

**DILTIAZEM HYDROCHLORIDE- diltiazem hydrochloride tablet**  
**Zydus Lifesciences Limited**

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**Diltiazem Hydrochloride Tablets, USP**  
**Rx only**

**SPL UNCLASSIFIED**

**PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

NDC 70771-1929-1 in bottles of 100 tablets

Diltiazem hydrochloride tablets USP, 30 mg

100 Tablets

Rx Only



Over Coding Template  
No Varnished Area (Do Not Print)  
(18 x 41 mm)

NDC 70771-1929-1

**Diltiazem  
Hydrochloride  
Tablets, USP**

**30 mg**

Diltiazem hydrochloride tablets may be  
swallowed whole, crushed, or chewed.  
Do not split diltiazem hydrochloride tablets.

**zydus**

**100 Tablets**  
**Rx only**

Each film coated tablet contains  
Diltiazem Hydrochloride, USP.....30 mg  
(equivalent to 27.6 mg diltiazem).  
**Usual dosage:** See package insert for full  
prescribing information.  
Dispense in a tight, light-resistant container as  
defined in the USP, with child-resistant closure.  
**This package is child-resistant.**  
**Keep this and all medicines out of  
the reach of children.**  
**Store at 25°C (77°F); excursions permitted  
to 15° to 30°C (59° to 86°F) [see USP  
Controlled Room Temperature]. Avoid  
excessive humidity.**  
Manufactured by: Zydus Lifesciences Ltd.,  
Baddi-173205, India

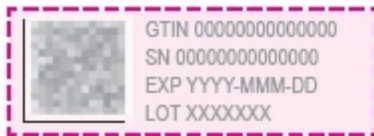
XXXXXXXX  
Rev.: 08/25

NDC 70771-1930-1 in bottles of 100 tablets

Diltiazem hydrochloride tablets USP, 60 mg

100 Tablets

Rx Only



Over Coding Template  
No Varnished Area (Do Not Print)  
(18 x 41 mm)

NDC 70771-1930-1

**Diltiazem  
Hydrochloride  
Tablets, USP**

**60 mg**

Diltiazem hydrochloride tablets may be  
swallowed whole, crushed, or chewed.

**zydus**

**100 Tablets  
Rx only**

Each film coated tablet contains  
Diltiazem Hydrochloride, USP..... 60 mg  
(equivalent to 55.1 mg diltiazem).  
**Usual dosage:** See package insert for full  
prescribing information.  
Dispense in a tight, light-resistant container as  
defined in the USP, with child-resistant closure.  
**This package is child-resistant.**  
**Keep this and all medicines out of  
the reach of children.**  
**Store at 25°C (77°F); excursions permitted  
to 15° to 30°C (59° to 86°F) [see USP  
Controlled Room Temperature]. Avoid  
excessive humidity.**  
**Manufactured by: Zydus Lifesciences Ltd.,  
Baddi-173205, India**

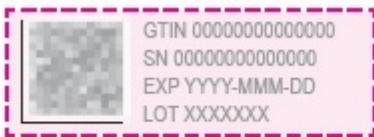
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Rev.: 08/25

NDC 70771-1931-1 in bottles of 100 tablets

Diltiazem hydrochloride tablets USP, 90 mg

100 Tablets

Rx Only



Over Coding Template  
No Varnished Area (Do Not Print)  
(18 x 41 mm)

NDC 70771-1931-1

# Diltiazem Hydrochloride Tablets, USP

**90 mg**

Diltiazem hydrochloride tablets may be swallowed whole, crushed, or chewed.

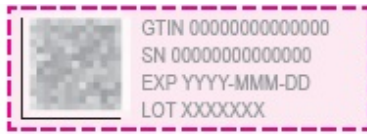
**zydus**

**100 Tablets**  
**Rx only**

Each film coated tablet contains Diltiazem Hydrochloride, USP.....90 mg (equivalent to 82.7 mg diltiazem).  
**Usual dosage:** See package insert for full prescribing information.  
Dispense in a tight, light-resistant container as defined in the USP, with child-resistant closure.  
**This package is child-resistant.**  
**Keep this and all medicines out of the reach of children.**  
**Store at 25°C (77°F); excursions permitted to 15° to 30°C (59° to 86°F) [see USP Controlled Room Temperature]. Avoid excessive humidity.**  
**Manufactured by: Zydus Lifesciences Ltd., Baddi-173205, India**

XXXXXXXX  
Rev.: 08/25

NDC 70771-1932-1 in bottles of 100 tablets  
Diltiazem hydrochloride tablets USP, 120 mg  
100 Tablets  
Rx Only



Over Coding Template  
No Varnished Area (Do Not Print)  
(18 x 50.5 mm)

XXXXXXXX  
Rev.: 08/25

NDC 70771-1932-1

# Diltiazem Hydrochloride Tablets, USP

120 mg

Diltiazem hydrochloride tablets may be  
swallowed whole, crushed, or chewed.

