VERAPAMIL HYDROCHLORIDE - verapamil hydrochloride injection, solution
Zydus Lifesciences Limited

Verapamil Hydrochloride Injection, USP
Single-Dose Fliptop Vial
Protect from light.
Rx only

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70771-1601-1

Verapamil Hydrochloride Injection, USP

5 mg/2 mL

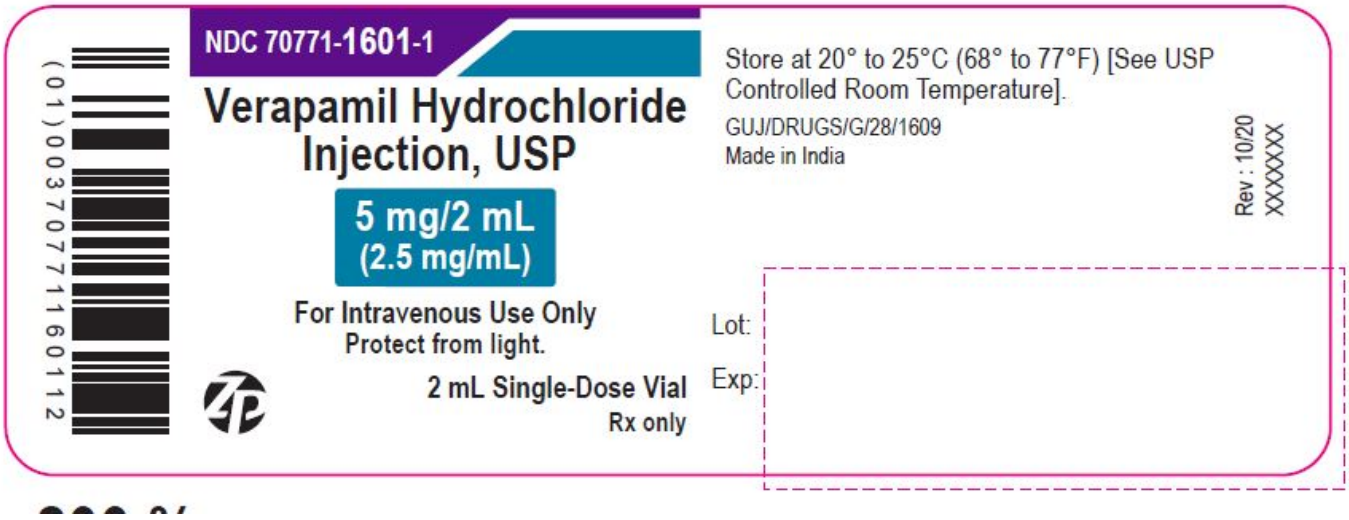
(2.5 mg/mL)

For Intravenous Use Only

Protect from light.

