XJOW- menthol gel Omom Pharmaceuticals, 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.

XJOW PAIN GEL

Drug Facts

Active Ingredients

Menthol USP 1.25%

Purpose

Cooling Pain Reliever

Uses: Temporary relief from minor aches and pains of sore muscles and joints associated with: • arthritis • simple backache • bruises • cramps • muscle strains and sprains.

Warnings: For external use only.

When using this product: • Use only as directed. • Do not bandage tightly or use with heating pad • Avoid contact with the eyes or mucous membranes • Do not apply to wounds or damaged skin • Do not use with other ointments, creams, sprays or liniments • Do not apply to irritated skin or if excessive irritation develops • Wash hands after use with cool water.

Stop use and ask a doctor if: • Condition worsens, or symptoms persist for more than 7 days, or clear up and reoccur within few days. • Redness is present • Irritation develops

If pregnant or breastfeeding: Ask a health professional before use.

Keep out of the reach of children: • If accidentally ingested, get medial help or contact a Poison Control Center immediately.

Directions: Adults and children 12 years of age and older: • Apply to the affected area. • Massage the dispensed gel, into the affected area until thoroughly absorbed. • Use as necessary. • Do not exceed 4 times a day. Children under 12 years of age: Consult physician

Inactive Ingredients: Acanthopanax Senticosus Root Extract, Amber Powder, Angelica Archangelica Root Oil, Carbomer, Carthamus Tinctorius Seed Oil, Cinnamomum Cassia Leaf Oil, Cnidium Officinale Root Extract, Commiphora Myrrha Resin, Corydalis Turtschaninovii Root Extract, Curcuma Longa Root Extract, Daeonnrops Draco Extract, Dipsacus Sylvestris Extract, Dryneria, Ethyl Alcohol, Ethylhexylglycerin, Eucommia Ulmoides Leaf Extract, Gardenia Florida Oil, Ledebouriella Divaricata Root Extract, Olibanum, Orchis Mascula Flower Extract, Paeonia Officinalis Callus Extract, Panax Notoginseng Root Extract, Phenoxyethanol, Prunus Persica Kernel Oil, Rheum Palmatum Root / Stalk Extract, Triethanolamine, Water (Aqua)

Other information: Store in a cool dry place with lid closed tightly.

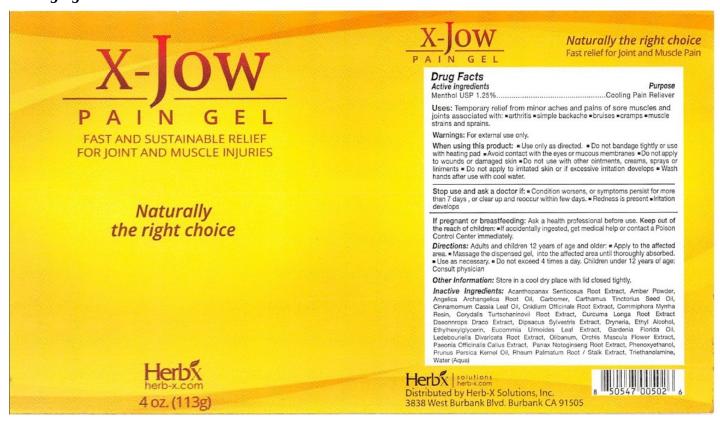
FAST AND SUSTAINABLE RELIEF FOR JOINT AND MUSCLE INJURIES

Naturally the right choice

Distributed by Herb-X Solutions, Inc.

3838 West Burbank Blvd. Burbank CA 91505

Packaging



XJOW

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69934-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10 EIP3A) (MENTHOL - UNII:L7T10 EIP3A)	MENTHOL, UNSPECIFIED FORM	1.25 g in 100 g	

Inactive Ingredients	
Ingredient Name	Strength
ELEUTHERO (UNII: ZQH6VH092Z)	

AMBER (UNII: 70J9Z0J26P)	
ANGELICA ROOT OIL (UNII: B25G881UOX)	
CARBOXYPOLYMETHYLENE (UNII: 0 A5MM307FC)	
SAFFLOWER OIL (UNII: 65UEH262IS)	
CHINESE CINNAMON LEAF OIL (UNII: 4U4V2F2E4Y)	
CNIDIUM OFFICINALE ROOT (UNII: 8S3OZD358J)	
MYRRH (UNII: JC71GJ1F3L)	
CORYDALIS YANHUSUO TUBER (UNII: 0TUP42692Z)	
TURMERIC (UNII: 856 YO1Z64F)	
DAEMONOROPS DRACO RESIN (UNII: 787Z7N9UCU)	
DIPSACUS FULLONUM ROOT (UNII: 4TJV827BQT)	
DRYNARIA FORTUNEI ROOT (UNII: 731W842X8Q)	
ALCOHOL (UNII: 3K9958V90M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EUCOMMIA ULMO IDES LEAF (UNII: AM272O881C)	
GARDENIA JASMINO IDES FRUIT (UNII: 7CTH8 MD549)	
SAPO SHNIKO VIA DIVARICATA ROOT (UNII: 8 H8 4LFK2QD)	
FRANKINCENSE (UNII: R9 XLF1R1WM)	
ORCHIS MASCULA FLOWER (UNII: 6H1JQK35LA)	
PAEONIA OFFICINALIS ROOT (UNII: 8R564U2E1P)	
PANAX NOTOGINSENG ROOT (UNII: GQX1C1175U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PRUNUS PERSICA SEED (UNII: V9 C8 1470 RR)	
RHEUM PALMATUM ROOT (UNII: G025DAL7CE)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

ı	Packaging	ackaging		
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:69934-001- 04	113 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 1/15/20 17	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	0 1/15/20 17	

Labeler - Omom Pharmaceuticals, Inc. (079869885)

Revised: 1/2017 Omom Pharmaceuticals, Inc.