

**VERAPAMIL HYDROCHLORIDE- verapamil hydrochloride tablet**  
**Zydus Lifesciences Limited**

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**Verapamil Hydrochloride Extended-Release Tablets, USP**  
**Rx only**

**SPL UNCLASSIFIED**

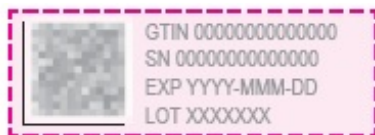
**PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

NDC 70771-1953-1 in bottles of 100 tablets

Verapamil Hydrochloride Extended-Release Tablets, USP 120 mg

100 Tablets

Rx Only



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

NDC 70771-1953-1

**Verapamil  
Hydrochloride  
Extended-Release  
Tablets, USP**

**120 mg**

zydus

100 Tablets  
Rx only

Each film coated extended-release tablet contains Verapamil Hydrochloride, USP, .....120 mg (equivalent to 111.08 mg verapamil).  
**Usual dosage:** See package insert for full prescribing information.  
Dispense in a tight, light-resistant, child-resistant container as defined in the USP.  
**This package is child-resistant.**  
**Keep this and all medicines out of the reach of children.**  
Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F) [See USP Controlled Room Temperature]. Protect from Light and Moisture.  
Dist. by: Zydus Pharmaceuticals (USA) Inc. Pennington, NJ 08534

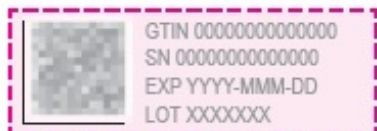
Rev.: 12/25

NDC 70771-1954-1 in bottles of 100 tablets

Verapamil Hydrochloride Extended-Release Tablets, USP 180 mg

100 Tablets

Rx Only



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

NDC 70771-1954-1

**Verapamil  
Hydrochloride  
Extended-Release  
Tablets, USP**

**180 mg**

zydus

100 Tablets  
Rx only

Each film coated extended-release tablet contains Verapamil Hydrochloride, USP .....180 mg (equivalent to 166.62 mg verapamil).  
**Usual dosage:** See package insert for full prescribing information.  
Dispense in a tight, light-resistant, child-resistant container as defined in the USP.  
**This package is child-resistant.**  
**Keep this and all medicines out of the reach of children.**  
Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F) [See USP Controlled Room Temperature]. Protect from Light and Moisture.  
Dist. by: Zydus Pharmaceuticals (USA) Inc. Pennington, NJ 08534

Rev.: 12/25

NDC 70771-1955-1 in bottles of 100 tablets

Verapamil Hydrochloride Extended-Release Tablets, USP 240 mg

100 Tablets

Rx Only



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

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NDC 70771-1955-1

# Verapamil Hydrochloride Extended-Release Tablets, USP

240 mg

Each film coated extended-release tablet contains Verapamil Hydrochloride, USP.....240 mg (equivalent to 222.16 mg verapamil).

Usual dosage: See package insert for full prescribing information.

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

This package is child-resistant.

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

Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F) [See USP Controlled Room Temperature]. Protect from Light and Moisture.

Dist. by: Zydus Pharmaceuticals (USA) Inc. Pennington, NJ 08534

Rev.: 12/25

**100 Tablets**  
Rx only

<b>VERAPAMIL HYDROCHLORIDE</b>			
verapamil hydrochloride tablet			
Product Information			
<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:70771-1953
<b>Route of Administration</b>	ORAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
<b>VERAPAMIL HYDROCHLORIDE</b> (UNII: V3888OEY5R) (VERAPAMIL - UNII:CJO37KU29)		VERAPAMIL HYDROCHLORIDE	120 mg
Inactive Ingredients			
	Ingredient Name	Strength	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)			
<b>SODIUM ALGINATE</b> (UNII: C269C4G2ZQ)			
<b>POVIDONE K30</b> (UNII: U725QWY32X)			
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)			

<b>HYPROMELLOSE 2910 (5 MPA.S)</b> (UNII: R75537T0T4)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	

### Product Characteristics

<b>Color</b>	BROWN (Beige)	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	11mm
<b>Flavor</b>		<b>Imprint Code</b>	61;5
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1953-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2026	
2	NDC:70771-1953-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2026	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA219252	05/21/2026	

## VERAPAMIL HYDROCHLORIDE

verapamil hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:70771-1954
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>VERAPAMIL HYDROCHLORIDE</b> (UNII: V3888OEY5R) (VERAPAMIL - UNII:CJ0037KU29)	VERAPAMIL HYDROCHLORIDE	180 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>SODIUM ALGINATE</b> (UNII: C269C4G2ZQ)	

<b>POVIDONE K30</b> (UNII: U725QWY32X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>HYPROMELLOSE 2910 (5 MPA.S)</b> (UNII: R75537T0T4)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	

### Product Characteristics

<b>Color</b>	BROWN (Beige)	<b>Score</b>	2 pieces
<b>Shape</b>	OVAL	<b>Size</b>	15mm
<b>Flavor</b>		<b>Imprint Code</b>	562
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1954-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2026	
2	NDC:70771-1954-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2026	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA219252	05/21/2026	

## VERAPAMIL HYDROCHLORIDE

verapamil hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:70771-1955
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>VERAPAMIL HYDROCHLORIDE</b> (UNII: V3888OEY5R) (VERAPAMIL - UNII:CJ0037KU29)	VERAPAMIL HYDROCHLORIDE	240 mg

### Inactive Ingredients

Ingredient Name	Strength
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<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>SODIUM ALGINATE</b> (UNII: C269C4G2ZQ)	
<b>POVIDONE K30</b> (UNII: U725QWY32X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>HYPROMELLOSE 2910 (5 MPA.S)</b> (UNII: R75537T0T4)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	

### Product Characteristics

<b>Color</b>	BROWN (Beige)	<b>Score</b>	2 pieces
<b>Shape</b>	OVAL (Capsule)	<b>Size</b>	18mm
<b>Flavor</b>		<b>Imprint Code</b>	563
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1955-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2026	
2	NDC:70771-1955-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2026	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA219252	05/21/2026	

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

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

### Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		677605858	ANALYSIS(70771-1953, 70771-1954, 70771-1955) , MANUFACTURE(70771-1953, 70771-1954, 70771-1955)