2 mL Single-Dose Vial

Rx only



NDC 70771-1601-5

Verapamil Hydrochloride Injection, USP

5 mg/2 mL

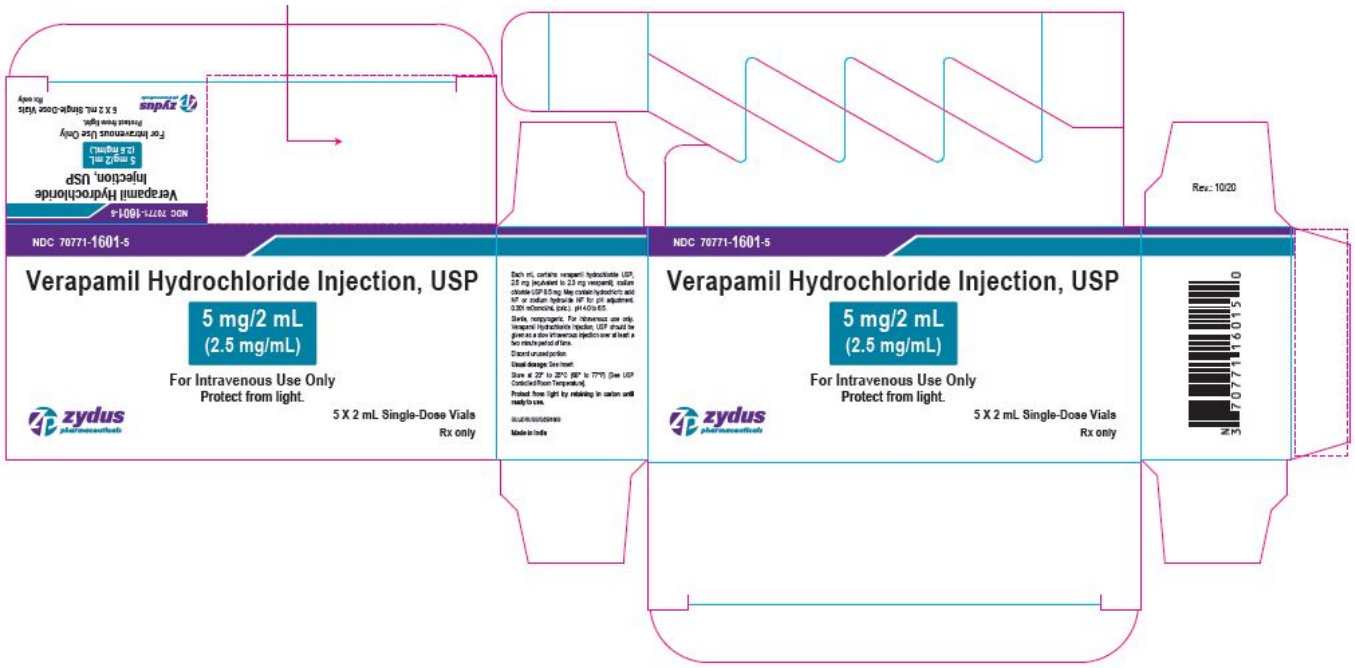
(2.5 mg/mL)

For Intravenous Use Only

Protect from light.

5 X 2 mL Single-Dose Vials

Rx only



NDC 70771-1601-7

Verapamil Hydrochloride Injection, USP

5 mg/2 mL

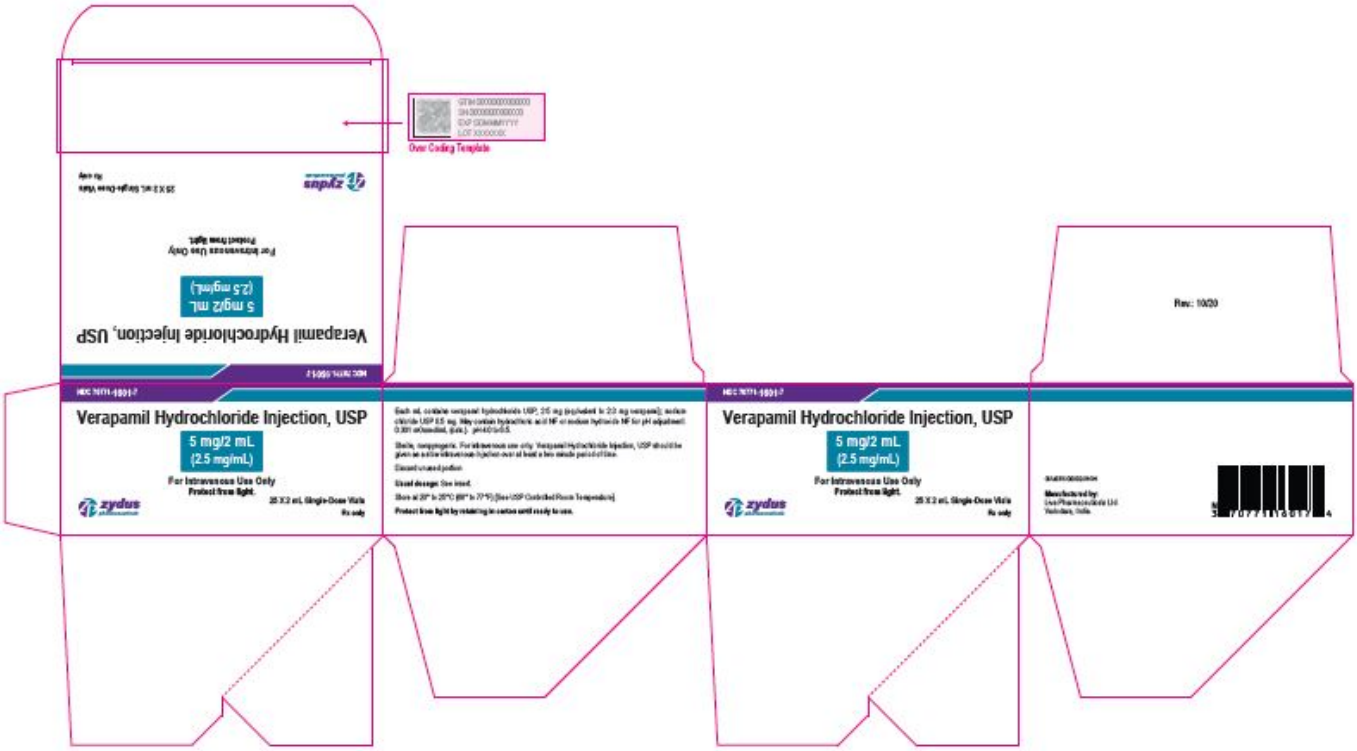
(2.5 mg/mL)

