DR. SPENSERS NATURAL HERBAL PAIN RELIEF - menthol cream
LTC Health Corp

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.

Active Ingredient

Menthol (Mentha piperita 3.5%, from leaf and aerial parts)

Purpose

Natural Herbal Pain Relief Cream

Keep out of reach of children

Keep out of reach of children.

Indications:

For the temporary relief of aches and pains of muscles and joints associated with arthritis, strains, sprains, bruises and backache.

Warnings:

For external use ONLY. If rash or irritation occurs, discontinue use. The application of eternal heat such as an electric heating pad may result in excessive skin irritation or skin burn. Avoid contact with eyes, mucous membranes, and damaged skin or open wound. If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a physician.

Do not use

if you are pregnant or breast feeding.

Directions:

Adults and children 12 years of age and older: Apply and massage a small amount 3 times over the area of pain. Repeat 3 to 4 times daily.

Non-medicinal Ingredients:

purified water, herbal extracts of : ledebouriella root, sophora root, acanthopanax root, knotweed root, chebula, dandelion, isopropyl alcohol, sunflower oil, emulsifying wax, ethoxydiglycol, isopropyl myristate, carbomer, optiphen plus, phenoxyethanol, BHT, sodium hydroxide

Dr. Spenser's Product Label

Dr. Spenser's™

Natural Herbal Pain Relief Cream

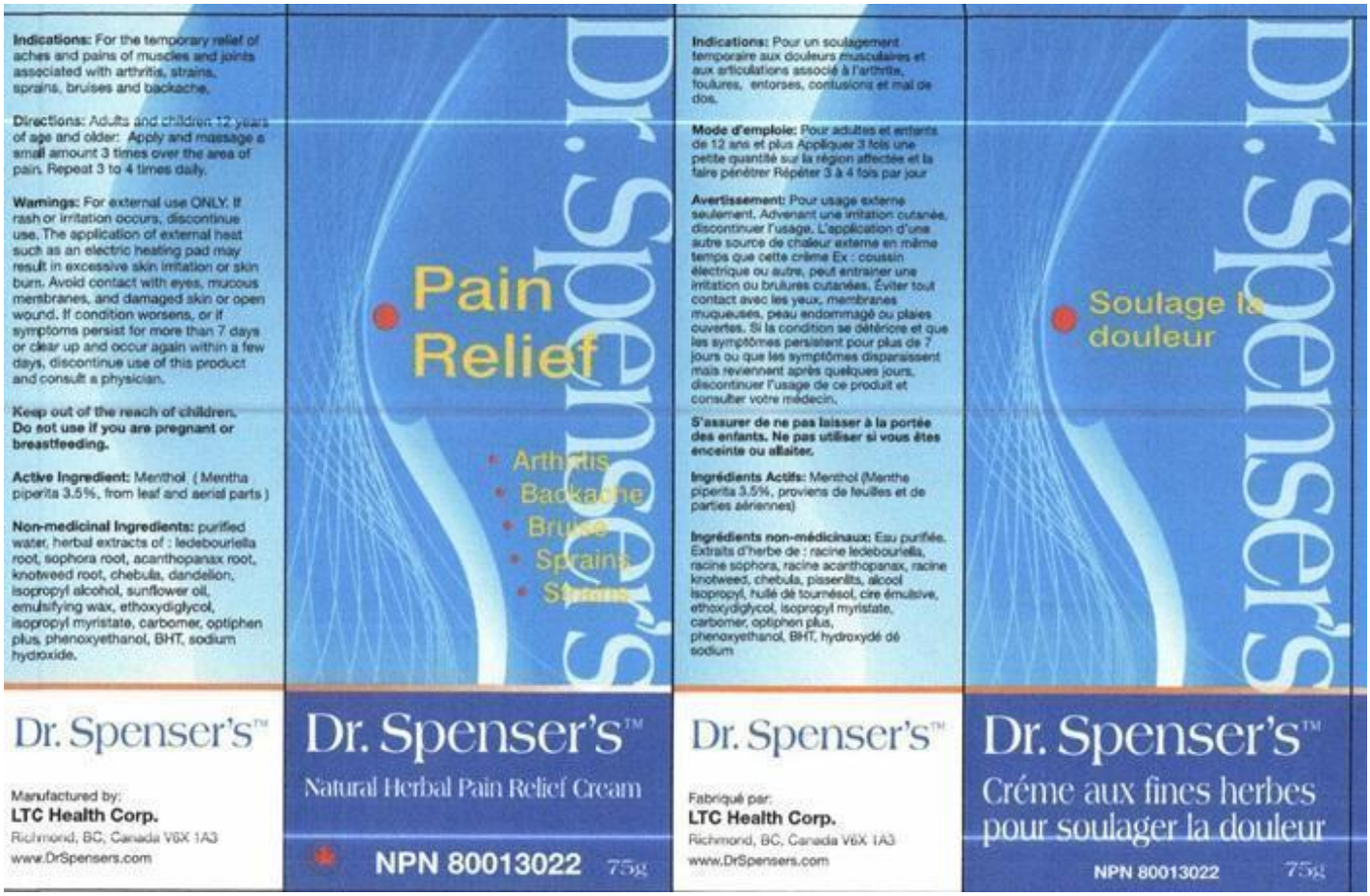
Pain Relief

Arthritis
Backache
Bruise
Sprains
Strains

NPN 80013022 75g

Manufactured by:
LTC Health Corp.
Richmond, BC, Canada V6X 1A3

www.DrSpensers.com



DR. SPENSERS NATURAL HERBAL PAIN RELIEF

menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69562-022
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	2.625 g in 75 g

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SAPOSHNIKOVIA DIVARICATA ROOT (UNII: 8H84LFK2QD)	
SOPHORA FLAVESCENS ROOT (UNII: IYR6K8KQ5K)	
ELEUTHERO (UNII: ZQH6VH092Z)	
PERSICARIA TINCTORIA LEAF (UNII: FU6582QMPV)	
TERMINALIA CHEBULA FRUIT (UNII: S8R4V700NK)	
TARAXACUM OFFICINALE (UNII: 39981FM375)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL SUCROSE CROSSLINKED) (UNII: Z135WT9208)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PEG-150 STEARATE (UNII: 7BSG7DF10Q)	
STEARETH-20 (UNII: L0Q8IK9E08)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69562-022-01	75 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part356	11/01/2009	

Labeler - LTC Health Corp (202782066)

Registrant - LTC Health Corp (202782066)

Establishment

Name	Address	ID/FEI	Business Operations
LTC Health Corp		202782066	manufacture(69562-022)

Revised: 1/2015 LTC Health Corp