

GLYBURIDE- glyburide tablet

DirectRX

GLYBURIDE

Description

-

Clinical Pharmacology

-

Indications and Usage

-

Contraindications

-

Precautions

-

Adverse Reactions

-

Overdosage

-

Dosage and Administration

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Package Label

D

Mfg For: Heritage Pharmaceuticals Inc.
Edison, NJ 07724
NDC 23155-058-01

**GLYBURIDE
5mg 60 Tabs**

Generic For: **MICRONASE**
Each tablet contains 5mg glyburide

Lot# Discard After: 01/20
Prod# 378-60

Alpharetta, GA 30005

AEER9
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

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NDC 61919-378-60

GLYBURIDE 5mg
NDC 61919-378-60 60 Tabs
Lot Exp Date 01/20
Mfg NDC 23155-058-01

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Mfg Lot: 12/3/2015

Packaged and Distributed By: **DIRECTRX**

GLYBURIDE

glyburide tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61919-378(NDC:23155-058)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYBURIDE (UNII: SX6K58TVWC) (GLYBURIDE - UNII: SX6K58TVWC)	GLYBURIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	

Product Characteristics

Color	blue	Score	2 pieces
Shape	CAPSULE	Size	9mm
Flavor		Imprint Code	I37
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61919-378-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	12/03/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090937	12/03/2015	

Labeler - DirectRX (079254320)

Establishment

Name	Address	ID/FEI	Business Operations
DirectRX		079254320	repack(61919-378)