GOOD NEIGHBOR PHARMACY IBUPROFEN- ibuprofen tablet, chewable Proficient Rx LP

Amerisource Bergen Junior Strength Ibuprofen Tablets, 100 mg Drug Facts

Active ingredient (in each tablet)

Ibuprofen 100 mg (NSAID)*

*nonsteroidal anti-inflammatory drug

Purposes

Pain reliever/fever reducer

Uses

temporarily:

- reduces fever
- relieves minor aches and pains due to the common cold, flu, sore throat, headaches and toothaches

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 your child:

- has had stomach ulcers or bleeding problems
- takes a blood thinning (anticoagulant) or steroid drug
- takes other drugs containing prescription or nonprescription NSAIDs [aspirin, ibuprofen, naproxen, or others]
- takes 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.

Sore throat warning: Severe or persistent sore throat or sore throat accompanied by high fever,

headache, nausea, and vomiting may be serious. Consult doctor promptly. Do not use more than 2 days or administer to children under 3 years of age unless directed by doctor.

Do not use

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

Ask a doctor before use if

- stomach bleeding warning applies to your child
- child has a history of stomach problems, such as heartburn
- child has problems or serious side effects from taking pain relievers or fever reducers
- child has not been drinking fluids
- child has lost a lot of fluid due to vomiting or diarrhea
- child has high blood pressure, heart disease, liver cirrhosis, kidney disease, or had a stroke
- child has asthma
- child is taking a diuretic

Ask a doctor or pharmacist before use if the child is

- 1. under a doctor's care for any serious condition
- 2. taking any other drug

When using this product

- mouth or throat burning may occur; give with food or water
- take with food or milk if stomach upset occurs

Stop use and ask a doctor if

- child experiences any of the following signs of stomach bleeding:
- feels faint
- vomits blood
- has bloody or black stools
- has stomach pain that does not get better
- child has symptoms of heart problems or stroke:
- chest pain
- trouble breathing
- weakness in one part or side of body
- slurred speech
- leg swelling
- the child does not get any relief within first day (24 hours) of treatment
- fever or pain gets worse or lasts more than 3 days
- redness or swelling is present in the painful area
- any new symptoms appear

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

- this product does not contain directions or complete warnings for adult use
- do not give more than directed
- find right dose on chart below. If possible, use weight to dose; otherwise use age.
- if needed, repeat dose every **6-8 hours**
- do not use more than 4 times a day

Dosing Chart				
Weight (lb)	Age (yr)	Tablets		
under 24	under 2	ask a doctor		
24-35	2-3	1		
36-47	4-5	1 ½		
48-59	6-8	2		
60-71	9-10	2 ½		
72-95	11	3		

Other information

- 1. phenylketonurics: contains phenylalanine 6 mg per tablet
- 2. store between 20-25°C (68-77°F)
- 3. do not use if printed seal under cap is broken or missing

Inactive ingredients

acesulfame potassium, ammonium glycyrrhizin, aspartame, carnauba wax, croscarmellose sodium, D&C red no. 27 aluminum lake, FD&C blue no. 1 aluminum lake, hypromellose, magnesium stearate, mannitol, natural and artificial flavors, silicon dioxide, sodium lauryl sulfate, soybean oil, succinic acid

Questions or comments?

1-800-719-9260

Principal Display Panel

Compare to Motrin® Junior Strength active ingredient

See New Warnings

FOR AGES 2 TO 11

junior strength

Ibuprofen tablets, 100 mg

Pain Reliever/Fever Reducer (NSAID)

Lasts up to 8 hours

24 grape-flavored chewable tablets

Relabeled by:

Proficient Rx LP





NDC 71205-423-24

Lot #:00000 Exp. 00/00/00 SN# MASTER

> Relabeled By: Proficient Rx LP Thousand Oaks, CA 91320

Ibuprofen 100mg

#24 Chewable Tablets

Each tablets contains: Ibuprofen 100 mg (NSAID)*
Pain reliever/ fever reducer *nonsteroidal
anti-inflammatory drug

Purple (Lavender), round, scored tablet with imprint code (L521). Grape flavor

Product ID: SI042324

FOR AGES 2 to 11

Dist. By: AmerisourceBergen 1300 Morris Drive Chesterbrook, PA 19087

Store between 20-25°C (68-77°F)

Keep medication out of the reach of children

GOOD NEIGHBOR PHARMACY IBUPROFEN

ORAL

ibuprofen tablet, chewable

Route of Administration

Product Information				
Product Type H	HUMAN OTC DRUG	Item Code (Source)	NDC:71205-423(NDC:46122-010)	

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
IBUPRO FEN (UNII: WK2XYI10 QM) (IBUPRO FEN - UNII:WK2XYI10 QM)	IBUPROFEN	100 mg	

Inactive Ingredients			
Ingredient Name	Strength		
ACESULFAME POTASSIUM (UNII: 230 V73Q5G9)			
AMMO NIUM GLYCYRRHIZATE (UNII: 3VRD35U26C)			
ASPARTAME (UNII: Z0H242BBR1)			
CARNAUBA WAX (UNII: R12CBM0 EIZ)			
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)			
D&C RED NO. 27 (UNII: 2LRS 185U6K)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MANNITOL (UNII: 3OWL53L36A)			
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)			
SODIUM LAURYL SULFATE (UNII: 368GB5141J)			

SO YBEAN O IL (UNII: 241ATL177A)
SUCCINIC ACID (UNII: AB6MNQ6J6L)

Product Characteristics				
Color	PURPLE (Lavender)	Score	2 pieces	
Shape	ROUND	Size	12mm	
Flavor	GRAPE	Imprint Code	L521	
Contains				

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71205-423-24	1 in 1 CARTON	03/11/2020	
1		24 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076359	10/19/2009	

Labeler - Proficient Rx LP (079196022)

Establishment			
Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-423), RELABEL(71205-423)

Revised: 3/2020 Proficient Rx LP