UP AND UP INFANTS ACETAMINOPHEN- acetaminophen suspension Target Corporation

Target Corporation Infants' Acetaminophen Drug Facts

Active ingredient (in each 5 mL)

Acetaminophen 160 mg

Purpose

Pain reliever/fever reducer

Uses

temporarily:

- reduces fever
- relieves minor aches and pains due to:
- the common cold
- flu
- headache
- sore throat
- toothache

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if your child takes

- more than 5 doses in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if your child has ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if your child has

liver disease

Ask a doctor or pharmacist before use if your child is

taking the blood thinning drug warfarin

When using this product

do not exceed recommended dose (see overdose warning)

Stop use and ask a doctor if

- new symptoms occur
- pain gets worse or lasts more than 5 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present. These could be signs of a serious condition.

Keep out of reach of children.

Overdose warning: In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical even if you do not notice any signs or symptoms.

Directions

- this product does not contain directions or complete warnings for adult use
- do not give more than directed (see overdose warning)
- shake well before using
- mL = milliliter
- find right dose on chart. If possible, use weight to dose; otherwise, use age.
- push air out of syringe. Insert syringe tip into bottle opening.
- flip bottle upside down. Pull yellow part of syringe to the first dose line and then push product back into bottle.
- pull yellow part of syringe until it reaches and stays at the correct dose
- dispense liquid slowly into child's mouth, toward inner cheek
- repeat dose every 4 hours while symptoms last
- do not give more than 5 times in 24 hours
- replace cap tightly to maintain child resistance

Dosing Chart					
Weight (lb)	Age (yr)	Dose (mL)*			
under 24	under 2 years	ask a doctor			

24-35 2-3 ye	ars 5 mL
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* or as directed by a doctor

Attention: use only enclosed syringe specifically designed for use with this product. Do not use any other dosing device.

Other information

- store at 20-25°C (68-77°F)
- do not use if printed neckband is broken or missing

Inactive ingredients

anhydrous citric acid, calcium sulfate, carrageenan, flavor, glycerin, hydroxyethyl cellulose, microcrystalline cellulose and carboxymethylcellulose sodium, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose, tribasic sodium phosphate

Questions or comments

1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredient in Infants' Tylenol[®] Oral Suspension for ages 2 to 3 years dye-free infants' acetaminophen 160 mg per 5 mL fever reducer/pain reliever oral suspension ibuprofen free alcohol free aspirin free paraben free CHERRY FLAVOR Use only with enclosed syringe See side panel for more information SYRINGE INCLUDED AGES 2 to 3 YEARS up&up™

2 FL OZ (59 mL)



UP AND UP INFANTS ACETAMINOPHEN

acetaminophen suspension

Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-266			
Route of Administration	ORAL					
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Active Ingredient/Active Moiety						

Inactive Ingredients Ingredient Name Strength ANHYDROUS CITRIC ACID (UNII: XF417D3P5L) Strength CALCIUM SULFATE, UNSPECIFIED FORM (UNII: WATODDB505) CARAGEENAN (UNII: SC69YCD2Y)) Strength GUYCERIN (UNII: DC6A3CO0X) Strength Strength HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D) Strength Strength MICROCRYSTALLINE CELULOSE (UNII: 0P1R32061U) Strength Strength CARBOXYMETHYL CELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311) Strength Strength PROPYLENE GLYCOL (UNII: S029167V3) WATER (UNII: 0590F0K00R) Strength Strength StoBIUM BENZOATE (UNII: 01/245FE5U) StoBIUM BENZOATE (UNII: 8KW8220702) Strength Strength StoBIUM PHOSPHATE, TRIBASIC (UNII: A752(30A6X) Imprint Code Strength Strength Product Characteristics Strength Strength Strength Strength Packaging Y Marketing Start Marketing End Date Date 1 10 CARTON 04/21/2015 Strength Date Date 1 1 Sp mL in 1 BOTHE; Type 0: Not a Combination 04/21/2015 Strength Date Date	Inactive Ingredier ANHYDROUS CITRIC AC CALCIUM SULFATE, UN CARRAGEENAN (UNII: 5C GLYCERIN (UNII: PDC6A3 HYDROXYETHYL CELLU MICROCRYSTALLINE CE CARBOXYMETHYLCELL PROPYLENE GLYCOL (U WATER (UNII: 059QF0KO SODIUM BENZOATE (UN SORBITOL SOLUTION (I SUCRALOSE (UNII: 96K6) SODIUM PHOSPHATE, T	Ingredient Name Ingredient Name CID (UNII: XF417D3PSL) ISPECIFIED FORM (UNII: WAT0DDB505) C69YCD2YJ) 3C00X) JLOSE, UNSPECIFIED (UNII: T4V6TWG28D) ELLULOSE (UNII: OP1R32D61U) JUOSE SODIUM, UNSPECIFIED (UNII: K679 JNII: 6DC9Q167V3) OR) NII: 0J245FE5EU) UNII: 8KW3E20702) UQ3Z D4)		160 mg in 5 m Strength
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Labeler - Target Corporation (006961700)

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Target Corporation