LANLIJEN MENTHOL PAIN RELIEF- menthol gel Jiangsu Chaben Medical Healthcare Technology Co.,Ltd

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.

Lanlijen, Menthol Pain Relief, 3fl.oz/89ml, Gel

Active ingredient

Menthol 4.0% w/w Purpose: Topical Analgesic

Purpose

Topical Analgesic

Uses

Temporary relief from minor aches and pains of sore muscles and joints associated with arthritic, backache, strains and sprains

Warnings

For external use only

Flammable, Keep away from excessive heat or open flame

Ask a doctor before use if you have Sensitive Skin

Stop use and ask a doctor if burning discomfort or excessive skin irritation develops, conditions worsen, or symptoms persist for more than 7 days, or clear up and reoccur within a few days

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When using this product

Avoid contact with eyes or mucous membranes • Do not apply to wounds or damaged skin • Do not apply to irritated skin or if excessive irritation develops • Do not use with other creams, sprays, ointments, or liniments • Do not use with heating pad or device • Store in a cool dry place • Do not bandage • Use only as directed • Wash hands after

use with cool water

- Do not apply to wounds or damaged skin Do not apply to irritated skin or if excessive irritation develops Do not use with other creams, sprays, ointments, or liniments Do not use with heating pad or device
- Store in a cool dry place

If pregnant or breast-feeding

Ask a health professional before use

Keep out of reach of children

If accidentally ingested, get medical help or contact a Poison Control Center immediately

Directions

Adults and children 12 years of age and older, Rub a thin film over affected areas not more 4 times daily; massage not necessary

Children under 12 years of age, Consult physician

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Inactive Ingredients

Alcohol, Aloe Barbadensis Leaf Extract, Arnica Montana Flower Extract, Boswellia Carterii Resin Extract, Camellia Sinensis Leaf Extract, Camphor, Carbomer, FD&C Blue 1, FD&C Yellow 5, Glycerin, llex Paraguariensis Leaf Extract, Isopropyl Myristate, Silica, Triethanolamine, Vitamin E, Water

Questions or Comments 86-513-89072216

E-mail: info@lanlijen.com www.lanlijen.com Tel:86-513-89072216

No Animal Testing, No NSAIDs, Ibuprofen, Aspirin or Salicylate

Package label. Principal display panel

Lanijen

FAST ACTING

PAIN-RELIEF GEL

LONG LASTING

MENTHOL PAIN RELIEF

QUICKLY ABSORBED, NON-GREASY WORKS NATURALLY WITH BODY

3FL.OZ | 89ML

COLD THERAPY PAIN RELIEF FOR SORE MUSCLES, BACKACHES, SORE JOINTS AND ARTHRITIS NDC 00000000000

Drug Facts

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Questions or Comments 86-513-89072216

Produced for: Jiangsu Chaben Medical Healthcare Technology Co.,Ltd Room 402, No.166 North Street, Nantong, Jiangsu, China 226001

E-mail: info@lanlijen.com www.lanlijen.com Tel:86-513-89072216

Made in China

No Animal Testing, No NSAIDs, Ibuprofen, Aspirin or Salicylate

* Trusted by pharmacists, physical therapists, chiropractors, massage therapists and podiatrists







LANLIJEN MENTHOL PAIN RELIEF

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:83050-232

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) MENTHOL 0.04 g in 1 g

Inactive Ingredients	
Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	
WATER (UNII: 059QF0KO0R)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
.ALPHATOCOPHEROL, D- (UNII: N9PR3490H9)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
GLYCERIN (UNII: PDC6A3C0OX)	
FRANKINCENSE (UNII: R9XLF1R1WM)	
ALCOHOL (UNII: 3K9958V90M)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
TROLAMINE (UNII: 903K93S3TK)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:83050-232- 30	90 g in 1 BOTTLE; Type 0: Not a Combination Product	12/15/2022			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part348	12/15/2022				

Labeler - Jiangsu Chaben Medical Healthcare Technology Co.,Ltd (615936219)

Registrant - Jiangsu Chaben Medical Healthcare Technology Co.,Ltd (615936219)

Establishment					
Name	Address	ID/FEI	Business Operations		
Shanghai Chuangshi Medical Technology (Group) Co., Ltd.		546872672	manufacture(83050-232)		

Revised: 12/2022 Jiangsu Chaben Medical Healthcare Technology Co.,Ltd