shaped tablets)



Product Informa	tion					
Product Type		HUMAN OTC DRUG	Item Code (S	ource)	NDC:51660-	493
Route of Administra	ation	ORAL				
Active Ingredier	nt/Active Moie	ty				
	Ingredient Name				trength	Strengtl
IBUPROFEN (UNII: W	K2XYI10QM) (IBU	IPROFEN - UNII:WK2XYI10	QM)	IB UPRO FEN		200 mg
PSEUDO EPHEDRINE HYDRO CHLO RIDE (UNII: 6 V9 V2RYJ8N) (PSEUDO EPHEDRINE PSEUDO EPHEDR			PSEUDOEPHEDRI HYDROCHLORIDE		30 mg	
Inactive Ingredi	ents					
		Ingredient Nam	e		5	Strength
PO VIDO NE, UNSPEC		9 GH9 4E)				
ACACIA (UNII: 5C540						
CALCIUM CARBONA	TE (UNII: H0 G9 3	79 FGK)				
CARNAUBA WAX (UI	NII: R12CBM0EIZ)					
CROSCARMELLOSE	E SODIUM (UNII: 1	M28OL1HH48)				
STARCH, CORN (UN	I: O8232NY3SJ)					
FD&C BLUE NO. 2 (U	JNII: L06K8R7DQ	K)				
FD&C RED NO. 40 (U	INII: WZB9127XO	A)				
FD&C YELLOW NO.	6 (UNII: H77VEI9	3A8)				
GELATIN, UNSPECIE	IED (UNII: 2G860	N327L)				
		00000 WAMW) (UNII: RFV	V2ET671P)			
KAOLIN (UNII: 24H4N	WX5CO)					
CROSPOVIDONE (12	20 .MU.M) (UNII:	58401960MK)				
PO WDERED CELLUI	LOSE (UNII: SMD	X3XO9M)				
PROPYLENE GLYCC	L (UNII: 6DC9Q1	67V3)				
SHELLAC (UNII: 46 N	l07B71O)					
SILICON DIOXIDE (U						
SODIUM BENZOATE		EU)				
SUCROSE (UNII: C15						
TALC (UNII: 7SEV7J4						
TITANIUM DIO XIDE		<i>?</i>)				
WHITE WAX (UNII: 70						
GUAR GUM (UNII: E8						
FERROSOFERRIC O						
		FIED (UNII: 3WJQ0SDW1A	r)			
CORN OIL (UNII: 847	0G57WFM)					
Product Charact	eristics					
Color	bro wn		Score		no score	

Flavor	I	Imprint Code		423				
Contains								
Packaging								
# Item Code	Package Description		Marketing Start Date	Marketing End Date				
1 NDC:51660-493-41 40 in 1 BLISTER PACK; Type 0: Not a Combination Product 10/13/2001								
Marketing Information								
Marketing Category	Application Number or Monograph C	itation I	Marketing Start Date	Marketing End Date				
ANDA	ANDA074567	10	0/13/2001					

Labeler - Ohm Laboratories Inc. (184769029)

Establishment							
Name	Address	ID/FEI	Business Operations				
Ohm Laboratories Inc.		184769029	MANUFACTURE(51660-493)				

Revised: 9/2020

Ohm Laboratories Inc.