GLYBURIDE- glyburide tablet
DirectRX

GLYBURIDE

Description

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Clinical Pharmacology

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Indications and Usage

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Contraindications

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Precautions

-

Adverse Reactions

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Overdosage

-

Dosage and Administration

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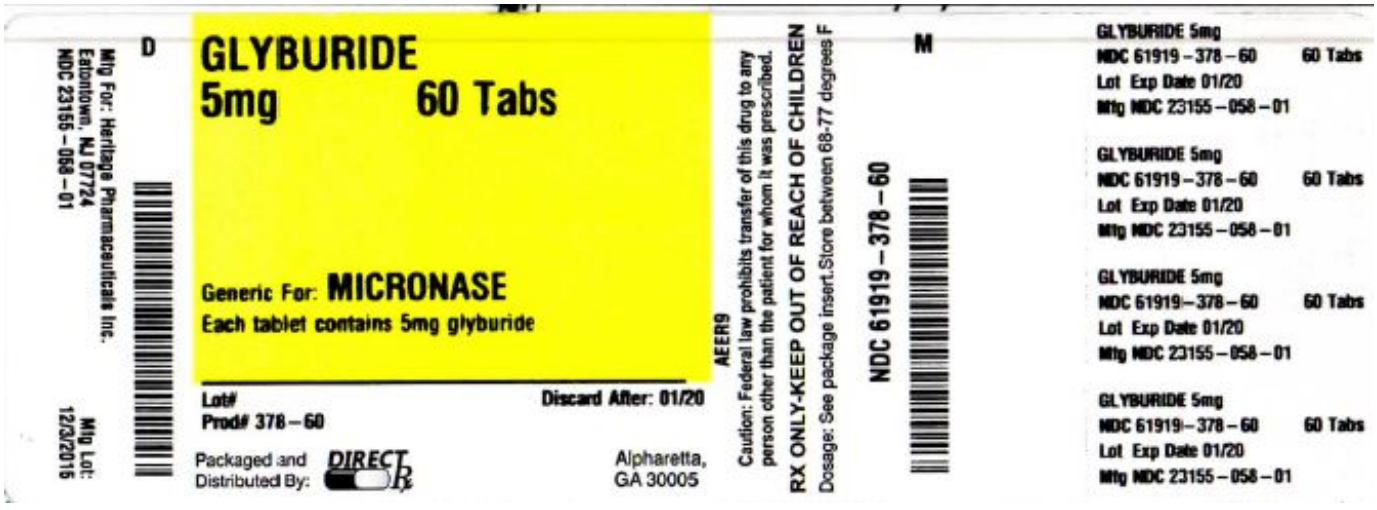
Package Label



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: H3R47K3TBD)	

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)