

WALGREENS URINARY PAIN RELIEF MAXIMUM STRENGTH- phenazopyridine hydrochloride tablet

Walgreens Co.

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

PHENAZOPYRIDINE HYDROCHLORIDE 97.5 MG

PHENAZOPYRIDINE HYDROCHLORIDE URINARY ANALGESIC

DO NOT EXCEED RECOMMENDED DOSAGE

ASK DOCTOR BEFORE USE

IF YOU HAVE KIDNEY DISEASE

ALLERGIES TO FOODS,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 A DOCTOR

IF YOUR SYMPTOMS LAST FOR MORE THAN 2 DAYS

YOU SUSPECT YOU ARE HAVING AN ADVERSE REACTION TO THE MEDICATION

ASK A HEALTH PROFESSIONAL BEFORE USE

IN CASE OF 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.

Lactose, magnesium silicate, magnesium stearate, microcrystalline cellulose,

pharmaceutical glaze, and sodium starch glycolate.

Adults and Children 12 years of age and over; take 2 tablets 3 times daily with a full glass of water,with or after meals as needed.

Children under 12 years of age; consult a doctor.

Do not use for more than 2 days (12 tablets) without consulting a Doctor.



WALGREENS URINARY PAIN RELIEF MAXIMUM STRENGTH

phenazopyridine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0113
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENAZOPYRIDINE HYDROCHLORIDE (UNII: 0EWG668W17) (PHENAZOPYRIDINE - UNII:K2J09EMJ52)	PHENAZOPYRIDINE HYDROCHLORIDE	97.5 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	
MAGNESIUM SILICATE (UNII: 9B9691B2N9)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	975
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0113-01	1 in 1 CARTON	04/05/2010	
1	NDC:0363-0113-12	12 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		04/05/2010	

Labeler - Walgreens Co. (008965063)**Registrant** - Reese Pharmaceutical Co (004172052)**Establishment**

Name	Address	ID/FEI	Business Operations
Reese Pharmaceutical Co		004172052	relabel(0363-0113) , repack(0363-0113) , manufacture(0363-0113)

Revised: 1/2019

Walgreens Co.