

**LOSARTAN POTASSIUM- losartan potassium tablet, film coated**  
**Zydus Lifesciences Limited**

**LOSARTAN POTASSIUM TABLETS**

**PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

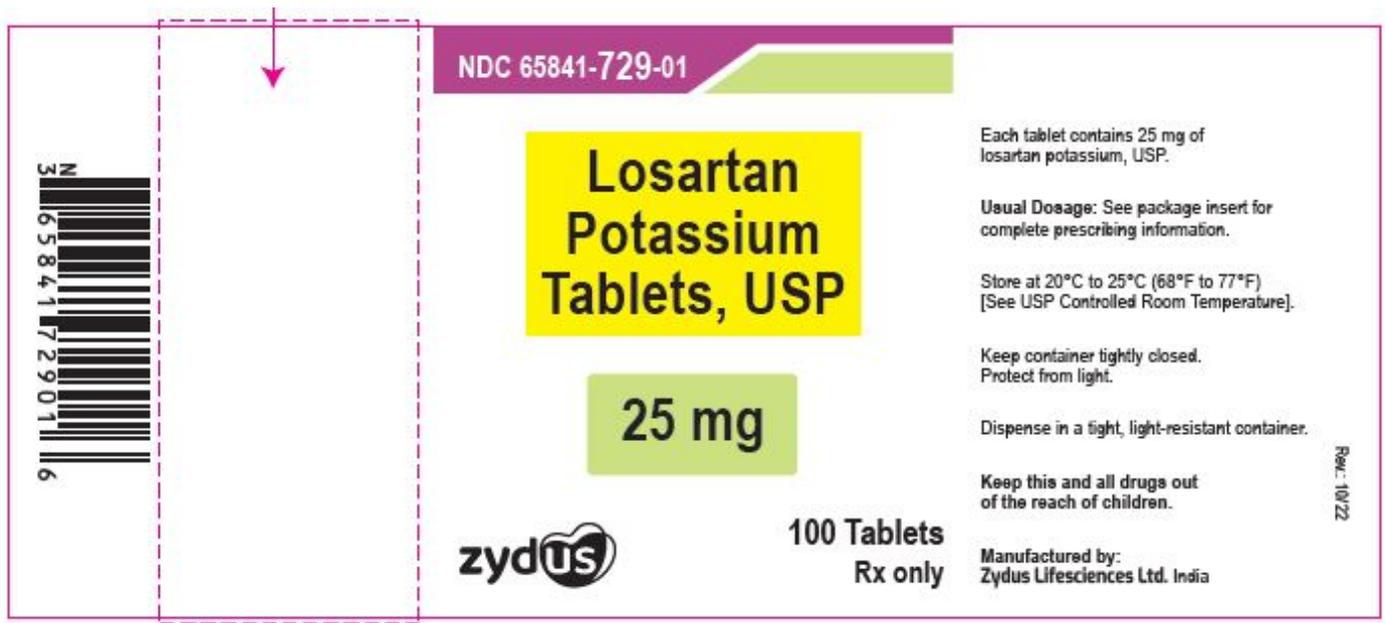
NDC 65841-729-01

Losartan Potassium Tablets USP, 25 mg

Rx Only

100 Tablets

Zydus



NDC 65841-730-01

Losartan Potassium Tablets USP, 50 mg

Rx Only

100 Tablets

Zydus

NDC 65841-730-01



**Losartan Potassium Tablets, USP**

**50 mg**

**zydus**

**100 Tablets**  
Rx only

Each tablet contains 50 mg of losartan potassium, USP.

**Usual Dosage:** See package insert for complete prescribing information.

Store at 20°C to 25°C (68°F to 77°F)  
[See USP Controlled Room Temperature].

Keep container tightly closed.  
Protect from light.

Dispense in a tight, light-resistant container.

Keep this and all drugs out of the reach of children.

Manufactured by:  
Zydus Lifesciences Ltd. India

Rev: 10/22

NDC 65841-731-01

Losartan Potassium Tablets USP, 100 mg

Rx Only

100 Tablets

Zydus

NDC 65841-731-01



**Losartan Potassium Tablets, USP**

**100 mg**

**zydus**

**100 Tablets**  
Rx only

Each tablet contains 100 mg of losartan potassium, USP.

