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

-

DRUG ABUSE AND DEPENDENCE

-

OVERDOSAGE

-

DOSAGE AND ADMINISTRATION

-

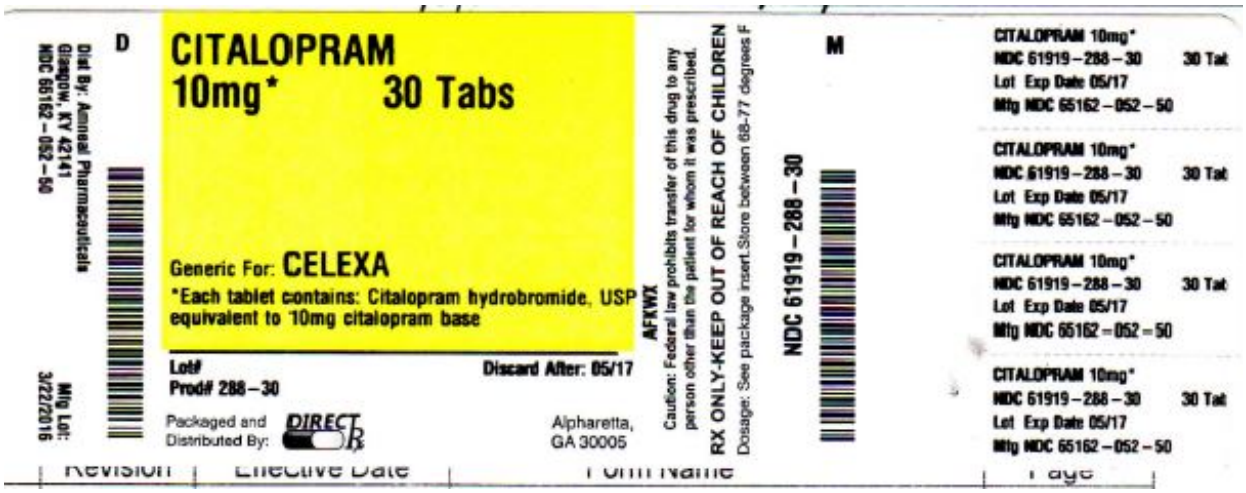
ANIMAL TOXICOLOGY

-

Medication Guide

-

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



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: I1E9D14F36) (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