TOPCARE CHILDRENS IBUPROFEN- ibuprofen suspension
Topco Associates LLC

Topco Associates LLC. Children’s Ibuprofen Drug Facts

Active ingredient (in each 5 mL)

Ibuprofen 100 mg (NSAID)*

*nonsteroidal anti-inflammatory drug

Purposes

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:

- 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

- under a doctor’s care for any serious condition
- taking any other drug

When using this product

- give 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
- wash dosage cup after each use

Dosing Chart		
Weight (lb)	**Age (yr)**	**Dose (mL)****
under 24 lbs	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 years	15 mL

**or as directed by a doctor

Other information

- **each 5 mL contains**: sodium 2 mg
- **do not use if printed neckband is broken or missing**
- store at 20-25°C (68-77°F)
- do not freeze

Inactive ingredients

anhydrous citric acid, D&C red #33, FD&C blue #1, FD&C red #40, glycerin, high fructose corn syrup, hypromellose, natural and artificial grape flavor, polysorbate 80, purified water, sodium benzoate, sorbitol solution, xanthan gum

Questions or comments?

1-888-423-0139

Principal Display Panel

TopCare® health

COMPARE TO CHILDREN'S MOTRIN® GRAPE FLAVOR ACTIVE INGREDIENT

children's Ibuprofen

ORAL SUSPENSION

100 mg per 5 mL

PAIN RELIEVER • FEVER REDUCER (NSAID)

Lasts up to 8 HOURS

Alcohol Free

Ages 2 to 11 Years

GRAPE FLAVOR

4 FL OZ (120 mL)

Drug Facts (continued)

- right before or after heart surgery

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TOPCO ASSOCIATES LLC
ELK GROVE VILLAGE
IL 60007
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QUESTIONS?
1-888-423-0139
topcare@topco.com
www.topcarebrand.com

Scan here for more information or call 1-888-423-0139

GLUTEN FREE

QUALITY GUARANTEED

TopCare health

children's Ibuprofen
ORAL SUSPENSION
100 mg per 5 mL
PAIN RELIEVER • FEVER REDUCER (NSAID)

NDC 36800-660-00

TopCare health

COMPARE TO CHILDREN'S MOTRIN® GRAPE FLAVOR ACTIVE INGREDIENT†

children's
Ibuprofen
ORAL SUSPENSION
100 mg per 5 mL
PAIN RELIEVER • FEVER REDUCER (NSAID)

Lasts up to 8 HOURS

Alcohol Free

4 FL OZ (120 mL)

†This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Children's Motrin® Grape Flavor.

0 36800 06098 2

Code Area

66026 88 C11

Important: Read all product information before using. Keep this box for important information. This product is intended for use in children.

Drug Facts

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Ibuprofen 100 mg (NSAID)*	Pain reliever/ fever reducer

*nonsteroidal anti-inflammatory drug

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TOPCARE CHILDRENS IBUPROFEN
ibuprofen suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:36800-660
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	100 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	PURPLE (opaque)	**Score**	
Shape		**Size**	
Flavor	GRAPE	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-660-26	1 in 1 CARTON	07/02/2002	02/28/2022
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:36800-660-28	1 in 1 CARTON	03/12/2013	11/30/2021
2		148 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:36800-660-00	1 in 1 CARTON	01/01/2020	
3		120 mL 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	ANDA074937	07/02/2002	

Labeler - Topco Associates LLC (006935977)

Revised: 5/2022

Topco Associates LLC