**Usual Dosage:** See package insert for complete prescribing information.

Store at 20°C to 25°C (68°F to 77°F)  
[See USP Controlled Room Temperature].

Keep container tightly closed.  
Protect from light.

Dispense in a tight, light-resistant container.

Keep this and all drugs out of the reach of children.

Manufactured by:  
Zydus Lifesciences Ltd. India

Rev: 10/22

## LOSARTAN POTASSIUM

losartan potassium tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:65841-729
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LOSARTAN POTASSIUM</b> (UNII: 3ST302B24A) (LOSARTAN - UNII:JMS50MPO89)	LOSARTAN POTASSIUM	25 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>HYDROXYPROPYL CELLULOSE, LOW SUBSTITUTED</b> (UNII: 2165RE0K14)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	

### Product Characteristics

<b>Color</b>	WHITE (WHITE TO OFF-WHITE)	<b>Score</b>	no score
<b>Shape</b>	CAPSULE (CAPSULE)	<b>Size</b>	8mm
<b>Flavor</b>		<b>Imprint Code</b>	Z;2
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-729-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
2	NDC:65841-729-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
3	NDC:65841-729-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
4	NDC:65841-729-24	10000 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
5	NDC:65841-729-06	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078243	10/04/2010	

# LOSARTAN POTASSIUM

losartan potassium tablet, film coated

## Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:65841-730
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LOSARTAN POTASSIUM</b> (UNII: 3ST302B24A) (LOSARTAN - UNII:JMS50MPO89)	LOSARTAN POTASSIUM	50 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>HYDROXYPROPYL CELLULOSE, LOW SUBSTITUTED</b> (UNII: 2165RE0K14)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	

## Product Characteristics

<b>Color</b>	WHITE (WHITE TO OFF-WHITE)	<b>Score</b>	2 pieces
<b>Shape</b>	CAPSULE (CAPSULE)	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	Z16
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65841-730-06	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
2	NDC:65841-730-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
3	NDC:65841-730-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
4	NDC:65841-730-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
5	NDC:65841-730-	10000 in 1 BOTTLE; Type 0: Not a Combination	10/04/2010	

24	Product	10/04/2010	
<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078243	10/04/2010	

## LOSARTAN POTASSIUM

losartan potassium tablet, film coated

<b>Product Information</b>			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:65841-731
Route of Administration	ORAL		

<b>Active Ingredient/Active Moiety</b>			
Ingredient Name	Basis of Strength	Strength	
LOSARTAN POTASSIUM (UNII: 3ST302B24A) (LOSARTAN - UNII:JMS50MPO89)	LOSARTAN POTASSIUM	100 mg	

<b>Inactive Ingredients</b>		
Ingredient Name	Strength	
HYPROMELLOSES (UNII: 3NXW29V3WO)		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)		
HYDROXYPROPYL CELLULOSE, LOW SUBSTITUTED (UNII: 2165RE0K14)		
STARCH, CORN (UNII: O8232NY3SJ)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		

<b>Product Characteristics</b>			
Color	WHITE (WHITE TO OFF-WHITE)	Score	no score
Shape	CAPSULE (CAPSULE)	Size	12mm
Flavor		Imprint Code	Z18
Contains			

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:65841-731-06	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
2	NDC:65841-731-16	90 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
3	NDC:65841-731-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
4	NDC:65841-731-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	10/04/2010	
5	NDC:65841-731-77	10 in 1 CARTON	10/04/2010	
5	NDC:65841-731-30	10 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
ANDA	ANDA078243	10/04/2010	

**Labeler** - Zydus Lifesciences Limited (918596198)

**Registrant** - Zydus Lifesciences Limited (918596198)

## Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		677605858	ANALYSIS(65841-729, 65841-730, 65841-731) , MANUFACTURE(65841-729, 65841-730, 65841-731)

## Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		918596198	ANALYSIS(65841-729, 65841-730, 65841-731) , MANUFACTURE(65841-729, 65841-730, 65841-731)

## Establishment

Name	Address	ID/FEI	Business Operations
Zydus Pharmaceuticals Limited		650173735	MANUFACTURE(65841-729, 65841-730, 65841-731) , ANALYSIS(65841-729, 65841-730, 65841-731)

Revised: 1/2026

Zydus Lifesciences Limited