CHILDRENS IBUPROFEN- ibuprofen suspension A-S Medication Solutions

Ibuprofen Oral Suspension, USP

Drug Facts

Active ingredient

(in each 5 mL)

Ibuprofen, USP 100 mg (NSAID)**

**nonsteroidal anti-inflammatory drug

Purpose

Pain reliever/fever reducer

Uses

temporarily:

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

Warnings

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

- hivesfacial swelling
- asthma (wheezing)shock
- skin reddeningrash
- 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

- 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

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
- shake well before using
- mL = milliliter
- find right dose on chart. If possible, use weight to dose; otherwise use age.
- use only enclosed dosing cup. Do not use any other dosing device.
- if needed, repeat dose every **6-8 hours**

- do not use more than 4 times a day
- replace original bottle cap to maintain child resistance

Dosing Chart			
Weight (lb)	Age (yr)	Dose (mL)*	
under 24	under 2 years	ask a doctor	
24-35 lbs	2-3 years	5 mL	
36-47 lbs	4-5 years	7.5 mL	
48-59 lbs	6-8 years	10 mL	
60-71 lbs	9-10 years	12.5 mL	
72-95 lbs	11 voars	15 mI	

^{*}or as directed by a doctor

Other information

- each 5 mL contains: sodium 2 mg
- store between 20° to 25°C (68° to 77°F)
- do not use if printed neckband is broken or missing
- see bottom panel for lot number and expiration date

Inactive ingredients

Original Berry: citric acid, D&C yellow #10, FD&C red #40, flavors, glycerin, hypromellose, polysorbate 80, purified water, sodium benzoate, sucrose, xanthan gum

Dye-Free Berry: citric acid, flavors, glycerin, hypromellose, polysorbate 80, purified water, sodium benzoate, sucrose, xanthan gum

Grape: citric acid, D&C red #33, FD&C blue #1, FD&C red #40, flavors, glycerin, hypromellose, polysorbate 80, purified water, sodium benzoate, sucrose, xanthan gum

Bubble Gum: artificial bubble gum flavor, citric acid, FD&C red #40, glycerin, hypromellose, polysorbate 80, purified water, sodium benzoate, sucrose, xanthan gum

Questions? 1-800-432-8534 between 9 am and 4 pm EST, Monday–Friday.

HOW SUPPLIED

Product: 50090-2528

NDC: 50090-2528-0 118 mL in a BOTTLE / 1 in a CARTON

Ibuprofen



CHILDRENS IBUPROFEN

ibuprofen suspension

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50090-2528(NDC:0472-1255)		

Route of Administration ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
IBUPRO FEN (UNII: WK2XYI10 QM) (IBUPRO FEN - UNII: WK2XYI10 QM)	IBUPROFEN	100 mg in 5 mL	

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)			
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
GLYCERIN (UNII: PDC6A3C0OX)			
HYPROMELLOSE 2910 (4000 MPA.S) (UNII: RN3152OP35)			
POLYSORBATE 80 (UNII: 6OZP39ZG8H)			
WATER (UNII: 059QF0KO0R)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SUCROSE (UNII: C151H8 M554)			
XANTHAN GUM (UNII: TTV12P4NEE)			

Other Ingredients			
Ingredient Kind	Ingredient Name	Quantity	
Does not contain	ALCOHOL (UNII: 3K9958V90M)	0 in 5 mL	

Product Characteristics

Color	ORANGE	Score
Shape		Size
Flavor	BERRY	Imprint Code
Contains		

]	Packaging				
#	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:50090-2528- 0	1 in 1 CARTON	10/18/2016		
1		118 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA074916	04/30/1999		

Labeler - A-S Medication Solutions (830016429)

Establishment				
Name	Address	ID/FEI	Business Operations	
A-S Medication Solutions		830016429	RELABEL(50090-2528)	

Revised: 8/2018 A-S Medication Solutions