UP AND UP IBUPROFEN- ibuprofen suspension Target Corporation

Target Corporation Children's Ibuprofen Oral Suspension 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, artificial mixed berry flavor, glycerin, high fructose corn syrup, hypromellose, polysorbate 80, purified water, sodium benzoate, sorbitol solution, xanthan gum

Questions?

Call 1-888-547-7400

Principal Display Panel

see new warnings

Compare to active ingredient in Children's Motrin® Dye-Free

dye-free

children's

ibuprofen

oral suspension

100 mg per 5 mL

pain reliever/fever reducer (NSAID)

alcohol free

non-staining

LASTS UP TO 8 HOURS

BERRY FLAVOR

AGES 2 TO 11 YEARS

4 FL OZ (118 mL)



(NSAID)

Important: Read all product information before

Drug Facts

Active ingredient Purposes

(in each 5 mL) buprofen 100 mg (NSAID)*..... ..Pain reliever/ fever reducer *nonsteroidal anti-inflammatory drug

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8

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2 TO 11

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Drug Facts (continued)

■ right before or after heart surgery

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natorqudi **children's** dye-free

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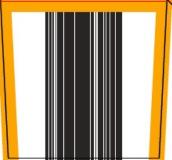
Inactive ingredients anhydrous citric acid, artificial mixed berry flavor, glycerin, high fructose corn syrup, hypromellose, polysorbate 80, purified water, sodium berzoate, sorbitol solution, xanthan gum

Questions? Call 1-888-547-7400

GLUTEN FREE

100% satisfaction quaranteed or

your money back.
relcome any questions you may have at et.com/comments or 1-800-910-6874 094 01 0636 R02 ID225432
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up&up

4 FL OZ (118 mL)



Code Area

UP AND UP IBUPROFEN

ibuprofen suspension

Product Information

Route of Administration

Product Type

HUMAN OTC DRUG

ORAL

Item Code (Source)

NDC:11673-685

Active Ingredient/Active Moiety

ı	Ingredient Name	Basis of Strength	Strength
ı	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)		
GLYCERIN (UNII: PDC6A3C0OX)		
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)		
HYPROMELLOSES (UNII: 3NXW29V3WO)		
POLYSORBATE 80 (UNII: 6 OZP39 ZG8 H)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SORBITOL (UNII: 506T60A25R)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Product Characteristics			
Color	WHITE	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-685-26	1 in 1 CARTON	07/10/2009	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11673-685-34	1 in 1 CARTON	03/12/2011	
2		237 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:11673-685-28	1 in 1 CARTON	08/24/2012	11/28/2014
3		148 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	07/10/2009	

Labeler - Target Corporation (006961700)

Revised: 10/2017 Target Corporation