#### IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE- ibuprofen and pseudoephedrine hydrochloride tablet, sugar coated The Kroger Company

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#### Ibuprofen and Pseudoephedrine Hydrochloride

#### Drug Facts

Active ingredients (in each caplet)	Purposes
Ibuprofen, USP 200 mg (NSAID) <sup>*</sup>	Pain reliever/fever reducer
Pseudoephedrine HCl, USP 30 mg	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

### 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 **1-800-632-6900** 

#### DISTRIBUTED BY THE KROGER CO. CINCINNATI, OHIO 45202

# PRINCIPAL DISPLAY PANEL - 40 Tablet Blister Pack Carton

COMPARE TO the active ingredients of ADVIL® COLD & SINUS \*\*\*See side panel

NDC 30142-423-41

Kroger<sub>ℝ</sub>

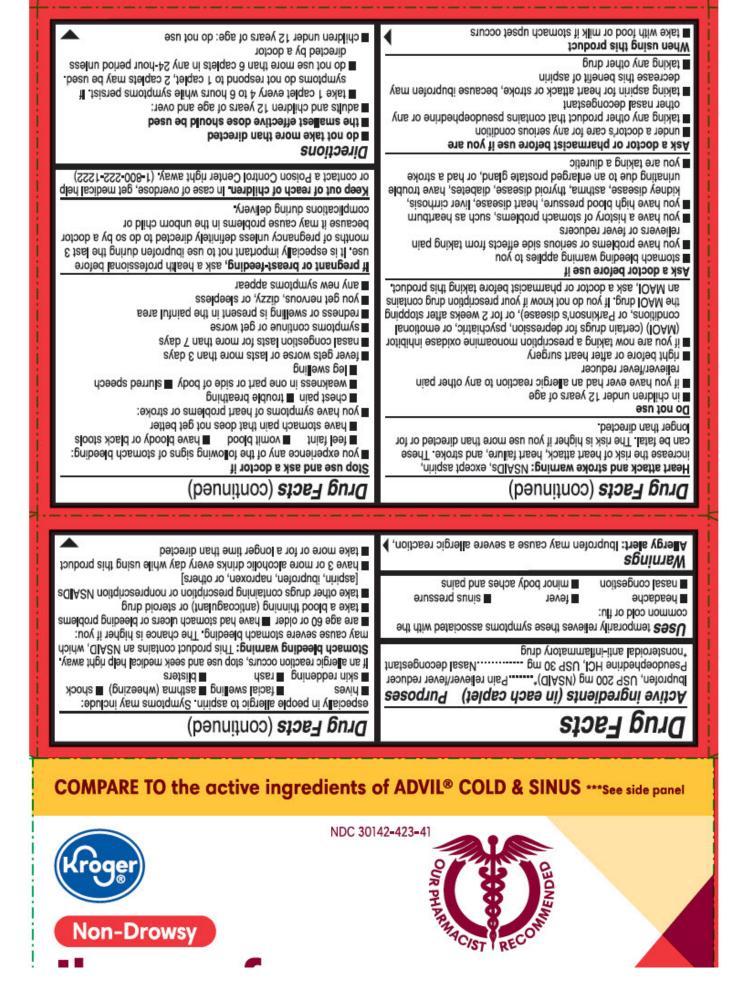
OUR PHARMACIST RECOMMENDED

Non-Drowsy Ibuprofen Cold & Sinus

Ibuprofen & 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 Fever

40 COATED CAPLETS\*\* \*\*oval-shaped tablets



# Ibuproten Cold & Sinus

Ibuprofen & Pseudoephedrine HCI Tablets, USP

Ibuprofen, USP 200 mg Pain Reliever/Fever Reducer **(NSAID)**\* Pseudoephedrine HCl, USP 30 mg Nasal Decongestant RELIEVES: Sinus Pressure

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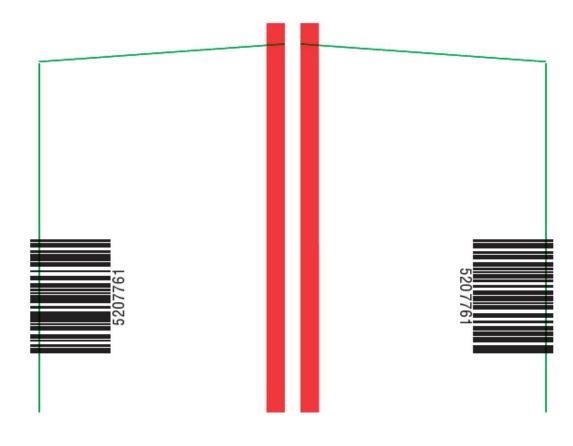
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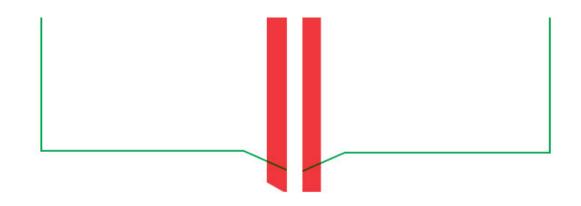
actual size

