EQUATE IBUPROFEN- ibuprofen tablet, chewable Wal-Mart Stores Inc

Wal-Mart Ibuprofen Tablets, 100 mg Drug Facts

Active ingredient (in each tablet)

Ibuprofen 100 mg (NSAID)*

*nonsteroidal anti-inflammatory drug

Purposes

Pain reliever/fever reducer

Uses

temporarily:

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

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 chances are 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

- mouth or throat burning may occur; give with food or water
- 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

- chew or crush tablets completely before swallowing
- this product does not contain directions or complete warnings for adult use
- do not give more than directed
- find right dose on chart below. If possible, use weight to dose; otherwise use age.
- if needed, repeat dose every **6-8 hours**
- do not use more than 4 times a day

Dosing Chart						
Weight (lb)	Age (yr)	Tablets				
under 24	under 2	ask a doctor				
24-35	2-3	1				
36-47	4-5	1 1⁄2				
48-59	6-8	2				
60-71	9-10	2 1/2				
72-95	11	3				

Other information

- phenylketonurics: contains phenylalanine 6 mg per tablet
- store at 20-25°C (68-77°F)
- do not use if printed seal under cap is broken or missing

Inactive ingredients

acesulfame potassium, ammonium glycyrrhizin, aspartame, carnauba wax, croscarmellose sodium, D&C red no. 27 aluminum lake, FD&C blue no. 1 aluminum lake, hypromellose, magnesium stearate, mannitol, natural and artificial flavors, silicon dioxide, sodium lauryl sulfate, soybean oil, succinic acid

Questions or comments?

1-888-287-1915

Principal Display Panel

equate[™]

Compare to Children's Motrin® active ingredient

FOR AGES 2 to 11 YEARS

children's

IBUPROFEN

Chewable Tablets, 100 mg

LASTS UP TO 8 HOURS

Pain Reliever/Fever Reducer (NSAID)

Chewables

Grape Flavor

Chew or crush tablets completely before swallowing

24 CHEWABLE TABLETS



EQUATE IBU buprofen tablet,								
Product Infor	mation							
Product Type		HUMAN OTC DRUG	ltem	Code (Source)	NDC:49	035-521	
		ORAL	nem	couc (bource,			
Route of Admin	istration	URAL						
Active Ingred	ient/Active	Moiety						
	Ingred	lient Name			Basis of St	rength	Strength	
BUPROFEN (UNII:	WK2XYI10QM) (I	BUPROFEN - UNII:WK2XYI10	QM)		IBUPROFEN		100 mg	
Inactive Ingre	dients							
		Ingredient Name				S	Strength	
ACESULFAME POT	TASSIUM (UNII:	230V73Q5G9)						
AMMONIUM GLYC	YCYRRHIZATE (UNII: 3VRD35U26C)							
ASPARTAME (UNII:	AME (UNII: Z0H242BBR1)							
CARNAUBA WAX (UNII: R12CBM0EIZ)								
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)								
D&C RED NO. 27 (UNII: 2LRS185U6K)								
FD&C BLUE NO. 1	. (UNII: H3R47K3	TBD)						
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)								
MAGNESIUM STEARATE (UNII: 70097M6I30)								
MANNITOL (UNII: 30WL53L36A)								
SILICON DIOXIDE	(UNII: ETJ7Z6XB	U4)						
SODIUM LAURYL S	SULFATE (UNII:	368GB5141J)						
SOYBEAN OIL (UN	II: 241ATL177A)							
SUCCINIC ACID (U	NII: AB6MNQ6J6	L)						
Product Chara	acteristics							
Color	PURPLE (L	avender)	Sc	core		2 pi	2 pieces	
Shape	ROUND		Siz	Size		12n	12mm	
Flavor	GRAPE		Im	Imprint Code		L52	L521	
Contains								
Packaging								
# Item Code	Pac	kage Description		Mark	ceting Start Date		eting End Date	
1 NDC:49035-521- 62	1 in 1 CARTON			10/25/20	011			
1	24 in 1 BOTTL Product	E; Type 0: Not a Combinati	on					
2 NDC:49035-521-	2 in 1 CARTON	l		10/25/20	011	06/30/20	21	

2	Product						
Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
ANDA	ANDA076359	10/25/2011					

Labeler - Wal-Mart Stores Inc (051957769)

Revised: 10/2022

Wal-Mart Stores Inc