

DOXYCYCLINE - doxycycline injection, powder, lyophilized, for solution
Zydus Lifesciences Limited

Doxycycline For Injection USP

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - DOXYCYCLINE 100 MG CONTAINER LABEL

NDC 70771-1121-1

Doxycycline for Injection, USP

100 mg per vial

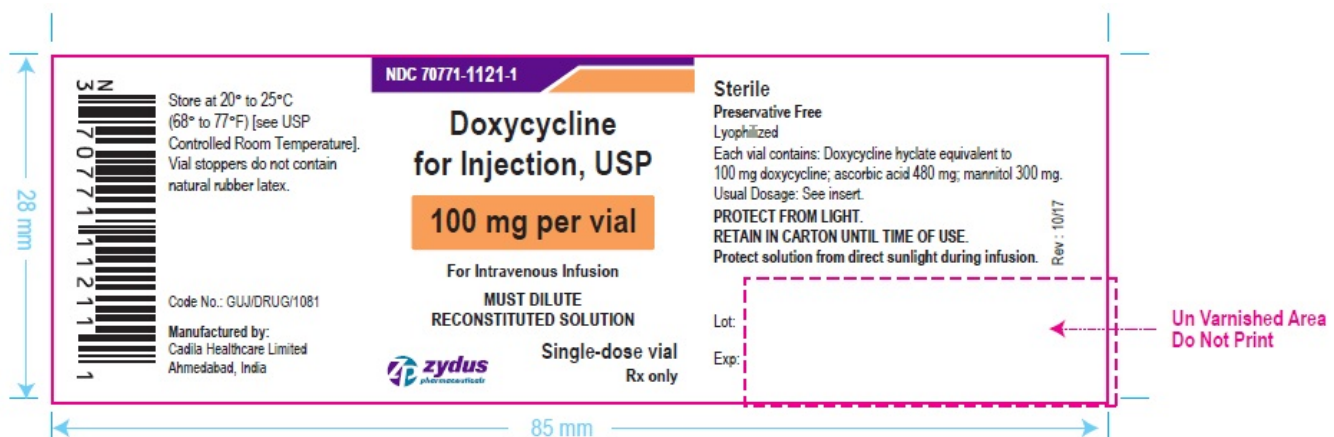
For Intravenous Infusion

MUST DILUTE RECONSTITUTED SOLUTION

Single-dose vial

Rx only

Zydus Pharmaceuticals



PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - DOXYCYCLINE 100 MG CARTON LABEL

NDC 70771-1121-6

Doxycycline for Injection, USP

100 mg per vial

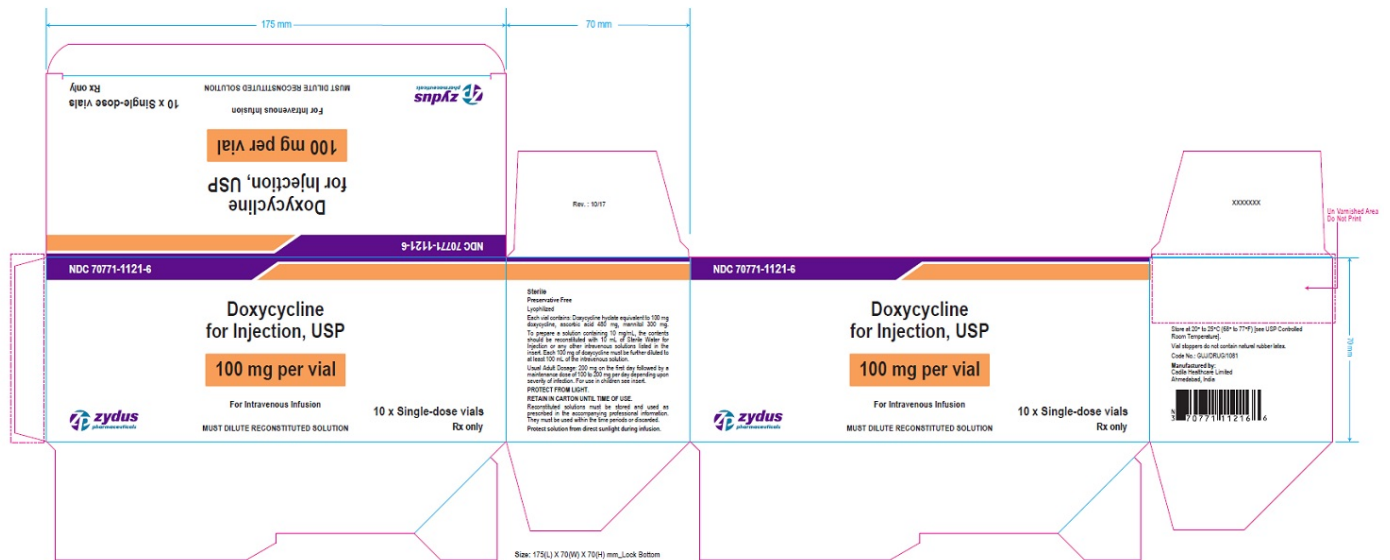
For Intravenous Infusion

MUST DILUTE RECONSTITUTED SOLUTION

10 x Single-dose vials

Rx only

Zydus Pharmaceuticals



PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - DOXYCYCLINE 200 MG CONTAINER LABEL

