

FIBRO FLEX- magnesia muriatica spray
MAGNESIUM DIRECT INC.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Fibro Flex

Drug Facts

Active Ingredients

Magnesia Muriatica 1X HPUS

Purpose

Joint and soft tissue pain

Keep out of reach of children.

Uses Temporarily relieves the symptoms of Fibromyalgia pain.

Warnings FOR EXTERNAL USE ONLY

If pregnant or breast feeding, ask a health professional before use. Consult a physician if symptoms persist for more than 7 days or worsen. **Stop use if:** Excessive irritation or burning of skin develops.

Directions Apply 4-8 sprays to areas with Fibromyalgia pain on the body. Rub thoroughly. For convenience, spray into hand first, then apply. May be used multiple times daily. **DO NOT** use more than 100 sprays in a 24 hour period.

Inactive Ingredients: water, calcium chloride, trace minerals.

Questions? 1-888-249-8574 M-F, 9-5, EST

Distributed by: Magnesium Direct

Alpharetta, GA 30005

WWW.FIBRO-FLEX.COM

Packaging

Stay Active
with Fibro Flex

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FF071501 NO EXPIRATION DATE NDC 61495-401-04

Fibro Flex™

Fibromyalgia
Gently Calms Pain
Improves Mobility

Stay Active

ODORLESS
CLEAR
LOTION

Homeopathic
4 fl. oz. (118mL)

FIBRO FLEX

magnesia muriatica spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61495-401
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MAGNESIUM CHLORIDE (UNII: 02F3473H9O) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM CHLORIDE	1 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61495-401-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/11/2015	

Labeler - MAGNESIUM DIRECT INC. (019855198)**Establishment**

Name	Address	ID/FEI	Business Operations
MAGNESIUM DIRECT INC.		019855198	manufacture(61495-401)

Revised: 7/2015

MAGNESIUM DIRECT INC.