For Intravenous Use Only

Protect from light.

25 X 2 mL Single-Dose Vials

Rx only



PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70771-1602-1

Verapamil Hydrochloride Injection, USP

10 mg/4 mL

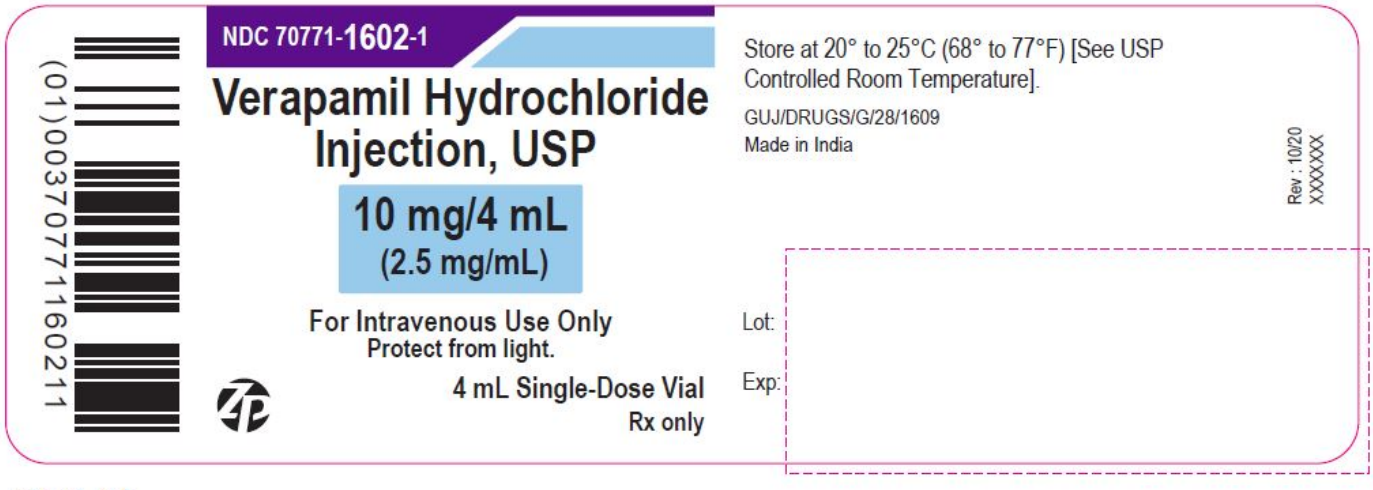
(2.5 mg/mL)

For Intravenous Use Only

Protect from light.

