# THERAWORX RELIEF MUSCLE CRAMP AND SPASM RELIEF- magnesium sulfate heptahydrate spray, metered 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.

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## Theraworx Relief Muscle Cramp and Spasm Relief Spray

## **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 Homeopathic Pharmacopoeia of the United States.

### Purpose

Muscle Soreness Relief

#### Uses

- prevents muscle cramps
- releases muscle tightness
- reduces muscle soreness

## Warnings

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

## When using this product

• avoid eye contact • do not store at temperature above 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.

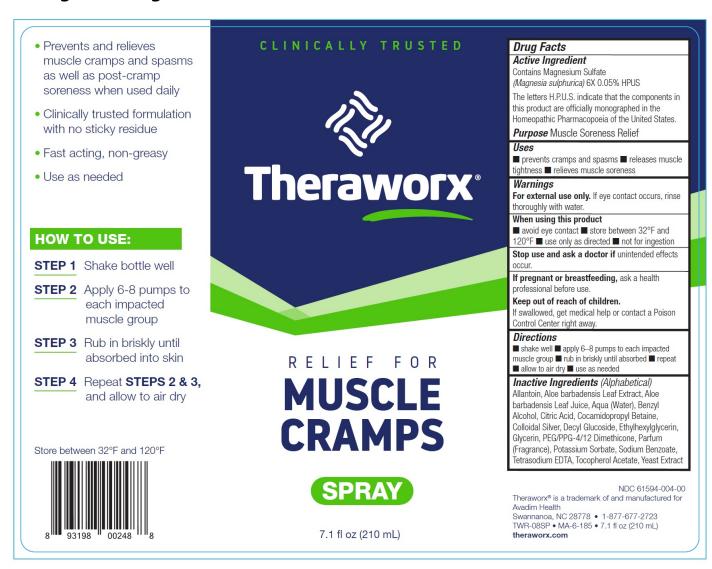
#### **Directions**

• shake well • apply 6-8 pumps onto entire muscle group(s) or area • briskly rub in until absorbed • use as needed • allow to air dry

## Inactive Ingredients

Aqua (Water), Cocamidopropyl Betaine, Aloe Barbadensis Leaf Juice, Colloidal Silver, Tocopheryl Acetate, Glycerin, Allantoin, Beta Glucan, Citrus Paradisi (Grapefruit) Fruit Extract, Lauryl Glucoside, Tetrasodium EDTA, PEG/PPG-4/12 Dimethicone, Methylparaben, Propylparaben, Parfum (Fragrance)

### Package Labeling:



#### THERAWORX RELIEF MUSCLE CRAMP AND SPASM RELIEF

magnesium sulfate heptahydrate spray, metered

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Г	100	JUCL		OI I	нa	LIUII

Product Type HUMAN OTC DRUG Item Code (Source) NDC:61594-004

Ingredient Name	<b>Basis of Strength</b>	Strength
	MAGNESIUM SULFATE HEPTAHYDRATE	6 [hp_X] in 210 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)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG/PPG-4/12 DIMETHICONE (UNII: JAN3585W85)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
EDETATE SODIUM (UNII: MP1J8420LU)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
YEAST, UNSPECIFIED (UNII: 3NY3SM6B8U)	

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:61594- 004-00	210 mL in 1 BOTTLE; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)	07/22/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/22/2017	

# Labeler - AVADIM HOLDINGS, INC (118512488)

## **Registrant - AVADIM HOLDINGS, INC (118512488)**

Revised: 6/2024 AVADIM HOLDINGS, INC