REMATEX- rematex pain relieving cream cream Home Aide Diagnostics, 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.

Veltrix Pain Relieiving Cream

ACTIVE INGREDIENTS

Lidocain 4%	
——————————————————————————————————————	
Topical analgesic	
Menthol 1%	
Topical analgesic	

INDICATIONS

For the temporary relif of minor aches and pains of muscles and joints associated with:

- Simple Backache
- Arthritis
- Strains
- Sprains
- Bruises

WARNINGS

FOR EXTERNAL USE ONLY.

- Avoid contact with eyes.
- If condition worsens, of if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a sphysician.
- Do not apply to wounds or damaged skin
- Do not bandage tightly

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 daily
- Leave patch on affected area for up to 8-hours
- Do not use patches for longer than 5 consecutive days unless directed by a physician.
- Store below 25 °C (77 °F). Avoid direct sunlight.

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

If pregnant of breast-feeding, ask a healthcare professional before use

• This product may cause allergic reaction in some individuals.

INACTIVE INGREDIENTS

aliphatic alcohol; stearic acid; vaseline; peregal; Carmellose Sodium; water;

• Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.

• Children under 2 years of age: consult a physician.

DRUG FACTS	
ACTIVE INGREDIENTS	
Lidocain 4%	Topical analger
Menthal 1%	Topical analges
INDICATIONS	
For the temporary relif of pain of muscles and joints associate	nd with:
• Minor Burns • Sunburn • Minor Cuts • Scrapes • Insect	Bites • Minor Skin Irritations
WARNINGS	
FOR EXTERNAL USE ONLY.	
Avoid contact with eyes.	
 If condition worsens, of if symptoms pensist for more than 7 days 	or clear up and occur again within a fe
days, discontinue use of this product and consult a s physician.	
 Do not apply to wounds or damaged skin 	
 Do not use in large quantities, particularly over raw surfaces or blin 	stered areas
DIRECTIONS	
Adults and children 2 years of age and older. Apply to affected area	not more than 3 to 4 times daily.
Children under 2 years of age: consult a physician.	
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 daily	
Leave patch on affected area for up to 8-hours	
Do not use patches for longer than 5 consecutive days unless direct	ed by a physician.
Store below 25 °C (77 °F). Avoid direct sunlight.	100,000
Keep out of reach of children. If s-allowed, get medical help or co	
If pregnant of breast-feeding, ask a healthcare professional before use	
 This product may cause allergic reaction in some individuals. 	
INACTIVE INGREDIENTS	
alighatic alcohol; stearic acid; vaseline; peregal; Carmellose; odium;	water





Distributed by: Home Aide Diagnostics, Inc. 1072 S. Powerline Road Deerfield Beach, FL 33442 1 (800) 915-0116



NDC #: 69379-0041-01

4 g in 100 g

Lot #: Exp Date:

Net Weight: 120 g

LIDOCAINE

REMATEX

rematex pain relieving cream cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69379-041

Route of Administration TOPICAL

LIDO CAINE (UNII: 98PI200987) (LIDO CAINE - UNII:98PI200987)

Active Ingredient/Active Moiety Ingredient Name Basis of Strength MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) MENTHOL 1 g in 100 g

Inactive Ingredients

Ingredient Name
Strength
STEARIC ACID (UNII: 4ELV7Z65AP)

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:69379-041- 01	120 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/01/2015		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/01/2015	

Labeler - Home Aide Diagnostics, Inc. (783518983)

Registrant - Zhejiang Bangli Medical Products Co. Ltd (421295875)

Establishment						
Name	Address	ID/FEI	Business Operations			
Zhejiang Bangli Medical Products Co. Ltd		421295875	manufacture(69379-041)			

Revised: 6/2015 Home Aide Diagnostics, Inc.