## Other information

# (continued) area (continued)







# **IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE**

ibuprofen and pseudoephedrine hydrochloride tablet, sugar coated

Product Information							
Product Type	HUMAN OTC DRUG	Item Code (S	Item Code (Source) NDC:30				
Route of Administration	ORAL						
Active Ingredient/Active N	Aniaty						
0	ngredient Name		Basis of Streng	th Strengt			
IBUPROFEN (UNII: WK2XYI10QM)	<u> </u>	M	IBUPROFEN	200 mg			
PSEUDO EPHEDRINE HYDRO CHI - UNII:7CUC9DDI9F)			PSEUDOEPHEDRINE HYDROCHLORIDE	30 mg			
Inactive Ingredients							
	Ingredient Name	2		Strength			
<b>ACACIA</b> (UNII: 5C5403N26O)							
CALCIUM CARBONATE (UNII: H0 G9379 FGK)							
CARNAUBA WAX (UNII: R12CBM0EIZ)							
STARCH, CORN (UNII: 08232NY3SJ)							
CROSCARMELLOSE SODIUM (U	JNII: M28OL1HH48)						
CROSPOVIDONE (15 MPA.S AT	<b>5%)</b> (UNII: 68401960MK)						
FD&C BLUE NO. 2 (UNII: L06K8F	R7DQK)						
FD&C RED NO. 40 (UNII: WZB9127XOA)							
FD&C YELLOW NO.6 (UNII: H77	VEI93A8)						
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)							
GUAR GUM (UNII: E8911637KE)							
GLYCERYL TRISTEARATE (UNII: P6OCJ2551R)							
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)							
	. , .	KAOLIN (UNII: 24H4NWX5CO)					
KAOLIN (UNII: 24H4NWX5CO)							
KAOLIN (UNII: 24H4NWX5CO) POLYETHYLENE GLYCOL, UNS	PECIFIED (UNII: 3WJQ0SDW1A	)					
	PECIFIED (UNII: 3WJQ0SDW1A	)					
KAOLIN (UNII: 24H4NWX5CO) POLYETHYLENE GLYCOL, UNS	<b>PECIFIED</b> (UNII: 3WJQ0SDW1A SMD1X3XO9M)	)					
KAOLIN (UNII: 24H4NWX5CO) POLYETHYLENE GLYCOL, UNS POWDERED CELLULOSE (UNII:	PECIFIED (UNII: 3WJQ0SDW1A SMD1X3XO9M) FZ989GH94E)	)					

sc	SODIUM BENZOATE (UNII: OJ245FE5EU)					
st	J <b>CROSE</b> (UNII: C153	1H8 M554)				
TALC (UNII: 7SEV7J4R1U)						
WHITE WAX (UNII: 7G1J5DA97F)						
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)						
FERROSOFERRIC OXIDE (UNII: XM0M87F357)						
P	roduct Charact	eristics				
С	olor	brown	Score		no score	
SI	ıape	OVAL (Caplets)	Size		14mm	
Fl	avor		Imprint C	ode	423	
С	Contains					
P	ackaging					
<b>P</b> #	ackaging Item Code	Package Description		Marketing Start Date	Marketing End Date	
#		U I		Marketing Start Date	Marketing End Date	
#	Item Code NDC:30142-423-41	U I	bination Produc	10/13/2001	Marketing End Date	
# 1	Item Code NDC:30142-423-41	1 in 1 CARTON	bination Produc	10/13/2001	Marketing End Date	
# 1	Item Code NDC:30142-423-41	1 in 1 CARTON	bination Produc	10/13/2001	Marketing End Date	
# 1 1	Item Code NDC:30 142-423-41	1 in 1 CARTON 40 in 1 BLISTER PACK; Type 0: Not a Com	bination Produc	10/13/2001	Marketing End Date	
# 1 1	Item Code NDC:30142-423-41	1 in 1 CARTON 40 in 1 BLISTER PACK; Type 0: Not a Com formation		10/13/2001 t		
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# 1 1 N N	Item Code NDC:30142-423-41	1 in 1 CARTON 40 in 1 BLISTER PACK; Type 0: Not a Com formation		10/13/2001 t		

# Labeler - The Kroger Company (006999528)

Registrant - Ohm Laboratories Inc. (184769029)

# Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(30142-423)

Revised: 3/2020

The Kroger Company