# RICORA- lidocaine hydrochloride, menthol patch Meds Direct Rx, Inc.

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.

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## Ricora Patch

## **DRUG FACTS:**

## **ACTIVE INGREDIENTS:**

Lidocaine HCL 4.00% Menthol 1.00%

Topical Anesthetic

External Analgesic

## **USES:**

For temporary relief of pain associated with minor: •cuts •scrapes •skin irritations

## **WARNINGS:**

- For external use only.
- Avoid contact with eyes.
- **Do not** apply on 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 breastfeeding,

contact physician prior to use.

## Do not

use in large quantities, particularly over raw surfaces or blistered areas.

## **DIRECTIONS:**

## (Adults and Children 12 years of age and over)

- 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 5 consecutive days.

• Children under 12 should consult physician prior to use.

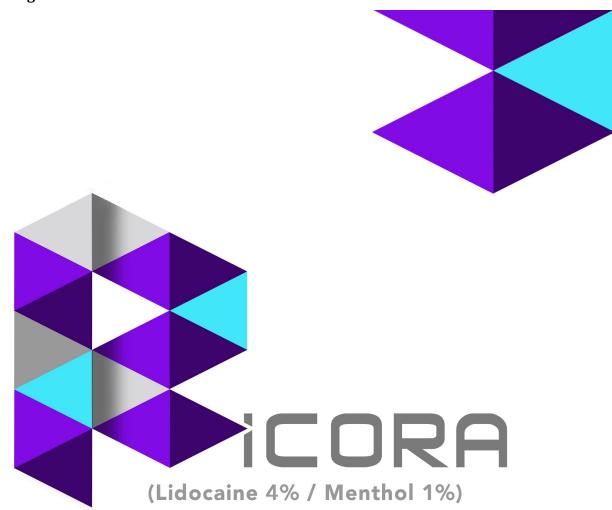
## **INACTIVE INGREDIENTS:**

Glyceryl Hydrogenated Rosinate, Hydrated Silica, Mineral Oil, PEG-400, Polyisobutene, Styrene/Isoprene Copolymer

Creates a warming sensation to the applied area.

Store below 25 degrees. Avoid direct sunlight.

## Package Lebeling:





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NDC: 69418-006-30

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Manufactured For: MEDS DIRECT RX INC

882 Third Avenue 10th Floor Suite 1000

Brooklyn, NY 11232

Questions or Comments call 855-480-MEDS

**Patches** 

Made in China

## **RICORA**

lidocaine hydrochloride, menthol patch

## **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69418-006
Route of Administration	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength	
LIDO CAINE HYDRO CHLO RIDE (UNII: V13007Z41A) (LIDO CAINE - UNII:98 PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 mg in 1 g	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
HYDRATED SILICA (UNII: Y6O7T4G8P9)		
MINERAL O IL (UNII: T5L8T28FGP)		
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)		

I	Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
1	NDC:69418-006-30	6 in 1 BOX			
1		5 in 1 POUCH			
1		2.3 g in 1 PATCH; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	11/21/2015		

# Labeler - Meds Direct Rx, Inc. (064053428)

Establishment				
Name	Address	ID/FEI	Business Operations	
Meds Direct Rx, Inc.		064053428	repack(69418-006), relabel(69418-006)	

Revised: 12/2015 Meds Direct Rx, Inc.