

LIDOCAINE- lidocaine patch
DIRECT RX

LIDOCAINE PATCH 5%

DESCRIPTION

-

CLINICAL PHARMACOLOGY

-

CLINICAL STUDIES

-

INDICATION AND USAGE

-

CONTRAINDICATIONS

-

WARNINGS

-

Close

PRECAUTIONS

-

ADVERSE REACTIONS

-

OVERDOSAGE

-

DOSAGE AND ADMINISTRATION

-

HANDLING AND DISPOSAL

-

PRINCIPAL DISPLAY PANEL

LIDOCAINE PATCH 5% 30 Patches

Generic For: **LIDODERM**
 Each adhesive patch contains: Lidocaine, 700mg (50mg per gram adhesive) in an aqueous base

Lot# **Prod# 779-30** Discard After: 07/17

Alpharetta, GA 30005

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

Keep used and unused patches out of the reach of children, pets, and others

Mfg For: Qualitest Pharmaceuticals
 Huntsville, AL 35811
 NDC 0603-1880-16

Mfg Lot: 3/3/2016

LIDOCAINE PATCH 5%
 NDC 61919-779-30 30 Patch
 Lot Exp Date 07/17
 Mfg NDC 0603-1880-16

LIDOCAINE PATCH 5%
 NDC 61919-779-30 30 Patch
 Lot Exp Date 07/17
 Mfg NDC 0603-1880-16

LIDOCAINE PATCH 5%
 NDC 61919-779-30 30 Patch
 Lot Exp Date 07/17
 Mfg NDC 0603-1880-16

LIDOCAINE PATCH 5%
 NDC 61919-779-30 30 Patch
 Lot Exp Date 07/17
 Mfg NDC 0603-1880-16

LIDOCAINE				
lidocaine patch				
Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61919-779(NDC:0603-1880)	
Route of Administration	CUTANEOUS			
Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	700 mg		
Inactive Ingredients				
Ingredient Name	Strength			
UREA (UNII: 8W8T17847W)				
GLYCERIN (UNII: PDC6A3C0OX)				
POLYVINYL ALCOHOL (UNII: 532B59J990)				
SODIUM POLYACRYLATE (250000 MW) (UNII: 05I15JN12J)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
GELATIN (UNII: 2G86QN327L)				
SORBITOL (UNII: 506T60A25R)				
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)				
TARTARIC ACID (UNII: W48881119H)				
KAOLIN (UNII: 24H4NWX5CO)				
METHYL PARABEN (UNII: A218C7HI9T)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
PROPYL PARABEN (UNII: Z8IX2SC10H)				
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)				
DIHYDROXYALUMINUM AMINO ACETATE (UNII: DO250MG0W6)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:61919-779-30	30 in 1 CARTON; Type 0: Not a Combination Product	03/03/2016	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
NDA	NDA020612	03/03/2016		

Labeler - DIRECT RX (079254320)

Establishment			
Name	Address	ID/FEI	Business Operations
DIRECT RX		079254320	repack(61919-779)

Revised: 3/2016

DIRECT RX