

**PHENAZOPYRIDINE HYDROCHLORIDE- phenazopyridine hydrochloride tablet, coated
DirectRX**

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.

PHENAZOPYRIDINE HYDROCHLORIDE

SPL Unclassified

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Description

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Clinical Pharmacology

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Indications and Usage

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Contraindications

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Adverse Reactions

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Precautions

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Dosage and Administration

Overdosage

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Package Label

Mfg For: Neulonwide Labs, LLC
 Berlin, NJ 08630
 NDC 42937-702-10

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PHENAZOPYRIDINE HYDROCHLORIDE
200mg 10 Tabs

Generic For: **PYRIDIUM**
 Each tablet contains: Phenazopyridine Hydrochloride, USP 200mg

Lot# Discard After: 07/17
 Prod# 098-10

Packaged and Distributed By: **DIRECT Rx**
 Alpharetta, GA 30005

AEG4K
 Caution: Federal law prohibits transfer of this drug to any person other than the patient for whom it was prescribed.
RX ONLY-KEEP OUT OF REACH OF CHILDREN
 Dosage: See package insert. Store between 68-77 degrees F.

NDC 61919-098-10

Medication should be taken with plenty of water. & May cause Discoloration of urine or feces

PHENAZOPYRIDINE HYDROCHLORIDE
 NDC 61919-098-10 10 Ta
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PHENAZOPYRIDINE HYDROCHLORIDE

phenazopyridine hydrochloride tablet, coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61919-098(NDC:42937-702)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENAZOPYRIDINE HYDROCHLORIDE (UNII: 0EWG668W17) (PHENAZOPYRIDINE - UNII:K2J09EMJ52)	PHENAZOPYRIDINE HYDROCHLORIDE	200 mg

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POVIDONES (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	brown (Reddish-brown)	Score	no score
Shape	ROUND	Size	10mm
Flavor		Imprint Code	702
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61919-098-10	10 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		12/09/2015	

Labeler - DirectRX (079254320)**Establishment**

Name	Address	ID/FEI	Business Operations
DirectRX		079254320	repack(61919-098)

Revised: 12/2015

DirectRX