

CHLORTHALIDONE- chlorthalidone tablet
Zydus Lifesciences Limited

Chlorthalidone Tablets, USP

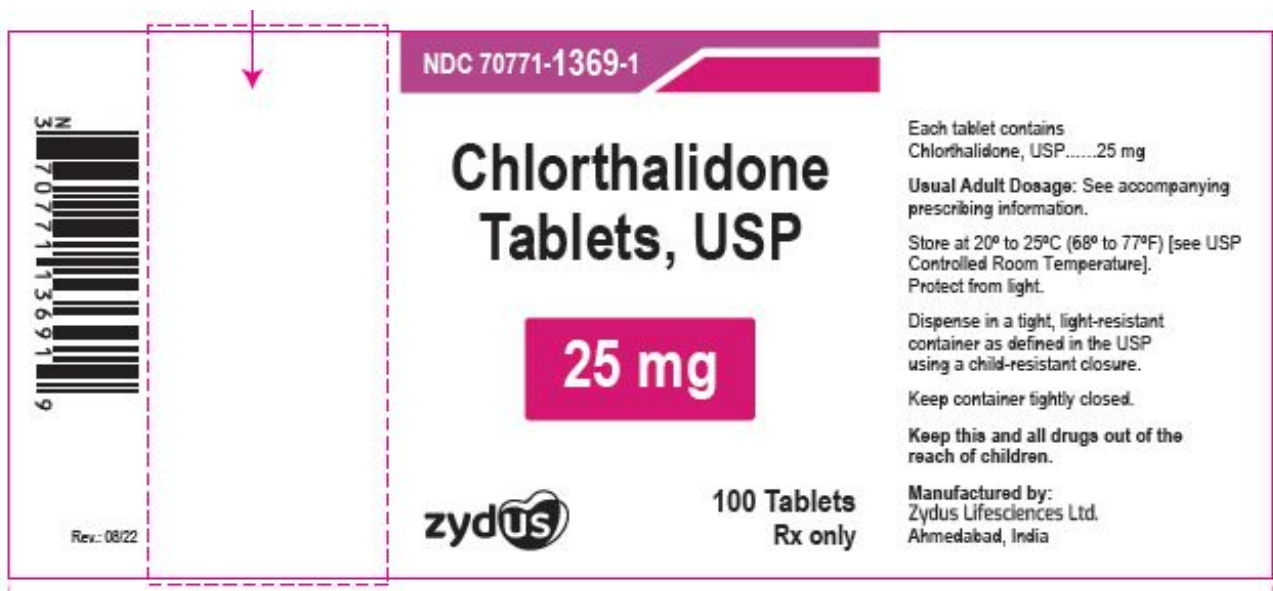
PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

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



Rev.: 08/22

Chlorthalidone Tablets, USP

50 mg

zydus

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

100 Tablets
Rx only

CHLORTHALIDONE

chlorthalidone tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1369
Route of Administration	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

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

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: 11/2024

Zydus Lifesciences Limited