

ALL-IN-ONE UTI EMERGENCY KIT- methenamine, sodium salicylate, and phenazopyridine hydrochloride
Bonafide Health, LLC dba Uqora

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Uqora All-In-One UTI Emergency Kit

Drug Facts

Active ingredients (in each tablet)

Methenamine 162mg

Sodium Salicylate 162.5mg (NSAID)*

*nonsteroidal anti-inflammatory drug

Purpose

Antibacterial

Analgesic (pain reliever)

Active ingredient (in each tablet)

Phenazopyridine Hydrochloride 99.5mg

Purpose

Urinary Tract Analgesic

Uses

UTI Infection Control

Temporarily relieves:

- pain and burning
- frequency and urgency of urination

UTI Pain Relief

Fast relief from urinary pain, burning, urgency and frequency associated with urinary tract infections. Treatment should not exceed 2 days; see Directions.

Warnings

Reye's Syndrome (UTI Infection Control Only)

Reye's syndrome:Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's syndrome, a rare but serious illness.

Stomach Bleeding Warning (UTI Infection Control Only)

Stomach bleeding warning:This product contains a nonsteroidal anti-inflammatory drug (NSAID), which may cause stomach bleeding. The chance is higher if you

- are 60 or older

- have stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing an NSAID (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

Do Not Exceed (*UTI Pain Relief Only*)

Do not exceed recommended dosage

Do Not Use

Do not use:

- If you are allergic to salicylates (including aspirin) (***UTI Infection Control only***)
- If you have Glucose-6-Phosphate Dehydrogenase (G6PD) deficiency unless approved by your doctor (***UTI Pain Relief only***)

Ask A Doctor Before Use

Ask a doctor before use if you have:

- frequent, burning urination for the first time (***UTI Infection Control only***)
- the stomach bleeding warning applying to you (***UTI Infection Control only***)
- history of stomach problems, such as heartburn (***UTI Infection Control only***)
- high blood pressure (***UTI Infection Control only***)
- heart disease (***UTI Infection Control only***)
- liver cirrhosis (***UTI Infection Control only***)
- bleeding problems (***UTI Infection Control only***)
- diuretic use (***UTI Infection Control only***)
- ulcers (***UTI Infection Control only***)
- kidney disease
- a sodium restricted diet (***UTI Infection Control only***)
- reached age 60 or older (***UTI Infection Control only***)
- allergies to foods, preservatives or dyes (***UTI Pain Relief only***)
- had a hypersensitive reaction to phenazopyridine (***UTI Pain Relief only***)

Ask A Doctor Or Pharmacist Before Use (*UTI Infection Control Only*)

Ask a doctor or pharmacist before use if you are:

- taking any other drug containing an NSAID (prescription or non-prescription)
- taking a blood thinning (anticoagulant), steroid, diabetes, gout or arthritis drug

When Using This Product

When using this product:

- do not take more than the recommended dosage (***UTI Infection Control only***)
- stomach upset may occur, taking this product with or after meals may reduce stomach upset (***UTI Pain Relief only***)
- your urine will become reddish-orange in color. This is not harmful, but care should be taken to avoid staining clothing or other items. (***UTI Pain Relief only***)

Stop Use And Ask A Doctor

Stop use and ask a doctor if:

- product has been used for 3 days (**UTI Infection Control only**)
- ringing in the ears (**UTI Infection Control only**)
- you experience any of the following signs of stomach bleeding: ■ feel faint ■ vomit blood ■ have bloody or black stool ■ have stomach pain or upset that gets worse or lasts (**UTI Infection Control only**)
- your symptoms last for more than 2 days (**UTI Pain Relief only**)
- you suspect you are having an adverse reaction to the medication (**UTI Pain Relief only**)

Long Term Administration (UTI Pain Relief Only)

Long-term administration of phenazopyridine hydrochloride has induced neoplasia in rats (large intestine) and mice (liver). Although no association between phenazopyridine hydrochloride and human neoplasia has been reported, adequate epidemiological studies along these lines have not been conducted.

If Pregnant or Breast Feeding

UTI Infection Control

If pregnant or breast feeding,ask a health professional before use. It is especially important not to use this product (which contains sodium salicylate) during the last 3 months of pregnancy unless definitely directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

UTI Pain Relief

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

Keep Out Of The Reach Of Children

Keep out of the reach of children.In case of overdose, get medical help or contact a Poison Control Center right away 1-800-222-1222

Directions**UTI Infection Control**

Adults and children 12 years and over:

- take 2 tablets with a full glass of water 3 times a day. Drink plenty of fluids.
- **Do not use for more than a 3 day period unless directed by a doctor.**

Children under 12 years: ask a doctor.

