

**CITALOPRAM- citalopram tablet, film coated
DIRECT RX**

CITALOPRAM 10mg 30 TABS

DESCRIPTION

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CLINICAL PHARMACOLOGY

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INDICATIONS AND USAGE

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CONTRAINDICATIONS

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WARNINGS

-

PRECAUTIONS

-

-

ADVERSE REACTIONS PART 2

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DRUG ABUSE AND DEPENDENCE

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OVERDOSAGE

-

DOSAGE AND ADMINISTRATION

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ANIMAL TOXICOLOGY

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Medication Guide

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PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

D **CITALOPRAM**
10mg* 30 Tabs

Generic For: **CELEXA**
*Each tablet contains: Citalopram hydrobromide, USP equivalent to 10mg citalopram base

Lot# **Prod# 288-30** Discard After: 05/17

Packaged and Distributed By: **DIRECT** Alphaletta, GA 30005

Dist. By: Amneal Pharmaceuticals
Gilead, KY 42141
NDC 65162-052-50

Mfg Lot: 3/22/2016

AFWXX

Caution: Federal law prohibits transfer of this drug to any person other than the patient for whom it was prescribed.

RX ONLY-KEEP OUT OF REACH OF CHILDREN

Dosage: See package insert. Store between 68-77 degrees F

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NDC 61919-288-30

CITALOPRAM 10mg*
NDC 61919-288-30 30 Tab
Lot Exp Date 05/17
Mfg NDC 65162-052-50

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REVISION | Effective Date | Lot/Package | Days

CITALOPRAM

citalopram tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61919-288(NDC:65162-052)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CITALOPRAM HYDROBROMIDE (UNII: IIE9D14F36) (CITALOPRAM - UNII:0DHU5B8D6V)	CITALOPRAM	10 mg

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STARCH, CORN (UNII: O8232NY3SJ)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
TALC (UNII: 7SEV7J4R1U)	
POVIDONE K12 (UNII: 333AG72FWJ)	

Product Characteristics

Color	orange	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	IP;52

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61919-288-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077289	03/22/2016	

Labeler - DIRECT RX (079254320)**Establishment**

Name	Address	ID/FEI	Business Operations
DIRECT RX		079254320	repack(61919-288)

Revised: 4/2016

DIRECT RX