

**MAXIMUN STRENGTH LIDOCAINE PATCH PLUS MENTHOL- lidocaine, menthol patch  
Meijer Distribution 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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**Meijer Maximun Strength Lidocaine Patch Plus Menthol**

**Drug Facts**

Lidocaine 4% Topical Anesthetic

Menthol 1% Topical Anesthetic

Topical Anesthetic

**USES**

Temporary relief of minor pain

**WARNINGS**

- For External use only. Use only as directed.
- Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
- More than one patch on your body at a time
- On cut, irritated or swollen skin
- On puncture wounds
- For more than one week without consulting a doctor
- If you are allergic to any active or inactive ingredients
- If pouch is damaged or opened.

**If pregnant or breast feeding**

- Contact a physician prior to use.

**WHEN USING:**

- Use only as directed
- Read and follow all directions and warnings on this carton
- Do not allow contact with the eyes
- Do not use at the same time as other topical analgesics
- Do not bandage tightly or apply local heat (such as heating pads) to the area of use
- Do not microwave
- Dispose of used patch in manner that always keeps product away from children and pets. Used patches still contain the drug product that can produce serious adverse effects if a child or pet chews or ingests this patch.

**Stop use and consult a doctor if**



# MAXIMUN STRENGTH LIDOCAINE PATCH PLUS MENTHOL

lidocaine, menthol patch

## Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-841
Route of Administration	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g in 100 g
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	1 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
DIHYDROXYALUMINUM AMINO ACETATE ANHYDROUS (UNII: 1K713C615K)	
GLYCERIN (UNII: PDC6A3C0OX)	
KAOLIN (UNII: 24H4NWX5CO)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL 1-(2-METHYLBUTYRATE) (UNII: 9Q5W5G6461)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
SODIUM POLYACRYLATE (250000 MW) (UNII: 05115JN12J)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0K00R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-841-05	5 in 1 POUCH	07/01/2018	
1		8 g in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/01/2018	

Labeler - Meijer Distribution Inc (006959555)

## Establishment

Name	Address	ID/FEI	Business Operations
Foshan Aqua Gel Biotech Co.,Ltd.		529128763	manufacture(41250-841)

Revised: 12/2017

Meijer Distribution Inc