UQORA MAXIMUM UTI PAIN RELIEF AND FLUSH CONVENIENCE BUNDLE-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 Maximum UTI Pain Relief + Flush Convenience Bundle

Uqora Maximum UTI Pain Relief

Drug Facts

Active ingredient (in each tablet)

Phenazopyridine Hydrochloride 99.5 mg

Purpose

Active ingredient (in each tablet)	***Purpose***
Phenazopyridine Hydrochloride 99.5mg	Urinary Tract Analgesic

Use

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

Warnings

Do not exceed recommended dosage

Do Not Use

Do not use:
- If you have Glucose-6-Phosphate Dehydrogenase (G6PD) deficiency unless approved by your physician

Ask A Doctor Before Use

Ask a doctor before use if you have:
- kidney disease
- allergies to foods, preservatives or dyes
- had a hypersensitive reaction to phenazopyridine

When Using This Product

When using this product:
- stomach upset may occur, taking this product with or after meals may reduce stomach upset

- your urine will become reddish-orange in color. This is not harmful, but care should be taken to avoid staining clothing or other items.

Stop Use And Ask A Doctor

Stop use and ask a doctor if:
- your symptoms last for more than 2 days
- you suspect you are having an adverse reaction to the medication

Long Term Administration

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

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

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
- 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

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

Additional Items

FLUSH Urinary Tract Drink Mix

2 packets

Dietary Supplement

Principal Display Panel

uqora

Founded by a UTI Sufferer

Get Relief

Get Ahead

Convenience Bundle Pack

UTI Pain Relief Max Strength*

Fast targeted relief for pain, burning & urgency

2 Day Supply

12 tablets

99.5 mg Phenazopyridine Hydrochloride

Packaged for convenience.

Not intended to replace medical care.

*Most powerful dose without an Rx.

&

Flush Urinary Tract Drink Mix

Clinical Strength

D-Mannose per serving

Flushes your urinary tract + immune support*

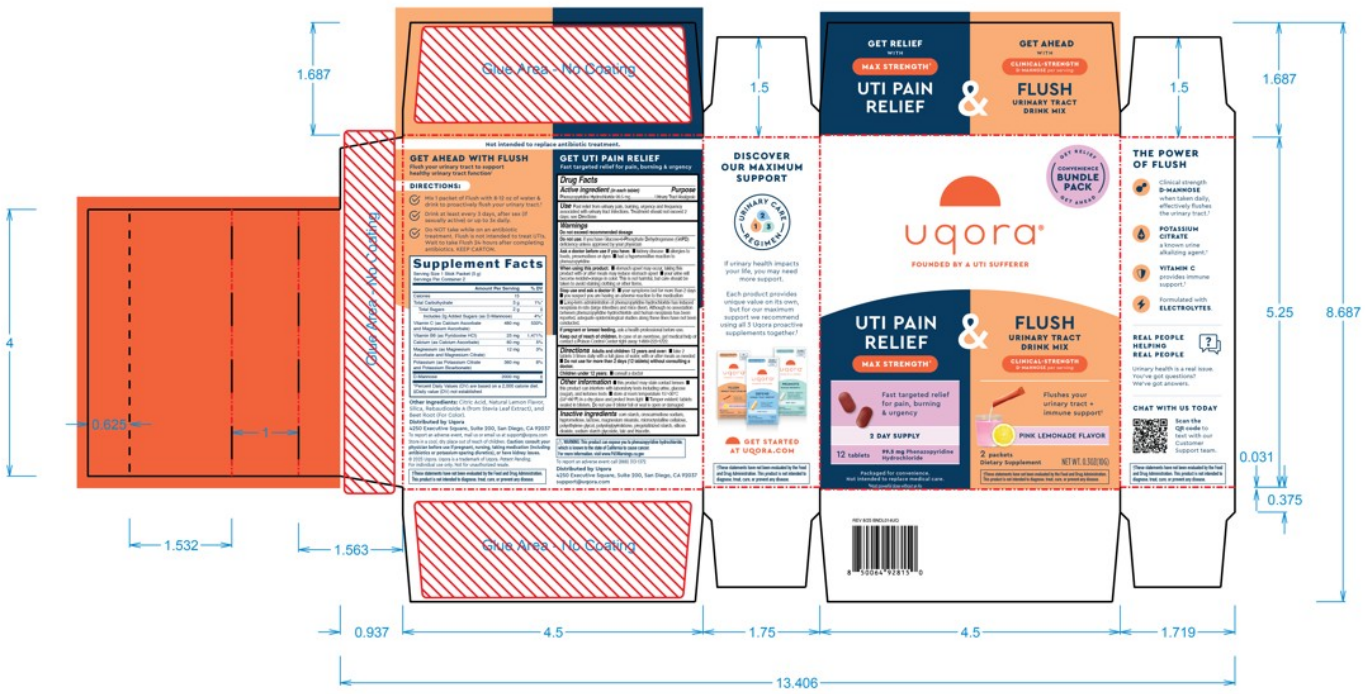
Pink Lemonade Flavor

2 packets

Dietary Supplement

Net Wt. 0.3OZ(10G

* These statements have not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure, or prevent any disease.



UQORA MAXIMUM UTI PAIN RELIEF AND FLUSH CONVENIENCE BUNDLE

phenazopyridine hydrochloride kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73712-600

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73712-600-12	1 in 1 KIT; Type 0: Not a Combination Product	11/10/2025	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 BLISTER PACK	12
Part 2	1 PACKET	2

Part 1 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: 3WQ0SDW1A)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
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)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

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	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
unapproved drug other		11/10/2025	

Part 2 of 2

FLUSH

d-mannose, potassium, magnesium calcium, vitamin b6, vitamin c powder

Product Information

Route of Administration	ORAL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
DIETARY SUPPLEMENT		11/10/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/10/2025	

Labeler - Bonafide Health, LLC dba Uqora (118327455)

Establishment

Name	Address	ID/FEI	Business Operations
Pharbest		557054835	manufacture(73712-600, 73712-114)