SIGNATURE CARE MAXIMUM STRENGTH URINARY PAIN RELIEFphenazopyridine hydrochloride tablet Safeway

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.

DRUG FACTS

Active ingredient (in each tablet)

Phenazopyridine Hydrochloride 99.5 mg.

Purpose

Urinary Analgesic

Warnings

Do not exceed recommended dosage

Ask doctor before use if you have

- kidney disease
- allergies to food, preservatives or dyes
- had a hypersensitive reaction to phenazopyridine

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 doctor if

- your symptoms last for more than 2 days
- you suspect you are having an adverse reaction to the medication

If pregnant or breast feeding,

Ask a health professional before use.

Keep out of reach of children

In case of an overdose, get medical help or contact a Poison Control Center right away.

Use

Fast relief from urinary pain, burning, urgency and frequency associated with urinary tract infections.

Inactive ingredients

Lactose, magnesium silicate, magnesium stearate, microcrystalline cellulose, pharmaceutical glaze, and sodium starch glycolate.

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
- children under 12 years: consult a doctor
- Do not use for more than 2 days (12 tablets) without consulting a doctor



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phenazopyridine hydrochloride tablet

Product Information

Product Type HUMAN OTC DRUG **Item Code (Source)** NDC:21130-792

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength

PHENAZOPYRIDINE HYDROCHLORIDE (UNII: 0EWG668W17) **PHENAZ OPYRIDINE** 99.5 mg **HYDROCHLORIDE**

(PHENAZ OPYRIDINE - UNII: K2J09EMJ52)

Inactive Ingredients			
Ingredient Name	Strength		
LACTOSE (UNII: J2B2A4N98G)			

MAGNESIUM SILICATE (UNII: 9B9691B2N9)

Product Characteristics

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

F	Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:21130- 792-24	1 in 1 CARTON	12/23/2019		
1		24 in 1 BLISTER PACK; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/23/2019	

Labeler - Safeway (009137209)

Registrant - Reese Pharmaceutical Co (004172052)

Revised: 12/2022 Safeway