UTI Pain Relief

Adults and children 12 years and over:

- take 2 tablets 3 times daily with a full glass of water, with or after meals as needed
- **Do not use for more than 2 days (12 tablets) without consulting a doctor.**

Children under 12 years:consult a doctor

Other Information

UTI Infection Control

- **each tablet contains:**sodium 24.4 mg
- store at 59-86°F (15-30°C) in a cool dry place
- protect from sunlight
- Tamper evident: tablets sealed in blisters. Do not use if blister foil or seal is open or damaged

UTI Pain Relief

- this product may stain contact lenses
- this product can interfere with laboratory tests including urine, glucose (sugar), and ketones tests
- store at room temperature 15°-30°C (59°-86°F) in a dry place and protect from light
- Tamper evident: tablets sealed in blisters. Do not use if blister foil or seal is open or damaged

Inactive Ingredients

UTI Infection Control

benzoic acid, croscarmellose sodium, fd&c red #40, fd&c yellow #6, hypromellose, magnesium stearate, methacrylic acid-ethyl acrylate copolymer, microcrystalline cellulose, silicon dioxide, stearic acid, sodium bicarbonate, sodium lauryl sulfate, talc, titanium dioxide, triacetin, triethyl citrate

UTI Pain Relief

corn starch, croscarmellose sodium, hypromellose, lactose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinylpyrrolidone, pregelatinized starch, silicon dioxide, sodium starch glycolate, talc and triacetin.

Distributed by Uqora, Inc.

3043 4th Avenue, San Diego, CA 92103

support@uqora.com

For questions or concerns please contact (888) 313-1372

Additional Items

2 diagnostic UTI test strips(Directions Enclosed)

Other information

- store at room temperature 59-86 F (15-30 C) out of direct sunlight. Do not

refrigerate or freeze

- For in-vitro (external) diagnostic use.

