KT RECOVERY PLUS PAIN RELIEF GEL- camphor, menthol gel KT Health LLC

KT Recovery Plus

Active Ingredients Purpose

Camphor (5%) External Analgesic

Menthol (5%) External Analgesic

Uses:

for the temporary relief of minor aches and pains of muscles and joints due to:

* simple backache * arthritis * sprains * strains * bruises

Warnings:

* for external use only * do not apply to wounds or damaged skin or bandage tightly * avoid contact with eyes * Keep out of reach of children. if swallowed, get medical help or contact a Poison Control Center immediately * 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 * Pregnancy-breast feeding warning: if pregnant or breast feeding, ask a health professional before use * do not bandage tightly

Inactive Ingredients:

distilled water, hamamelis virginiana (witch hazel extract),

ethanol alcohol, arnica Montana, oleyl alcohol and

zanthoxylum alatum (scezhuan pepper), Bio-saccharide

Gum -1, glycerin, ammonium acryloyldimethyltaurate/VP

copolymer, beta cyclodextrin, menthol, potassium hydroxide,

mannitol, cellulose chromium, hydroxide green, tocopheryl

acetate (vitamin E), hydroxypropyl methycellulose

Directions:

for adults and children 2 years and older; apply to affected area not more than 3 to 4 times daily. "Children under 2 years of age: consult a doctor." Apply generously to affected and surrounding areas. Rub in well. Use 1 application for minor pain, 2 for medium and 3 applications for severs symptoms. Allow to dry between applications

(usually just 2-3 minutes).

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An image/jpeg of the container label is included in this section.

KT RECOVERY PLUS PAIN RELIEF GEL

camphor, menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73044-102(NDC:10842-102)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	50 mg in 1 mL	
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	50 mg in 1 mL	

Inactive Ingredients				
Ingredient Name	Strength			
ZANTHOXYLUM BUNGEANUM FRUIT (UNII: 3CIP16A418)				
OLEYL ALCOHOL (UNII: 172F2WN8DV)				
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
MANNITOL (UNII: 30WL53L36A)				

EUPHORBIA ANTISYPHILITICA WHOLE (UNII: 82A88H0RIQ)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
MODIFIED CORN STARCH (1-OCTENYL SUCCINIC ANHYDRIDE) (UNII: 461P5CJN6T)	
ARNICA MONTANA (UNII: O80TY208ZW)	
WATER (UNII: 059QF0KO0R)	
WITCH HAZEL (UNII: 101I4J0U34)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYGLYCERYL-10 CAPRYLATE (UNII: YS396CQX5C)	
BIOSACCHARIDE GUM-1 (UNII: BB4PU4V09H)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73044- 102-03	5 mL in 1 PACKET; Type 0: Not a Combination Product	05/01/2019	
2	NDC:73044- 102-01	100 mL in 1 TUBE; Type 0: Not a Combination Product	05/01/2019	
3	NDC:73044- 102-02	89 mL in 1 BOTTLE, WTH APPLICATOR; Type 0: Not a Combination Product	05/01/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/01/2019	

Labeler - KT Health LLC (807008037)

Establishment				
Name	Address	ID/FEI	Business Operations	
Dynamic Blending Specialists		085704438	manufacture(73044-102)	

Establishment			
Name	Address	ID/FEI	Business Operations
United Laboratories Manufacturing, LLC		807878116	manufacture(73044-102) , relabel(73044-102)

Revised: 11/2024 KT Health LLC