

CLOPIDOGREL- clopidogrel tablet, film coated
Cadila Healthcare Limited

CLOPIDOGREL TABLETS

Manufactured by:
Cadila Healthcare Ltd.
India.

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

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

Rx only
90 Tablets

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

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 MPAS AT 5%) (UNII: 68401960MK)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
HYDROXYPROPYL CELLULOSE (1600000 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 - Cadila Healthcare Limited (918596198)

Registrant - Cadila Healthcare Limited (918596198)

Establishment

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

Revised: 8/2020

Cadila Healthcare Limited