# M-PAP- acetaminophen liquid VERITYRX, LLC

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M-PAP

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# **Acetaminophen Liquid**

160 mg/5 mL

Pain Reliever / Fever Reducer

Sugar Free / Alcohol Free

Aspirin Free

CHERRY FLAVOR

**Drug Facts** 

# **Active ingredient**

(in each 5 mL teaspoonful)

Acetaminophen, USP 160 mg .....

# **Purpose**

Pain reliever/fever reducer

### Uses

- reduces fever
- temporarily relieves minor aches and pains due to:
  - Headache
  - muscular aches
  - backache
  - arthritis
  - the common cold
  - toothache
  - menstrual cramps
  - reduces fever

## Warnings

**Liver Warning:**This product contains acetaminophen.

Severe liver damage may occur if you take:

- more than 8 teaspoonfuls (40 mL) in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen

3 or more alcoholic drinks every day while using this product

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.

- for more than 10 days for pain unless directed by a doctor
- for more than 3 days for fever unless directed by a doctor
- if you are allergic to acetaminophen or any of the inactive ingredients in this product.

#### Ask a doctor before use

if you have health issues especially liver disease.

# Ask a doctor or pharmacist before use

if you are taking other drugs, including the blood thinner warfarin.

# When using this product do no exceed recommended dose (see Overdose warning)

# Stop use and ask a doctor if

- new symptoms occur such as rash, hives, itching or hoarseness
- redness or swelling is present
- pain gets worse or lasts for more than 10 days
- fever gets worse or lasts for more than 3 days
- symptoms do not improve

These could be signs of a serious condition.

# If pregnant or breast-feeding,

ask a health professional before use.

# Keep out of reach of children. Overdose warning

Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical even if you do not notice any signs or symptoms.

## **Directions**

# Do not exceed recommended dosage.

- adults and children 12 years of age and older: Take 3 teaspoonfuls (15 mL) every 5 hours; do not exceed 15 teaspoonfuls (75 mL) in 24 hours
- **children under 12 years of age:**Under the direct guidance of a licensed professional, doctor, or pharmacist.

## Other Information

• each 5 mL contains: Sodium 3 mL

If dispensed, dispense in a tight, light resistant cotainer with a child-resistant cap. Store at 20°C to 25°C (68°F to 77°F), excursions permitted between 15°C and 30°C (between 59°F and 86°F).

# **Inactive Ingredients**

Anhydrous citric acid, FD&C Blue #1, FD&C Red #40, flavor, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sodium benzoate, sorbitol solution.

## **Questions?**

Any adverse reactions contact FDA (888) 463-6332 or www.FDA.gov/medwatch

Distributed by:

VERITYRX, LLC

20225 NE 16th Place

Miami, FL 33179

## PRINCIPAL DISPLAY PANEL

M-PAP

Acetaminophen Liquid

160 mg/5 mL

Cherry Flavor



# M-PAP

acetaminophen liquid

D	ro	d	11/	+	In	fo	rm	ati	ion

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83939-0011(NDC:58657-524)
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Route of Administration ORAL

# **Active Ingredient/Active Moiety**

Ingredient Name	<b>Basis of Strength</b>	Strength
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	160 mg in 5 mL

ACETAMINOPHEN (UNII: 302091119D) (ACETAMINOPHEN - UNII: 302091119D) ACETAMINOPHEN

Inactive Ingredients				
Ingredient Name	Strength			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
WATER (UNII: 059QF0KO0R)				
SODIUM BENZOATE (UNII: 0 245FE5EU)				

SACCHARIN SODIUM (UNII: SB8ZUX40TY)

SORBITOL (UNII: 506T60A25R)

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

Product Characteristics				
Color		Score		
Shape		Size		
Flavor	CHERRY	Imprint Code		
Contains				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:83939- 0011-1	5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product	11/11/2024		
2	NDC:83939- 0011-2	10 in 1 BOX, UNIT-DOSE	11/10/2025		
2		5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			
3	NDC:83939- 0011-3	40 in 1 BOX, UNIT-DOSE	11/10/2025		
3		5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			
4	NDC:83939- 0011-4	50 in 1 BOX, UNIT-DOSE	11/11/2024		
4		5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			
5	NDC:83939- 0011-5	100 in 1 BOX, UNIT-DOSE	11/10/2025		
5		5 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	04/01/2024		

# Labeler - VERITYRX, LLC (097807737)

Revised: 11/2025 VERITYRX, LLC