4 mL Single-Dose Vial

Rx only



NDC 70771-1602-5

Verapamil Hydrochloride Injection, USP

10 mg/4 mL

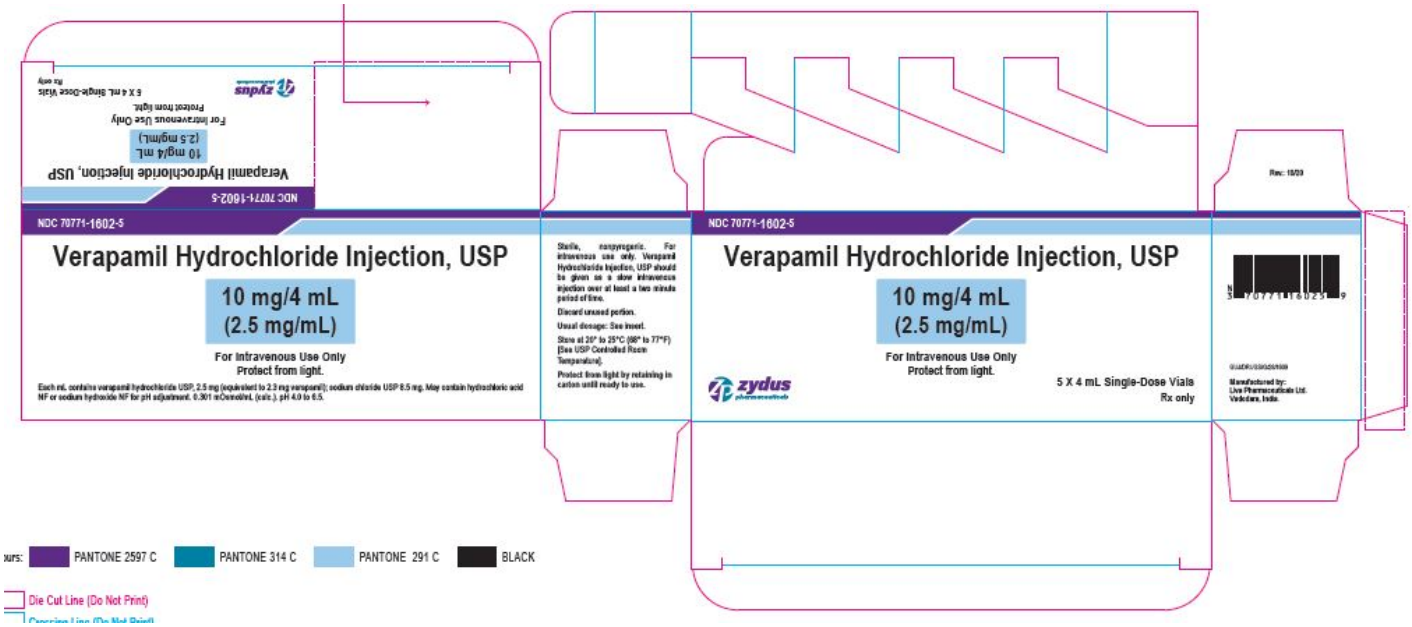
(2.5 mg/mL)

For Intravenous Use Only

Protect from light.

5 X 4 mL Single-Dose Vials

Rx only



VERAPAMIL HYDROCHLORIDE

verapamil hydrochloride injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:70771-1601
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
VERAPAMIL HYDROCHLORIDE (UNII: V3888OEY5R) (VERAPAMIL - UNII:CJ0O37KU29)	VERAPAMIL HYDROCHLORIDE	2.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

SODIUM CHLORIDE (UNII: 451W47IQ8X)	8.5 mg in 1 mL
WATER (UNII: 059QF0KO0R)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1601-7	25 in 1 CARTON	10/22/2020	
1	NDC:70771-1601-5	5 in 1 CARTON		
1	NDC:70771-1601-1	2 mL in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214215	10/22/2020	

VERAPAMIL HYDROCHLORIDE

verapamil hydrochloride injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:70771-1602
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
VERAPAMIL HYDROCHLORIDE (UNII: V3888OEY5R) (VERAPAMIL - UNII:CJ0O37KU29)	VERAPAMIL HYDROCHLORIDE	2.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	8.5 mg in 1 mL
WATER (UNII: 059QF0KO0R)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:70771-1602-5	5 in 1 CARTON	10/22/2020	
1	NDC:70771-1602-1	4 mL in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214215	10/22/2020	

Labeler - Zydus Lifesciences Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		873671928	MANUFACTURE(70771-1601, 70771-1602) , ANALYSIS(70771-1601, 70771-1602)

Revised: 11/2022 Zydus Lifesciences Limited