

**DOLLAR GENERAL MAXIMUM STRENGTH URINARY PAIN RELIEF-
phenazopyridine hydrochloride tablet
Dolgencorp, Inc.**

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



DOLLAR GENERAL MAXIMUM STRENGTH URINARY PAIN RELIEF

phenazopyridine hydrochloride tablet

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:55910-631 |
| 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 |
|--|----------|
| LACTOSE (UNII: J2B2A4N98G) | |
| MAGNESIUM SILICATE (UNII: 9B9691B2N9) | |

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:55910-631-12 | 1 in 1 CARTON | 10/03/2019 | |
| 1 | | 12 in 1 BLISTER PACK; Type 0: Not a Combination Product | | |
| 2 | NDC:55910-631-24 | 1 in 1 CARTON | 03/12/2020 | |
| 2 | | 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 | | 10/03/2019 | |

Labeler - Dolgencorp, Inc. (068331990)

Registrant - Reese Pharmaceutical Co (004172052)

Revised: 12/2022

Dolgencorp, Inc.