CHILDRENS IBUPROFEN - ibuprofen suspension/ drops
Chain Drug Marketing Association Inc.

Quality Choice Children's Ibuprofen Oral Suspension 634

ACTIVE INGREDIENT (in each 5 mL)

Ibuprofen, USP 100 mg (NSAID)*
*nonsteroidal anti-inflammatory drug

PURPOSE

Pain reliever/fever reducer

USE(S)

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
- asthma (wheezing)
- skin reddening
- facial swelling
- shock
- 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

- take with food or milk if stomach upset occurs

STOP USE AND ASK 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 the 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 years	15 mL

*or as directed by a doctor

OTHER INFORMATION

- **each 5 mL contains:** sodium 2 mg
- store between 20-25ºC (68-77ºF)

- **do not use if carton is opened or if printed inner seal imprinted with "SEALED for YOUR PROTECTION" is broken or missing.**

INACTIVE INGREDIENTS

anhydrous citric acid, D&C yellow no. 10, FD&C red no. 40, flavor, glycerin, polysorbate 80, pregelatinized starch, purified water, sodium benzoate, sucrose, xanthan gum.

PRINCIPAL DISPLAY PANEL

NDC83324-012-04

QUALITY CHOICE

***Compare to the Active Ingredient in Children's Motrin®**
Children's Ibuprofen

Pain Reliever/Fever Reducer
Oral Suspension, USP (NSAID)
100 mg per 5 mL
For Ages 2 to 11 years

Lasts up to 8 Hours

Alcohol Free

Mixed Berry Flavor
4 FL OZ (120 mL)

Active ingredient (in each 5 mL) ***Purpose***
Ibuprofen, USP 100 mg (NSAID)*....Pain reliever/fever reducer
*nonsteroidal anti-inflammatory drug

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

PEEL HERE FOR CONTINUED IMPORTANT INFORMATION

NDC 82424-012-04

Compare to the Active Ingredient in Children's Motrin®

Children's Ibuprofen

Pain Reliever/Fever Reducer

Oral Suspension, USP (NSAID)
100 mg Per 5 mL
For Ages 2 to 11 Years
Lasts up to 8 Hours
Alcohol Free

Mixed Berry Flavor

4 FL OZ (120 mL)

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

Do not use if printed inner seal imprinted with "SEALED for YOUR PROTECTION" is broken or missing

This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., owner of the registered trademark Motrin®.

Distributed by CLIMA, Inc.
Novi, MI 48375
www.qualitychoice.com
Questions 800-935-2362

MFG# 82041 REV 0823

LOT:

EXP:

UNWARNISHED AREA TO BE LEFT IN ASK FOR LOT & EXP INFORMATION

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 2 years of age unless directed by a 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 reliever/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 the 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 ■ the right dose is in dose. 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

NON PRINTING AREA

NON PRINTING AREA

Dosing Chart

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Other information

■ **each 5 mL contains:** sodium 2 mg
■ store between 20-25°C (68-77°F)
■ **do not use if printed inner seal imprinted with "SEALED for YOUR PROTECTION" is broken or missing.**

Inactive ingredients anhydrous citric acid, D&C yellow no. 10, FD&C red no. 40, flavor, glycerin, polyacrylate 30, pregelatinized starch, purified water, sodium benzoate, sucrose, xanthan gum

CHILDRENS IBUPROFEN

ibuprofen suspension/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-012
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 YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	

POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCROSE (UNII: C151H8M554)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	ORANGE	**Score**	
Shape		**Size**	
Flavor	BERRY (Mixed Berry)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-012-04	1 in 1 CARTON	10/25/2023	
1		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	ANDA210149	10/25/2023	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Establishment

Name	Address	ID/FEI	Business Operations
Guardian Drug Company		119210276	MANUFACTURE(83324-012)

Revised: 10/2023 Chain Drug Marketing Association Inc.