

CLOPIDOGREL- clopidogrel tablet, film coated
Zydus Lifesciences Limited

CLOPIDOGREL TABLETS

Manufactured by:
Cadila Healthcare Ltd.
India.

SPL MEDGUIDE

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 70771-1062-9 in bottle of 90 Tablets
Clopidogrel Tablets USP, 75 mg
Rx only
90 TABLETS

ZyGenerics
NDC 70771-1062-9
CLOPIDOGREL
Tablets, USP
75 mg

Each tablet contains 97.830 mg of clopidogrel bisulfate USP equivalent to 75 mg of clopidogrel base

Usual Adult Dosage: See package insert for complete prescribing information.

Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature].
Dispense in a tight container.

KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.

Manufactured by:
Cadila Healthcare Ltd.
Ahmedabad, India

PHARMACIST : Dispense the Medication Guide provided separately to each patient.

Rx only
90 Tablets

Lot: _____
Exp: _____
Rev.: 02/17

CLOPIDOGREL

clopidogrel tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70771-1062
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLOPIDOGREL BISULFATE (UNII: 08I79HTP27) (CLOPIDOGREL - UNII:A74586SNO7)	CLOPIDOGREL	75 mg

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CROSPVIDONE (15 MPA.S AT 5%) (UNII: 68401960MK)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
HYDROXYPROPYL CELLULOSE (160000 WAMW) (UNII: RFW2ET671P)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	PINK (PINK)	Score	no score
Shape	ROUND (ROUND)	Size	9mm
Flavor		Imprint Code	379
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70771-1062-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	02/08/2017	
2	NDC:70771-1062-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/08/2017	
3	NDC:70771-1062-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	02/08/2017	
4	NDC:70771-1062-0	1000 in 1 BOTTLE; Type 0: Not a Combination Product	02/08/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA201686	02/08/2017	

Labeler - Zydus Lifesciences Limited (918596198)

Registrant - Zydus Lifesciences Limited (918596198)

Establishment

Name	Address	ID/FEI	Business Operations
Zydus Lifesciences Limited		918596198	ANALYSIS(70771-1062) , MANUFACTURE(70771-1062)

Revised: 10/2023

Zydus Lifesciences Limited