

**CHLORTHALIDONE - chlorthalidone tablet**  
**Zydus Lifesciences Limited**

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**Chlorthalidone Tablets, USP**

**PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

NDC 70771-1369-1

Chlorthalidone tablets USP ,25 mg

Rx only

100 tablets



NDC 70771-1370-1

Chlorthalidone tablets USP, 50 mg

Rx only

100 tablets

NDC 70771-1370-1



**Chlorthalidone  
Tablets, USP**

**50 mg**

**zydus**

**100 Tablets  
Rx only**

Each tablet contains  
Chlorthalidone, USP.....50 mg

Usual Adult Dosage: See accompanying  
prescribing information.

Store at 20° to 25°C (68° to 77°F) [see USP  
Controlled Room Temperature].  
Protect from light.

Dispense in a tight, light-resistant  
container as defined in the USP  
using a child-resistant closure.

Keep container tightly closed.

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

Manufactured by:  
Zydus Lifesciences Ltd.  
Ahmedabad, India

Rev.: 08/22

## CHLORTHALIDONE

chlorthalidone tablet

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:70771-1369
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORTHALIDONE (UNII: Q0MQD1073Q) (CHLORTHALIDONE - UNII:Q0MQD1073Q)	CHLORTHALIDONE	25 mg

### Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

### Product Characteristics

<b>Color</b>	YELLOW (LIGHT YELLOW)	<b>Score</b>	no score
<b>Shape</b>	ROUND (ROUND)	<b>Size</b>	6mm
<b>Flavor</b>		<b>Imprint Code</b>	Z25
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1369-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2019	
2	NDC:70771-1369-0	1000 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2019	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207813	05/15/2019	

## CHLORTHALIDONE

chlorthalidone tablet

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1370
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORTHALIDONE (UNII: Q0MQD1073Q) (CHLORTHALIDONE - UNII:Q0MQD1073Q)	CHLORTHALIDONE	50 mg

### Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STARCH, CORN (UNII: O8232NY3S)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

### Product Characteristics

Color	WHITE (WHITE)	Score	2 pieces
Shape	ROUND (ROUND)	Size	7mm
Flavor		Imprint Code	Z;50
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:70771-1370-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2019	
2	NDC:70771-1370-0	1000 in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2019	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207813	05/15/2019	

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

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

## Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		863362789	ANALYSIS(70771-1369, 70771-1370) , MANUFACTURE(70771-1369, 70771-1370)

Revised: 8/2022

Zydus Lifesciences Limited