**100 Tablets**  
Rx only

Each film coated tablet contains  
Diltiazem Hydrochloride, USP.....120 mg  
(equivalent to 110.3 mg diltiazem).

**Usual dosage:** See package insert for full  
prescribing information.

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

**This package is child-resistant.**

**Keep this and all medicines out of  
the reach of children.**

**Store at 25°C (77°F); excursions permitted  
to 15° to 30°C (59° to 86°F) [see USP  
Controlled Room Temperature]. Avoid  
excessive humidity.**

**Manufactured by: Zydus Lifesciences Ltd.,  
Baddi-173205, India**

<b>DILTIAZEM HYDROCHLORIDE</b>			
diltiazem hydrochloride tablet			
Product Information			
<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:70771-1929
<b>Route of Administration</b>	ORAL		
Active Ingredient/Active Moiety			
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>	
<b>DILTIAZEM HYDROCHLORIDE</b> (UNII: OLH94387TE) (DILTIAZEM - UNII:EE92BBP03H)	DILTIAZEM HYDROCHLORIDE	30 mg	
Inactive Ingredients			
<b>Ingredient Name</b>	<b>Strength</b>		
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)			
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)			
<b>HYPROMELLOSE 2910 (5 MPA.S)</b> (UNII: R75537T0T4)			
<b>HYPROMELLOSE 2208 (100 MPA.S)</b> (UNII: B1QE5P712K)			

<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>HYPROMELLOSE 2910 (6 MPA.S)</b> (UNII: 0WZ8WG20P6)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>FD&amp;C BLUE NO. 1 ALUMINUM LAKE</b> (UNII: J9EQA3S2JM)	
<b>D&amp;C YELLOW NO. 10 ALUMINUM LAKE</b> (UNII: CQ3XH3DET6)	

### Product Characteristics

<b>Color</b>	GREEN	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	8mm
<b>Flavor</b>		<b>Imprint Code</b>	1867
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1929-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2026	
2	NDC:70771-1929-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2026	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA219253	05/25/2026	

## DILTIAZEM HYDROCHLORIDE

diltiazem hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:70771-1930
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DILTIAZEM HYDROCHLORIDE</b> (UNII: OLH94387TE) (DILTIAZEM - UNII:EE92BBP03H)	DILTIAZEM HYDROCHLORIDE	60 mg

### Inactive Ingredients

Ingredient Name	Strength
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<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>HYPROMELLOSE 2910 (5 MPA.S)</b> (UNII: R75537T0T4)	
<b>HYPROMELLOSE 2208 (100 MPA.S)</b> (UNII: B1QE5P712K)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>HYPROMELLOSE 2910 (6 MPA.S)</b> (UNII: 0WZ8WG20P6)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	

### Product Characteristics

<b>Color</b>	WHITE	<b>Score</b>	2 pieces
<b>Shape</b>	ROUND	<b>Size</b>	10mm
<b>Flavor</b>		<b>Imprint Code</b>	18;68
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1930-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2026	
2	NDC:70771-1930-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2026	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA219253	05/25/2026	

## DILTIAZEM HYDROCHLORIDE

diltiazem hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:70771-1931
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DILTIAZEM HYDROCHLORIDE</b> (UNII: OLH94387TE) (DILTIAZEM - UNII:EE92BBP03H)	DILTIAZEM HYDROCHLORIDE	90 mg

## Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)	
HYPROMELLOSE 2208 (100 MPA.S) (UNII: B1QE5P712K)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
TALC (UNII: 7SEV7J4R1U)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)	

## Product Characteristics

Color	GREEN	Score	2 pieces
Shape	OVAL (Capsule shaped)	Size	17mm
Flavor		Imprint Code	18;69
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1931-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2026	
2	NDC:70771-1931-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2026	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA219253	05/25/2026	

## DILTIAZEM HYDROCHLORIDE

diltiazem hydrochloride tablet

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1932
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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<b>DILTIAZEM HYDROCHLORIDE</b> (UNII: OLVH94387TE) (DILTIAZEM - UNII:EE92BBP03H)	DILTIAZEM HYDROCHLORIDE	120 mg
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### Inactive Ingredients

Ingredient Name	Strength
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>HYPROMELLOSE 2910 (5 MPA.S)</b> (UNII: R75537T0T4)	
<b>HYPROMELLOSE 2208 (100 MPA.S)</b> (UNII: B1QE5P712K)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>HYPROMELLOSE 2910 (6 MPA.S)</b> (UNII: 0WZ8WG20P6)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	

### Product Characteristics

<b>Color</b>	WHITE	<b>Score</b>	2 pieces
<b>Shape</b>	OVAL (Capsule shaped)	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	18;70
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1932-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2026	
2	NDC:70771-1932-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2026	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA219253	05/25/2026	

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

**Registrant** - Zydus Lifesciences Limited (677605858)

### Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		677605858	ANALYSIS(70771-1929, 70771-1930, 70771-1931, 70771-1932) , MANUFACTURE(70771-1929, 70771-1930, 70771-1931, 70771-1932)

