LIDOPATCH- lidocaine, menthol patch
Wuhan Bingbing Pharmaceutical Co., Ltd

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Lidopatch

WARNINGS SECTION
For external use only. Do not use if you are allergic or sensitive to lidocaine or menhol. Do not use if pouch is damaged or opened.

PURPOSE

Temporarily relieves minor pain associated with: arthritis, simple backache, bursitis, tendonitis, muscle straints, sprains and bruises.

ACTIVE INGREDIENT SECTION
Lidocaine HCL 3.6% - Topical analgesic
Menthol 1.25% - Topical analgesic

STOP USE SECTION
Stop Use and ask doctor if:
excessive redness or irritation is present
Pain persists for more than 7 days
symptoms cear up and occur again within a few days
conditions worsens
If pregnant or breasfeading, ask a health professional before use. keep out of reach of children. If swallowed, get medical help or contact a poison control center right away.

DOSAGE & ADMINISTRATION SECTION

Directions: Aduts apply patch to affected area for a maximum of 12 hours.
Do not use more than 1 patch every 24 hours.
children : consult your physician
remove protective film, gently apply to affected area.

INACTIVE INGREDIENT SECTION

Aloe Barbadensis, Diazolidinyl Urea, EDTA Disodium Salt, Glycerin, Iodpropynyl, Butylcarbonate, Methylparaben, Polysorbate 80, Propylparaben, Sodium Polyacrylate, Water

Usage
Aduts apply patch to affected area for a maximum of 12 hours.
Do not use more than 1 patch every 24 hours

KEEP OUT OF REACH OF CHILDREN.

LIDOPATCH

lidocaine, menthol patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70267-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	0.036 mg
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.0125 mg

Inactive Ingredients	
Ingredient Name	**Strength**
COBALT DISODIUM EDETATE (UNII: 3EY1Y2QRLI)	
GLYCERIN (UNII: PDC6A3C0OX)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
WATER (UNII: 059QF0KO0R)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLPARABEN (UNII: A2I8C7HI9T)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70267-001-02	1 in 1 PACKAGE; Type 0: Not a Combination Product	03/02/2016	

2	NDC:70267-001-01	30 in 1 PACKAGE; Type 0: Not a Combination Product	03/02/2016	
3	NDC:70267-001-03	30 in 1 PACKAGE; Type 0: Not a Combination Product	03/02/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	03/02/2016	

Labeler - Wuhan Bingbing Pharmaceutical Co., Ltd (529613802)

Establishment

Name	Address	ID/FEI	Business Operations
Wuhan Bingbing Pharmaceutical Co., Ltd.		529613802	manufacture(70267-001)

Revised: 1/2019 Wuhan Bingbing Pharmaceutical Co., Ltd