UP AND UP CHILDRENS ACETAMINOPHEN- acetaminophen suspension Target Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Target Corporation Children's 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

- pain gets worse or lasts more than 5 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- 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 below. If possible, use weight to dose; otherwise, use age.
- remove the child protective cap and squeeze your child's dose into the dosing cup
- repeat dose every 4 hours while symptoms last
- do not give more than 5 times in 24 hours

Weight (lb)	Age (yr)	Dose (mL)*
under 24	under 2 years	ask a doctor
24-35	2-3 years	5 mL
36-47	4-5 years	7.5 mL
48-59	6-8 years	10 mL
60-71	9-10 years	12.5 mL
72-95	11 years	15 mL

^{*}or as directed by a doctor

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

Other information

- each 5 mL contains: sodium 3 mg
- 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, FD&C red #40, flavor, glycerin, high fructose corn syrup, hydroxyethyl cellulose, microcrystalline cellulose and carboxymethylcellulose sodium, propylene glycol, propylparaben, purified water, sodium benzoate, sorbitol solution, tribasic sodium phosphate

Questions or comments?

Call 1-888-547-7400

Principal Display Panel

Compare to active ingredient in Children's Tylenol®

Oral Suspension

for ages 2 to 11 years

children's acetaminophen 160 mg per 5 mL

oral suspension

fever reducer/pain reliever

ibuprofen free

alcohol free

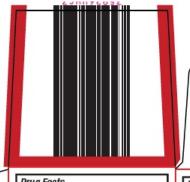
aspirin free

STRAWBERRY FLAVOR

DOSAGE CUP INCLUDED

AGES 2-11 YEARS

4 fl oz (118 mL)



children's acetaminophen 160 mg/5 mL

oral suspension fever reducer/pain reliever





NDC 11673-759-26

Compare to active ingredient in Children's Tylenol® **Oral Suspension****

for ages 2 to 11 years

children's acetaminophen 160 mg per 5 mL

oral suspension fever reducer/pain reliever

aspirin free



children's

acetaminophen

160 mg per 5 mL

DIRECTIONS: STEP 1. Shake well before using



STEP 2. Remove child protective



STEP 3. Squeeze you child's dose into the dosing cup



Please see dosing chart or ask a doctor for your child's correct dose.

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Drug Facts (continued)

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Questions or comments? Call 1-888-547-7400

"This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Children's Tylenol® Oral Suspension.

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4 fl oz (118 mL)





INCLUDED

AGES 2-11

75926 UW C7

UP AND UP CHILDRENS ACETAMINOPHEN

acetaminophen suspension

Product Information

Product Type

HUMAN OTC DRUG Item Code (Source)

NDC:11673-759

Route of Administration

ORAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	160 mg in 5 mL

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
CALCIUM SULFATE, UNSPECIFIED FORM (UNII: WAT0 DDB505)		
CARRAGEENAN (UNII: 5C69 YCD2YJ)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
GLYCERIN (UNII: PDC6A3C0OX)		
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)		
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)		
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679 OBS 311)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SORBITOL (UNII: 506T60A25R)		
SODIUM PHO SPHATE, TRIBASIC (UNII: A752Q30 A6 X)		

Product Characteristics			
Color	RED	Score	
Shape		Size	
Flavor	STRAWBERRY	Imprint Code	
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11673-759-26	1 in 1 CARTON	07/16/2011		
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:11673-759-28	1 in 1 CARTON	07/16/2011	06/18/2014	
2		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
OTC monograph not final	part343	07/16/2011	

Labeler - Target Corporation (006961700)

Revised: 12/2019 Target Corporation