NDC 70771-1122-1

Doxycycline for Injection, USP

200 mg per vial

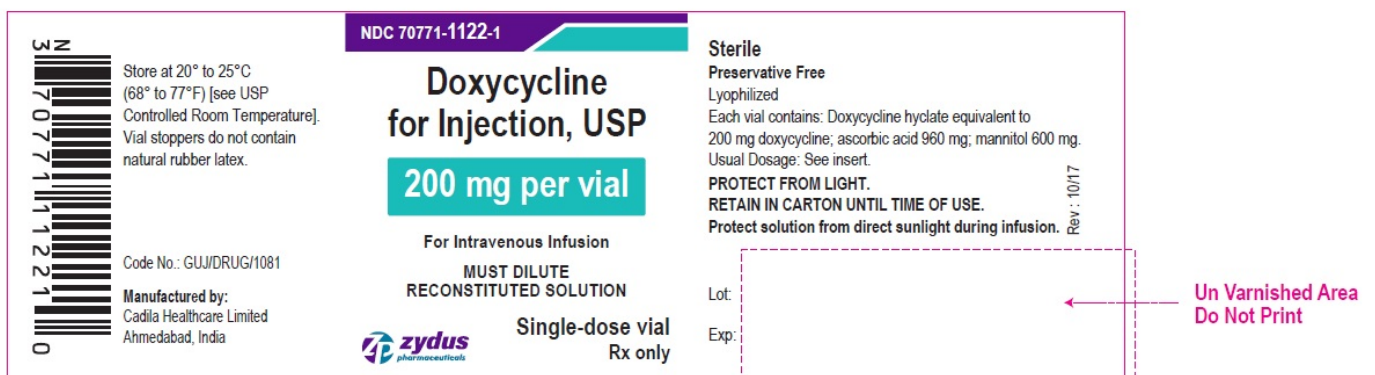
For Intravenous Infusion

MUST DILUTE RECONSTITUTED SOLUTION

Single-dose vial

Rx only

Zydus Pharmaceuticals



PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - DOXYCYCLINE 200 MG CARTON LABEL

NDC 70771-1122-1

Doxycycline for Injection, USP

200 mg per vial

For Intravenous Infusion

MUST DILUTE RECONSTITUTED SOLUTION

Single-dose vial

Rx only

Zydus Pharmaceuticals



DOXYCYCLINE

doxycycline injection, powder, lyophilized, for solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1121
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
DOXYCYCLINE HYCLATE (UNII: 19XTS3T51U) (DOXYCYCLINE ANHYDROUS - UNII:334895S862)		DOXYCYCLINE ANHYDROUS	100 mg in 10 mL	
Inactive Ingredients				
Ingredient Name		Strength		
ASCORBIC ACID (UNII: PQ6CK8PD0R)		480 mg in 10 mL		
MANNITOL (UNII: 3OWL53L36A)		300 mg in 10 mL		
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1121-6	10 in 1 CARTON	02/01/2018	
1	NDC:70771-1121-1	10 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	ANDA207757	02/01/2018		

DOXYCYCLINE

doxycycline injection, powder, lyophilized, for solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1122
Route of Administration	INTRAVENOUS		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
DOXYCYCLINE HYCLATE (UNII: 19XTS3T51U) (DOXYCYCLINE ANHYDROUS - UNII:334895S862)		DOXYCYCLINE ANHYDROUS	200 mg in 20 mL
Inactive Ingredients			
Ingredient Name		Strength	
ASCORBIC ACID (UNII: PQ6CK8PD0R)		960 mg in 20 mL	
MANNITOL (UNII: 3OWL53L36A)		600 mg in 20 mL	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1122-1	1 in 1 CARTON	02/01/2018	
1		20 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	ANDA207757	02/01/2018	

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

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		918596198	ANALYSIS(70771-1121, 70771-1122) , MANUFACTURE(70771-1121, 70771-1122)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		873671928	MANUFACTURE(70771-1121, 70771-1122) , ANALYSIS(70771-1121, 70771-1122)

Revised: 11/2022

Zydus Lifesciences Limited