ARTHRITIS WONDER- menthol cream
G2 Products LLC

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.

Arthritis Wonder

Active ingredient

Menthol 1.25%

Purpose

External analgesic

Uses

For the temporary relief of minor aches and pains of muscles and joints associated with ● arthritis ● simple backache ● strains ● bruises ● sprains

Warnings

For external use only.

Flammable: Keep away from fire or flame.

When using this product

• avoid contact with the eyes • do not bandage tightly

Do not use

● on wounds or damaged skin with a heating pad

Stop use and ask a doctor if

• condition worsens • symptoms persist for more than 7 days • symptoms clear up and occur again within a few days

Keep out of the reach of children

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

Directions

• Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily • Children under 2 years of age: consult a doctor

Inactive ingredients:

Alcohol, Caprylic/Capric Triglyceride, Caprylyl Glycol, Cetearyl Alcohol, Cetyl Alcohol, Dimethicone, Ethoxydiglycol, Ethylhexylglycerin, Glyceryl Oleate, Glyceryl Stearate, Hexylene Glycol, Lecithin, Mentha Piperita (Peppermint) Oil, Phenoxyethanol, Potassium Olivoyl Hydrolyzed Oat Protein, Propanediol, Pullulan, Sclerotium Gum, Sodium Phytate, Water, Wogonin, Xanthan Gum.

Principal Display Panel

ARTHRITIS WONDER

menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71893-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.4125 g in 113 g

Inactive Ingredients	
Ingredient Name	**Strength**
DIMETHICONE (UNII: 92RU3N3Y1O)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
PROPANEDIOL (UNII: 5965N8W85T)	
PULLULAN (UNII: 8ZQ0AYU1TT)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
WATER (UNII: 059QF0KO0R)	
WOGONIN (UNII: POK93PO28W)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALCOHOL (UNII: 3K9958V90M)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CLAVICEPS PURPUREA SCLEROTIUM (UNII: 01G9XEA93N)	
HEXASODIUM PHYTATE (UNII: ZBX50UG81V)	

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:71893-101-20	85 g in 1 APPLICATOR; Type 0: Not a Combination Product	04/26/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	04/06/2020	

Labeler - G2 Products LLC (080963729)

Revised: 3/2020 G2 Products LLC