IBUPROFEN ORAL - ibuprofen oral suspension Camber Consumer Care Inc

Ibuprofen Oral Suspension USP, 100 mg/5 mL (OTC)

ACTIVE INGREDIENT(S)

(in each 5 mL)

Ibuprofen 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
- 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 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
- \cdot 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 20-25°C (68-77°F)
- do not use if carton is opened or printed bottle neckband is broken or missing
- see bottom panel for lot number and expiration date

INACTIVE INGREDIENT SECTION

Berry flavor (with dye)

Acesulfame potassium, anhydrous citric acid, carboxymethylcellulose sodium, D&C Yellow #10, FD&C Red #40, flavors, glycerin, microcrystalline cellulose, polysorbate 80, propylene glycol, purified water, sodium benzoate, sucrose, and xanthan gum.

Berry flavor (dye free)

Acesulfame potassium, anhydrous citric acid, carboxymethylcellulose sodium, flavors, glycerin, microcrystalline cellulose, polysorbate 80, propylene glycol, purified water, sodium benzoate, sucrose, and xanthan gum.

Grape flavor

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Acesulfame potassium, anhydrous citric acid, carboxymethylcellulose sodium, D&C Red #33, FD&C Blue #1, flavors, glycerin, microcrystalline cellulose, polysorbate 80, propylene glycol, purified water, sodium benzoate, sucrose, and xanthan gum.

Bubble Gum flavor

Acesulfame potassium, anhydrous citric acid, carboxymethylcellulose sodium, D&C Red #33, FD&C Red #40, flavors, glycerin, microcrystalline cellulose, polysorbate 80, propylene glycol, purified water, sodium benzoate, sucrose, and xanthan gum.

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QUESTIONS SECTION

Call 1-888-588-1418 from 9 AM to 5 PM EST, Monday-Friday.



Distributed by:

Camber Consumer Care, Inc. Piscataway, NJ 08854, USA

PRINCIPAL DISPLAY PANEL

Ibuprofen oral suspension berry flavor with dye container carton



Ibuprofen oral suspension berry flavor with dye free container carton







Purpose

- ^ - ^

Children's **Ibuprofen**

Oral Suspension (NSAID)

For Ages 2 to 11 Years

CAMBER

Drug Facts (continued)

- 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:

- stomach bleedings are so that the control of the co

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 up the force 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
- use only enclose unsurp supported 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

Drug Facts (continued)

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

Inactive ingredients
Acesufiame potassium, anhydrous citric acid, carboxymethylcellulose sodium, D&C Red #33, FD&C Red #40, flavors, glycerin, microcrystalline cellulose, polysorbate 80, proplene glycol, purified water, sodium benzoale, sucrose, and xanthan gum

Questions or comments? all 1-888-588-1418 from 9 AM to 5 PM EST,

This product is not manufactured or distributed by McNeil nsumer Healthcare, distributor of Children's Motrin®.



Distributed by:
Camber Consumer Care, Inc.
Piscataway, NJ 08854, USA

BX-100

NDC 69230-310-11

See New Warnings

pare to the active ingredient in Children's Motrin®**

For ages 2 to 11 years

Children's **Ibuprofen**

Oral Suspension (NSAID) 100 mg per 5 mL

Pain Reliever/ Fever Reducer



Alcohol Free

4 FL OZ (118 mL)

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

Drug Facts

Active ingredient (in each 5 mL)

lbuprofen 100 mg (NSAID)*..... *nonsteroidal anti-inflammatory drug Pain reliev

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:

■ shock ■ skin reddening ■ rash
■ bisters
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

- Do not use
 if the child has ever had an allergic reaction to lbuprofen 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 beauthurs.
- entit thas a instory or storiest proteins, soon as hearthurn
 child has problems or serious side effects from taking
 which has been reducers
 a child less not been driving fluids
 child has lost a lot of fluid due to vomiting or diarrhea
 thinks high blood pressure, heart disease, liver
 cirrhoss, kidney disease, or had a stroke



Bubble Gum Flavor

IBUPROFEN ORAL

ibuprofen oral suspension

Product Information HUMAN OTC DRUG NDC:69230-308 **Product Type** Item Code (Source) **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	
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
GLYCERIN (UNII: PDC6A3C0OX)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)		
POLYSORBATE 80 (UNII: 60ZP39ZG8H)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SUCROSE (UNII: C151H8M554)		
XANTHAN GUM (UNII: TTV12P4NEE)		

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

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
NDC:69230-308-	1 in 1 CARTON	11/23/2018			
1	118 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	ANDA210602	11/23/2018		

IBUPROFEN ORAL

ibuprofen oral suspension

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69230-309

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
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)	
GLYCERIN (UNII: PDC6A3C0OX)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCROSE (UNII: C151H8M554)	
XANTHAN GUM (UNII: TTV12P4NEE)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Product Characteristics			
Color	purple	Score	
Shape		Size	
Flavor	GRAPE	Imprint Code	
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:69230-309- 11	1 in 1 CARTON	09/18/2019		
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:69230-309- 12	1 in 1 CARTON	09/18/2019		
2		237 mL in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing In	formation		
Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
ANDA	ANDA210602	09/18/2019	

IBUPROFEN ORAL

ibuprofen oral suspension

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69230-310

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
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)

GLYCERIN (UNII: PDC6A3C0OX)

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)

POLYSORBATE 80 (UNII: 60ZP39ZG8H)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

WATER (UNII: 059QF0KO0R)

SODIUM BENZOATE (UNII: OJ245FE5EU)

SUCROSE (UNII: C151H8M554)

XANTHAN GUM (UNII: TTV12P4NEE)

FD&C RED NO. 40 (UNII: WZB9127XOA)

D&C RED NO. 33 (UNII: 9DBA0SBB0L)

Product Characteristics

Color	pink	Score
Shape		Size
Flavor	BUBBLE GUM	Imprint Code

Contains

			in	

_	· uctuaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:69230-310- 11	1 in 1 CARTON	09/18/2019				
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product					

2	NDC:69230-310- 12	1 in 1 CARTON	09/18/2019	
2		237 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	ANDA210602	09/18/2019			

IBUPROFEN ORAL

ibuprofen oral suspension

Product	Informati	on
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:69230-311

Route of Administration ORAL

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

Inactive Ingredients	
Ingredient Name	Strength
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)	
GLYCERIN (UNII: PDC6A3C0OX)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCROSE (UNII: C151H8M554)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics					
Color		Score			
Shape		Size			
Flavor	BERRY	Imprint Code			
Contains					

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:69230-311- 11	1 in 1 CARTON	09/18/2019				
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product					
2	NDC:69230-311- 12	1 in 1 CARTON	09/18/2019				
2		237 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	ANDA210602	09/18/2019				

Labeler - Camber Consumer Care Inc (079539968)

Establishment			
Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(69230-308, 69230-309, 69230-310, 69230-311)

Revised: 1/2022 Camber Consumer Care Inc