CHILDREN ACETAMINOPHEN ORAL SOLUTION- acetaminophen liquid VERITYRX. LLC

APAP Child 160MG/5mL

Active ingredient (per 5 mL teaspoonful)

Acetaminophen 160 mg

Purpose

Pain Reliever/Fever Reducer

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 thoat 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 is allergic to acetaminophe or any of the inactive indreadients in this product.

Ask a doctor before use if your child hasliver disease.

Ask a doctor before use if your child istaking the blood thinning drug warfarin.

When using this product: Do not exceed recommended dose.

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.

Uses

- temporarily relieves minor aches and pains due to:
- headache
- toothache

- the common cold
- temporarily reduces fever
- fli
- sore throat

If pregnant or breast-feeding, ask a health professional before use

Keep out of reach of children. In case of overdose, get medical help or contact a Paoison Control Center right away. Quick medical attention is critical for adults/children even if you do not notice any signs or symptoms.

Directions

- do not take more than directed
- find dose in chart below
- if possible, use weight to dose, otherwise use age
- dosage may be repeasted every 4 hours, or as directed by a doctor
- do not give more than 5 doses in 24 hours

Weight (lbs)	under 24	24-35	36-47	48-59	60-71	72-95
Age (years)	under 2	2 to 3	4 to 5	6 to 8	9 to 10	11
Dosage	consult	1 tsp (5	1 1/2 tsp	2 tsp (10	2 1/2 tsp	3 tsp (15
Teaspoonful (tsp)	Physician	mL)	(7.5 mL)	mL)	(12.5 mL)	mL)

Inactive ingredients: artificial and natural cherry flavor, citric acid, FD&C Red #40, methylparaben, polyethylene glycol, propylene glycol, propylparaben, purified water, sodium citrate, sucralose



CHILDREN ACETAMINOPHEN ORAL SOLUTION

acetaminophen liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83939-0010(NDC:0904-7014)	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII: 36209ITL9D)	ACETAMINOPHEN	160 mg in 5 mL		

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
METHYLPARABEN (UNII: A218C7H19T)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			
WATER (UNII: 059QF0KO0R)			
SODIUM CITRATE (UNII: 1Q73Q2JULR)			

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:83939- 0010-1	1 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product	11/10/2025		
2	NDC:83939- 0010-2	10 in 1 BOX, UNIT-DOSE	11/10/2025		
2		1 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			
3	NDC:83939- 0010-3	40 in 1 BOX, UNIT-DOSE	11/10/2025		
3		1 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			
4	NDC:83939- 0010-4	50 in 1 BOX, UNIT-DOSE	11/10/2025		
4		1 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			
5	NDC:83939- 0010-5	100 in 1 BOX, UNIT-DOSE	11/10/2025		
5		1 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M013	10/01/2019		

Labeler - VERITYRX. LLC (097807737)

Revised: 11/2025 VERITYRX. LLC