WELL LIDO GEL PATCH- lidocaine patch
True Marker Pharmaceuticals, Inc.

Active Ingredients	**Purposes**
Lidocaine 4%	Topical Analgesic

Uses

For temporary relief of pain.

Warnings

For External Use only.

Stop use and ask a doctor

- Localized skin reaction occur, such as rash, itching, redness, irritation, pain, swelling and blistering
- Condition worsens
- Symptoms persist for more than 7 days
- Symptoms clear up and occur again within a few days

Keep out of the reach of children
If swallowed, get medical help or contact Poison Control Center right away.

Do Not Use
• More than 1 patch a time
• On wounds or damaged skin
• With a heating pad
• If you are allergic to any ingredients of this product

If pregnant or Breast-Feeding
Ask a health professional before use

When Using this product
• Use only as directed
• Avoid contact with eyes, mucous membranes or rashes
• Do not bandage tightly

Directions

Adult and children 12 years of age and over:
• Clean and dry affected area
• Remove film from patch and apply to the skin
• Apply 1 patch at a time to affected area, not more that 3 to 4 times a daily
• Remove patch from the skin after at most 8-hour application
• Children under 12 year of age. Consult a doctor.

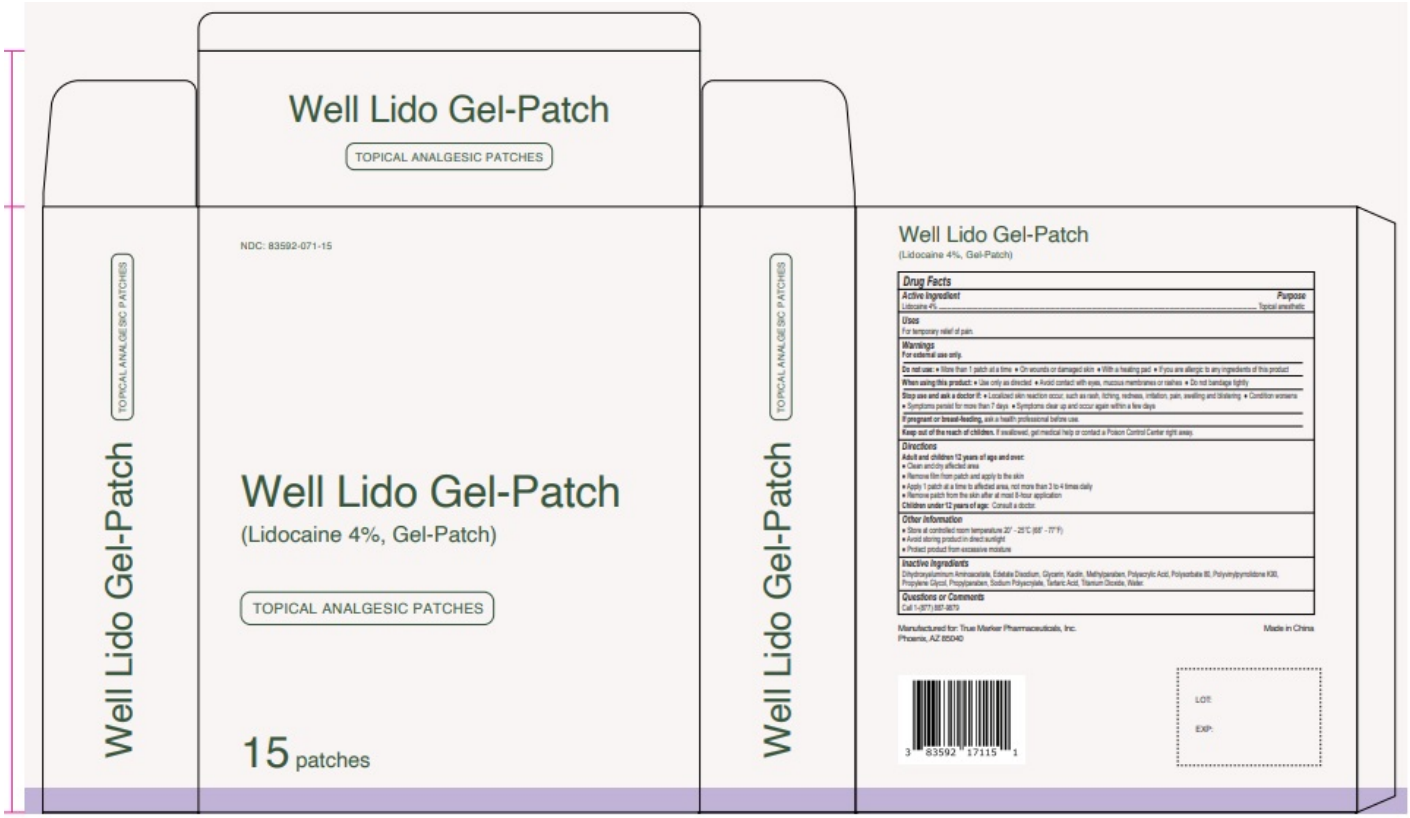
Inactive Ingredients

Dihydroxyaluminum aminoacetate, Edetate disodium, Glycerin, Kaolin, Methylparaben, Polyacrylic acid, Polysorbate 80, Polyvinylpyrrolidone K90, Propylene glycol, propylparaben, sodium polyacrylate, tartaric acid, titanium dioxide, water.

Other Information

- Store at controlled room temperature 68º-77ºF (20º-25ºC)
- Avoid storing product in direct sunlight
- Protect product from excessive moisture

Product label



WELL LIDO GEL PATCH

lidocaine patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83592-071
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	4 g

Inactive Ingredients

Ingredient Name	Strength
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
KAOLIN (UNII: 24H4NWX5CO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE K90 (UNII: RDH86HJV5Z)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
TARTARIC ACID (UNII: W4888I119H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83592-071-15	15 in 1 BOX	09/12/2024	
1		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	09/12/2024	

Labeler - True Marker Pharmaceuticals, Inc. (119046582)

Revised: 9/2024 True Marker Pharmaceuticals, Inc.