PANADOL- acetaminophen suspension Haleon US Holdings LLC

Drug Facts

Active ingredient (in each 5 mL)

Acetaminophen 160 mg

Purposes

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
 - colds
 - flu
 - headache
 - toothache
- temporarily reduces fever

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.

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 you are allergic to acetaminophen or any of the inactive ingredients in this product

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

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
- redness or swelling is present
- any new symptoms appear

These could be signs of a serious condition.

Keep out of reach of children.

Overdose warning:

Taking more than the recommended dose can cause serious health problems. In case of overdose, get medical help or contact a Poison Control Center right away. 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 take more than directed(see overdose warning)
- find right dose in chart. If possible use weight to dose; otherwise, use age
- if needed, repeat dose every 4 hours while symptoms persist or as directed by a doctor
- do not take more than 5 doses in 24 hours, unless directed by a doctor
- use only with enclosed pre-marked measuring syringe for accuracy. Do not use any other dosing device.

Age	Weight	Dosage
under 2 yrs	under 24 lbs	ask a doctor
2-3 yrs	24-35 lbs	5 mL

Other information

store below 30°C (86°F)

Inactive ingredients

benzoic acid, FD&C red no. 40, flavor, glycerin, hydrochloric acid*, polyethylene glycol, potassium sorbate, propylene glycol, purified water, sodium hydroxide*, sodium saccharin, sorbitol solution * contains one or more of these ingredients

Questions or comments?

1-800-455-7139(English/Spanish) weekdays

Keep Carton

Principal Display Panel

NDC 0135-0539-02

Panadol ®

Infants'

ACETAMINOPHEN

160 mg per 5 mL

LIQUID

Pain Reliever

Fever Reducer

SEE NEW WARNINGS INFORMATION

Ages 2-3 years

Fast relief of fever and pain

Gentle on your stomach

- Ibuprofen free
- No sugar added
- Aspirin free

artificial

raspberry flavor

Dosing Device Inside

I.85 fl oz (54.7 mL)

Tamper Evident Feature: Do not use if printed overwrap is missing or broken.

READ AND KEEP CARTON FOR COMPLETE INFORMATION

GlaxoSmithKlineConsumer Healthcare, L.P.

Moon Township, PA 15108

Made in Canada

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USING THE CONVENIENT DOSING DEVICE

- Shake the bottle well. Push the dosing device firmly into the bottle's neck.
- **Turn**the bottle upside down. Holding the device in place, gently pull the plunger out to draw the correct dose.
- With the bottle in an upright position, remove the device from the plug and bottle by gently **pulling**it. Place the end of the device into the child's mouth. Administer dose by pressing the plunger slowly.

After use, replace the cap. **Wash**the device in warm water and allow to dry.

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102012XB (A134044)



PANADOL

acetaminophen suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0135-0539
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	
BENZOIC ACID (UNII: 85KN0B0MIM)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
GLYCERIN (UNII: PDC6A3C0OX)		
HYDROCHLORIC ACID (UNII: QTT17582CB)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
SORBITOL (UNII: 506T60A25R)		

Product Characteristics			
Color	red (light red)	Score	
Shape		Size	
Flavor	RASPBERRY	Imprint Code	
Contains			

Packaging				
# Ite	m Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC: 0539	0135- -02	1 in 1 CARTON	10/15/2012	
1		54.7 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 Drug	M013	10/15/2012	

Revised: 2/2024 Haleon US Holdings LLC