LIDENZA PATCH- lidocaine hydrochloride patch
Patchwerx Labs, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Lidenza Patch

Warnings

For external use only

Avoid contact with eyes

Do not applly to open wounds or damaged skin

If symptoms persist for more than seven days, discontinue use and consult physician

If swallowed, consult physician

Do not bandage tightly

If pregnant or breast feeding, contact physician prior to use

Do not use in large quantities, particularly over raw surfaces or blistered areas

Keep out of reach of children

Lidocaine HCL 4.00%

Menthol 1.00%

aloe barbadensis leaf, water, arnica montan extract, boswella serrata extract, camelia sinensis leaf extract, carbomer, ethylhexylglycerine, glycerine, isopropyl myristate, PEG 8, phenoxyethanol, polysorbate 80, sodium lauryl sulfate, triethanolamine, fd&c blue #1, fd&c yellow #5

clean and dry affected area

remove patch from backing and apply to affected area

use only one patch at a time, and maximum of four patches/day

leave patch on affected area for up to 8 hours

do not use patches for longer than five consecutive days

children under 12 should consult physician prior to use

Temporary relief of pain associated with minor cuts, scrapes and minor skin irritations

Topical anesthetic

External analgesic

Lidenza Patch

Lidenza Patch **Rx Only**

Lidenza Patch

(Lidocaine 4% / Menthol 1%)

QTY : 15 patches

NDC 69329-017-15

RX Only

Lidenza Patch **RX Only**

Lidenza Patch

DRUG FACTS:		
ACTIVE INGREDIENTS:		
Lidocaine HCL	4.00%	Topical Anesthetic
Menthol	1.00%	External Analgesic

USES:
For Temporary relief of pain associated with minor cuts, scrapes and minor skin irritations.

WARNINGS:
- For external use only.
- Avoid contact with eyes.
- Do not apply to open wounds or damaged skin.
- If symptoms persist for more than seven days, discontinue use and consult physician.
- **Keep out of reach of children.** If swallowed, consult physician.
- Do not bandage tightly.
- If pregnant or breast feeding, contact physician prior to use.
- Do not use in large quantities, particularly over raw surfaces or blistered areas.

DIRECTIONS:
- Clean and dry affected area.
- Remove patch from backing and apply to affected area.
- Use only one patch at a time, and maximum of four patches/day.
- Leave patch on affected area for up to 8-hours.
- Do not use patches for longer than five consecutive days.
- Children under 12 should consult physician prior to use.

OTHER INGREDIENTS:
Aloe Barbadensis Leaf (Aloe Vera Juice) Gel, Aqua (Deionized Water), Arnica Montana Extract, Boswella Serrata Extract, Camellia Sinensis Leaf (Green Tea) Extract, Carbomer, Ethylhexylglycerin, Glycerin, Isopropyl Myristate, PEG-8, Phenoxyethanol, Polysorbate-80, Sodium Lauryl Sulfate, Triethanolamine, FD&C Blue #1, FD&C Yellow #5.

Store below 25 degrees. Avoid direct sunlight.

Manufactured for: Patchwerx Labs, Inc
Las Vegas, NV 89118
For Questions or Comments call +1 (800) 590-8070

Made in USA / Patent Pending

69329 0715

LIDENZA PATCH

lidocaine hydrochloride patch

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:69329-017(NDC:49430-017)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ARNICA MONTANA (UNII: O80TY208ZW)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BOSWELLIA SACRA WHOLE (UNII: 8O600AZL0W)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CARBOMER 1342 (UNII: 809Y72KV36)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PEG-8 GLYCERYL ISOSTEARATE (UNII: 74QQ5X3KL1)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TRIETHANOLAMINE BENZOATE (UNII: M3EN4GC19W)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69329-017-15	100 g in 1 PACKAGE		
2	NDC:69329-017-25	100 g in 1 PACKAGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/01/2013	

Labeler - Patchwerx Labs, Inc. (079584480)

Revised: 11/2014 Patchwerx Labs, Inc.