LAO WEI WOOD LOCK OIL- methyl salicylate, menthol oil RFX Pharmaceutical 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.

RFX Pharmaceutical Co Ltd

Active Ingredients

Methyl Salicylate 5% Menthol 16%

Purpose

Methyl Salicylate	External Analgesic
Menthol	External Analgesic

Uses

Temporarily relieves minor aches and pains of muscles and joints due to: simple backaches, arthritis, strains, bruises and sprains.

Directions

Adults and children 2 years of age and older: apply 3 to 4 drops of medicated oil to the affected area 1 to 2 times daily. Children under 2 years of age: Do not use, consult your physician.

Warnings

For external use only. Do not use on wounds, irritated or damaged skin, otherwise than as directed. When using this product, avoid contact with the eyes or mucous membranes. Do not bandage tightly.

Keep out of reach of children.

Keep out of reach of children to avoid accidental poinsoning. If swallowed, get medical help or contact a Poison Center right away.

Stop use and ask a doctor if

Condition worsens

Symptoms persist for more than 7 days

Symptoms clear up and occur again within a few days

Excessive irritation of the skin develops

Nausea, vomitting, abdominal discomfort, diarrhea, or skin rash occurs

When using for pain of arthritis: pain persits for more than 10 days, redness is present, in conditions affecting children 12 years of age.

Other Information

Keep container tightly closed. Store at 16 to 30°C (69 to 86°F).

Inactive Ingredients

CLOVE OIL, TURPENTINE OIL, SAFFLOWER, CORYDALIS AMBIGUA TUBER, ANGELICA DAHURICA ROOT, SAPOSHNIKOVIA DIVARICATA ROOT, FRANKINCENSE, MYRRH, DIPSACUS ASPER ROOT, SPATHOLOBUS SUBERECTUS STEM, PANAX NOTOGINSENG WHOLE, LIGUSTICUM WALLICHII ROOT

Questions or Comments? (626)401-1866, M-F 9:00 am to 5:00 pm

Manufactured by RFX Pharmaceutical Co.; Ltd No. 18, Dachongli Qianjian Road, Wuzhou, Guangxi, China

Distributed by WEON PHARMACEUTICAL GROUP COMPANY LIMITED. RM A05017/F SUNRISE IND BLDG, 10 HONG MAN ST CHAI WAN, HONG KONG

Imported by Lucky Mart Inc. 2210 N. Seaman Ave, South El Monte, CA 91733, U.S.A Phone: (626)401-1866

Drug Facts



LAO WEI WOOD LOCK	OIL								
methyl salicylate, menthol oil									
Product Information									
Product T ype	HUMAN OTC DRUG	Item Code (Source) NDC		NDC:76	:76206-301				
Route of Administration	TOPICAL								
Active Ingredient/Active Moiety									
Ingredient Name Basis of Strength					Strength				
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:0414PZ4LPZ) METHYL SALICYLAT									
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) MENTHOL					16 g in 100 mL				
Inactive Ingredients									
Ingredient Name					Strength				

CLOVE OIL (LINII, F							
CLOVE OIL (UNII: 5	578389	D6D0)					
TURPENTINE OIL (TURPENTINE OIL (UNII: C5H0QJ6V7F)						
SAFFLOWER (UNII:	4VBL	71TY4Y)					
CORYDALIS AMBIO	GUA T	UBER (UNII: 1DN1EU584R)					
ANGELICA DAHUR	ICA RO	DOT (UNII: 1V63N2S972)					
SAPO SHNIKO VIA DIVARICATA ROOT (UNII: 8H84LFK2QD)							
FRANKINCENSE (U	NII: R9	XLF1R1WM)					
MYRRH (UNII: JC710	GJ1F3L)					
DIPSACUS ASPER R	ROOT	(UNII: LB1GQP4253)					
SPATHOLOBUS SU	BERE	CTUS STEM (UNII: N51VZ363BA)					
PANAX NOTOGINS	ENG V	VHOLE (UNII: E7XOU43ESD)					
LIGUSTICUM WALLICHII ROOT (UNII: R81AD159QS)							
	Licin						
Packaging							
		Package Description	Marketing Start Date	Marketing End Date			
Packaging		Package Description L in 1 BOTTLE, GLASS; Type 0: Not a Combination	U	U			
 Packaging Item Code NDC:76206-301- 	38 m	Package Description L in 1 BOTTLE, GLASS; Type 0: Not a Combination	U	•			
 Packaging Item Code NDC:76206-301- 	38 m Pro d	Package Description L in 1 BOTTLE, GLASS; Type 0: Not a Combination uct	U	•			
<pre>P>ckaging f Item Code NDC:76206-301- 38</pre>	38 m Prod	Package Description L in 1 BOTTLE, GLASS; Type 0: Not a Combination uct	U	U			
Packaging # Item Code 1 NDC:76206-301- 38	38 m Prod	Package Description L in 1 BOTTLE, GLASS; Type 0: Not a Combination uct	Date	Date			

Labeler - RFX Pharmaceutical Co Ltd (530620871)

Registrant - RFX Pharmaceutical Co Ltd (530620871)

Establishment

Name	Address	ID/FEI	Business Operations
RFX Pharmaceutical Co Ltd		530620871	manufacture(76206-301)

Revised: 8/2015

RFX Pharmaceutical Co Ltd