

LINEZOLID - linezolid tablet, film coated
Zydus Lifesciences Limited

LINEZOLID TABLETS

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70771-1111-1

Linezolid Tablets, 600 mg

Rx only

100 tablets

ZyGenerics
NDC 70771-1111-1
Linezolid Tablets
600 mg
Rx only
100 Tablets

Each film-coated tablet contains 600 mg of linezolid.
Usual Dosage: See package insert for full prescribing information.
Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature]. Protect from light.
Dispense in tight, light-resistant container as defined in the USP.
KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.
Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India

Lot: Exp: Rev.: 08/17
No Varnish

LINEZOLID

linezolid tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1111
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LINEZOLID (UNII: ISQ9I6J12J) (LINEZOLID - UNII:ISQ9I6J12J)	LINEZOLID	600 mg

Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
CROSPVIDONE (UNII: 2S7830E561)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	WHITE (WHITE TO OFF-WHITE)	Score	no score
Shape	OVAL (OVAL)	Size	19mm
Flavor		Imprint Code	413
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1111-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	08/09/2017	
2	NDC:70771-1111-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	08/09/2017	
3	NDC:70771-1111-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	08/09/2017	
4	NDC:70771-1111-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	08/09/2017	
5	NDC:70771-1111-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	08/09/2017	
6	NDC:70771-1111-0	1000 in 1 BOTTLE; Type 0: Not a Combination Product	08/09/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206097	08/09/2017	

Labeler - Zydus Lifesciences Limited (918596198)

Registrant - Zydus Lifesciences Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		863362789	ANALYSIS(70771-1111) , MANUFACTURE(70771-1111)

Revised: 10/2022

Zydus Lifesciences Limited