Principal Display Panel

uqora

Urinary Care

Maximum Strength UTI Pain Relief Included

ALL-IN-ONE

UTI EMERGENCY KIT

2 Day Supply Includes:

Test Strips

- Test for a possible UTI at home in 2 minutes
- 2 Test Strips

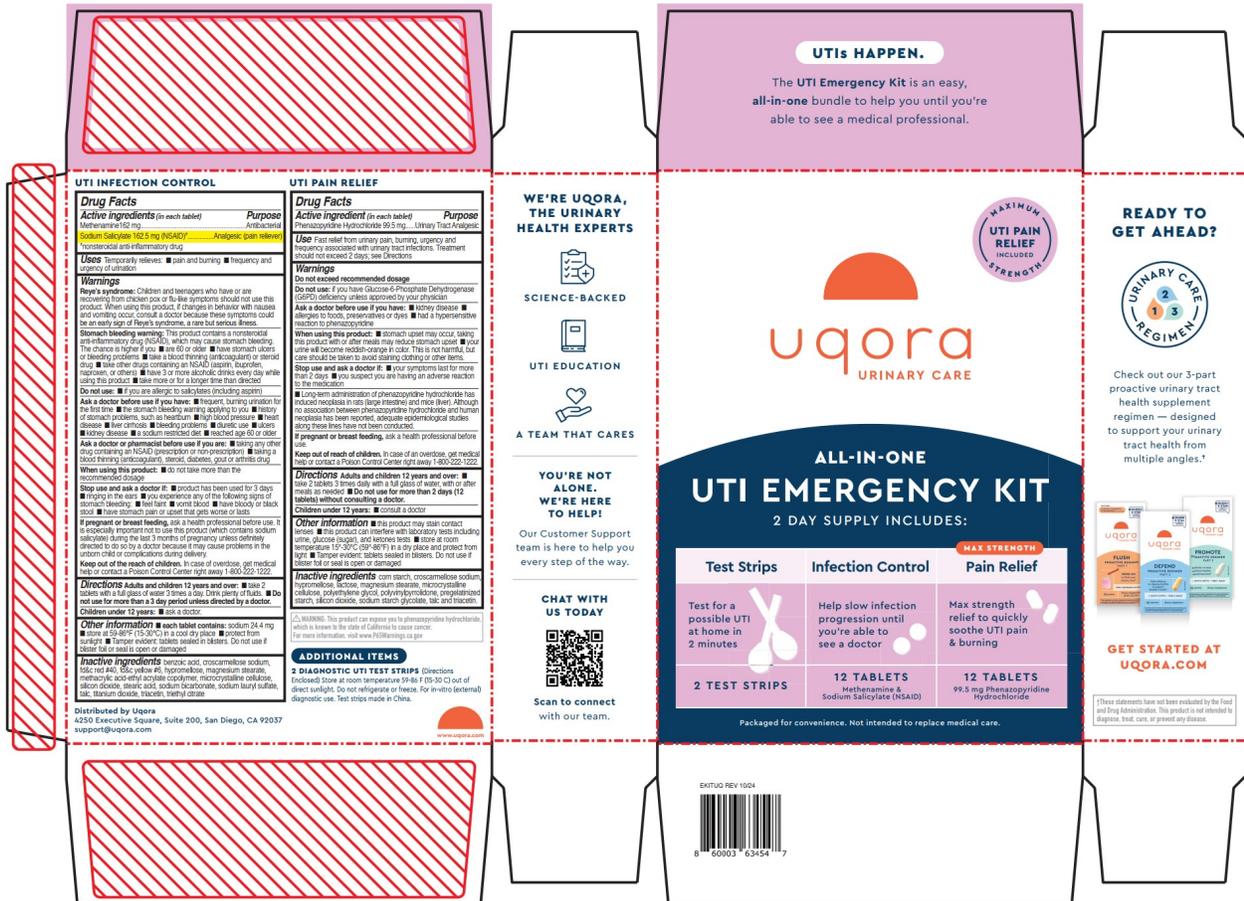
Infection Control

- Helps slow infection progression until you're able to see a doctor
- 12 Tablets (Methenamine & Sodium Salicylate (NSAID))

Max Strength Pain Relief

- Max strength relief to quickly soothe UTI pain & burning
- 12 Tablets (99.5mg Phenazopyridine Hydrochloride)

Packaged for convenience. Not intended to replace medical care.



ALL-IN-ONE UTI EMERGENCY KIT

methenamine, sodium salicylate, and phenazopyridine hydrochloride kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73712-501
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73712-501-24	1 in 1 CARTON; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)	10/17/2022	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BLISTER PACK	12
Part 2	1 BLISTER PACK	12

Part 1 of 2

UQORA ANTIBACTERIAL PLUS URINARY PAIN RELIEF

methenamine, sodium salicylate tablet

Product Information

Item Code (Source) NDC:73712-804

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
METHENAMINE (UNII: J50OIX95QV) (METHENAMINE - UNII:J50OIX95QV)	METHENAMINE	162 mg
SODIUM SALICYLATE (UNII: WQ1H85SYP) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SODIUM SALICYLATE	162.5 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHACRYLIC ACID-ETHYL ACRYLATE COPOLYMER (1:1) TYPE A (UNII: NX76LV5T8J)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	11mm
Flavor		Imprint Code	PH061
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73712-	12 in 1 BLISTER PACK; Type 9: Other Type of Part 3 Combination		

804-12	Product (e.g., Drug/Device/Biological Product)		
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/17/2022	

Part 2 of 2

UQORA MAXIMUM UTI PAIN RELIEF

phenazopyridine hydrochloride tablet

Product Information

Item Code (Source)	NDC:73712-114
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENAZOPYRIDINE HYDROCHLORIDE (UNII: 0EWG668W17) (PHENAZOPYRIDINE - UNII:K2J09EMJ52)	PHENAZOPYRIDINE HYDROCHLORIDE	99.5 mg

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6130)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TALC (UNII: 7SEV7J4R1U)	
TRIACETIN (UNII: XHX3C3X673)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	brown	Score	no score
Shape	OVAL	Size	9mm
Flavor		Imprint Code	p99
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73712-114-12	1 in 1 CARTON		
1		12 in 1 BLISTER PACK; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/17/2022	

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/17/2022	

Labeler - Bonafide Health, LLC dba Uqora (118327455)**Establishment**

Name	Address	ID/FEI	Business Operations
Pharbest		557054835	manufacture(73712-501)

Revised: 11/2025

Bonafide Health, LLC dba Uqora