LIDOCAINE- lidocaine patch
DIRECT RX

LIDOCAINE PATCH 5%

DESCRIPTION
-

CLINICAL PHARMACOLOGY
-

CLINICAL STUDIES
-

INDICATION AND USAGE
-

CONTRAINDICATIONS
-

WARNINGS
-

Close
PRECAUTIONS
-

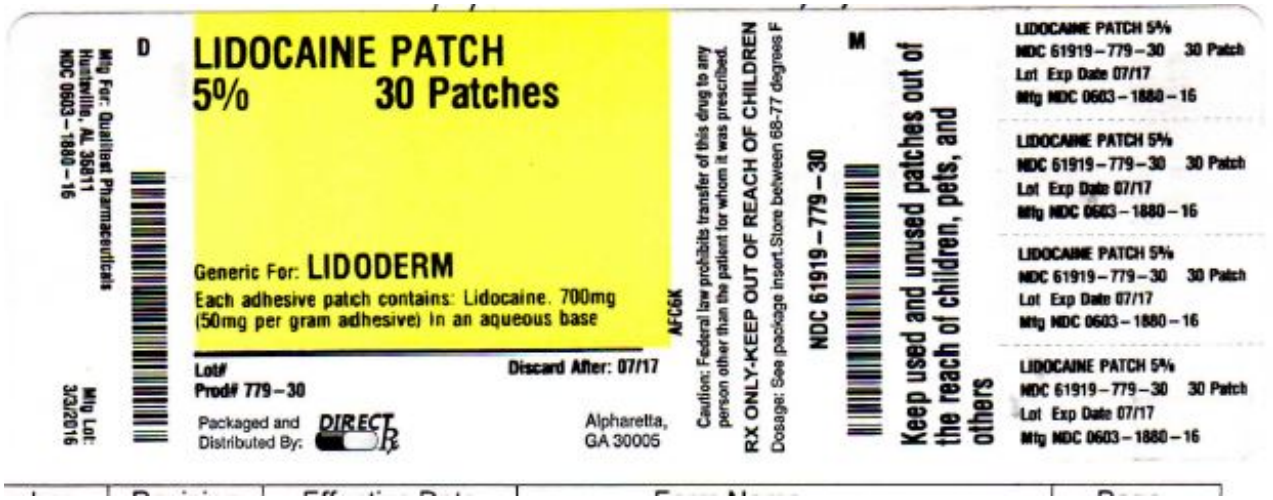
ADVERSE REACTIONS
-

OVERDOSAGE
-

DOSAGE AND ADMINISTRATION
-

HANDLING AND DISPOSAL
-

PRINCIPAL DISPLAY PANEL



LIDOCAINE

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 (2500000 MW) (UNII: 05I15JNI2J)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GELATIN (UNII: 2G86QN327L)	
SORBITOL (UNII: 506T60A25R)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
TARTARIC ACID (UNII: W4888I119H)	
KAOLIN (UNII: 24H4NWX5CO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
DIHYDROXYALUMINUM AMINOACETATE (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