THERAWORX RELIEF FOR FOOT CRAMPS ROLL ON- magnesium sulfate cream AVADIM HOLDINGS, 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.

Theraworx Relief for Foot Cramps Roll On

Drug Facts

Active Ingredient

Contains Magnesium Sulfate [(Magnesia sulphurica) 6X 0.05% HPUS The letters H.P.U.S. indicate that the components in this product are officially monographed in the Homepathic Pharmacopoeia of the United States.

Purpose

Muscle Soreness Relief

Uses

• prevents cramps and spasms • releases muscle tightness • relieves muscle soreness

Warnings

For external use only. If eye contact occurs, rinse thoroughly with water.

When using this product

• avoid eye contact • store between 32°F and 120°F • use only as directed • not for ingestion

Stop use and ask a doctor if

unintended effects occur.

If pregnant or breastfeeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

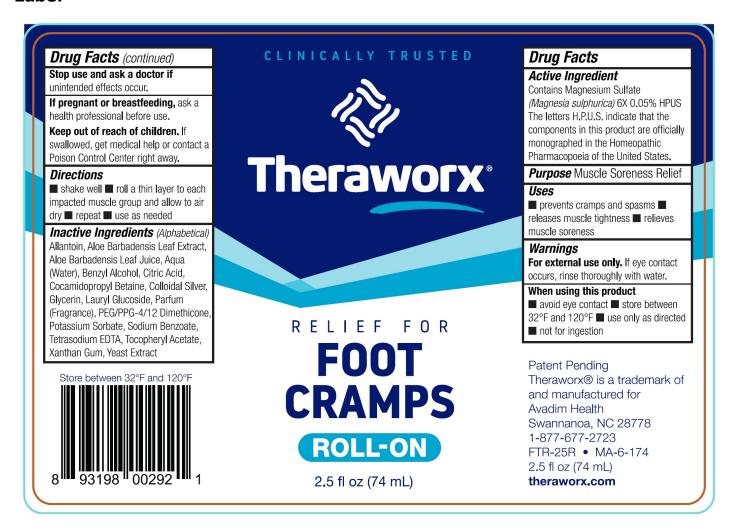
Directions

• shake well • roll a thin layer to each impacted muscle group and allow to air dry •

Inactive Ingredients (Alphabetical)

Allantoin, Aloe Barbadensis Leaf Extract, Aloe Barbadensis Leaf Juice, Aqua (Water), Benzyl Alcohol, Citric Acid, Cocamidopropyl Betaine, Colloidal Silver, Glycerin, Lauryl Glucoside, Parfum (Fragrance), PEG/PPG-4/12 Dimethicone, Potassium Sorbate, Sodium Benzoate, Tetrasodium EDTA, Tocopheryl Acetate, Xanthan Gum, Yeast Extract

Label



THERAWORX RELIEF FOR FOOT CRAMPS ROLL ON magnesium sulfate cream Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration TOPICAL Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM SULFATE, UNSPECIFIED FORM	6 [hp_X] in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
ALLANTOIN (UNII: 344S277G0Z)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
WATER (UNII: 059QF0KO0R)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)		
SILVER (UNII: 3M4G523W1G)		
GLYCERIN (UNII: PDC6A3C0OX)		
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)		
PEG/PPG-4/12 DIMETHICONE (UNII: JAN3585W85)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
EDETATE SODIUM (UNII: MP1J8420LU)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
XANTHAN GUM (UNII: TTV12P4NEE)		
YEAST, UNSPECIFIED (UNII: 3NY3SM6B8U)		

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:61594-041- 00	1 in 1 CARTON	04/01/2024			
1		74 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		04/01/2024			

Labeler - AVADIM HOLDINGS, INC. (118512488)

Revised: 1/2024 AVADIM HOLDINGS, INC.