PLERIXAFOR- plerixafor injection
Zydus Lifesciences Limited

PLERIXAFOR injection, for subcutaneous use

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70771-1776-1

Plerixafor Injection – 1.2 mL vial label

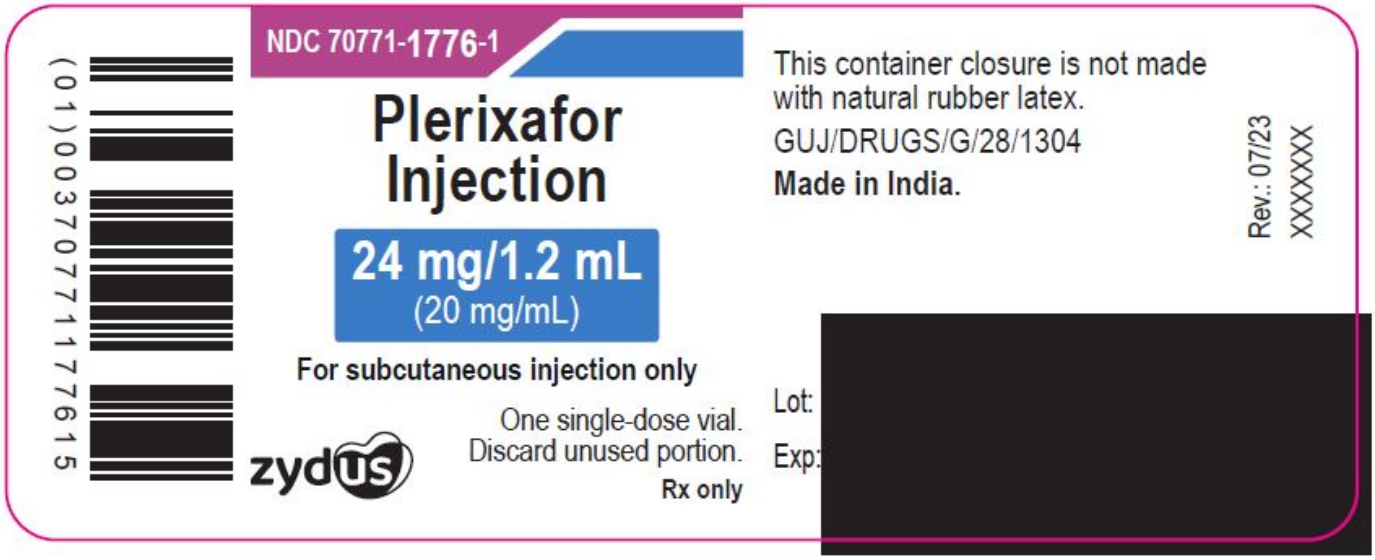
24 mg/1.2 mL

(20 mg/mL)

For subcutaneous injection only

For single-dose only

Rx only



NDC 70771-1776-1

Carton contains one vial of

Plerixafor Injection

24 mg/1.2 mL

(20 mg/mL)

For subcutaneous injection only

See package insert for dosage and administration

For single-dose only

Rx only

Over Coding Te
No Varnished A
(25 x 50 mm)

OR

PLERIXAFOR

plerixafor injection

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:70771-1776
Route of Administration	SUBCUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PLERIXAFOR (UNII: S915P5499N) (PLERIXAFOR - UNII:S915P5499N)	PLERIXAFOR	24 mg in 1.2 mL

Inactive Ingredients	
Ingredient Name	**Strength**
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	5.9 mg in 1.2 mL
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1776-1	1 in 1 CARTON	07/28/2023	
1		1.2 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA208980	07/28/2023	

Labeler - Zydus Lifesciences Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		650348852	MANUFACTURE(70771-1776) , ANALYSIS(70771-1776)

Revised: 7/2023 Zydus Lifesciences Limited