

**SUPER WHITE STUFF PAIN RELIEF- menthol cream
BLUE SPRING WELLNESS**

SUPER WHITE STUFF Pain Relief Cream

Active Ingredient:

Menthol, USP (1.4%)

Purpose:

Topical Analgesic

Uses:

Temporary relief minor aches, pains associated with arthritis, simple backache, sprains, strains, bruises.

Warnings:

For external use only.

Do not use

on wounded, damaged or irritated skin.

When using this product

avoid contact with eyes or mucous membranes, do not bandage tightly.

Stop use and ask a doctor if

you experience a rash and/or a reaction, condition worsens, or if symptoms persist for more than 10 days or clear up and occur again within few days.

If Pregnant or Breast-Feeding

ask a health professional before use. **Consult a doctor** before use on children under 12 if arthritis conditions are present. For all other listed uses, consult a doctor before use on children under 2.

Keep out of reach of children.

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

Directions:

Apply to affected area not more than 3 or 4 times a day.

Inactive Ingredients:

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Anthemis Nobilis (Roman Chamomile) Oil, Aqua (Deionized Water), Argania Spinosa (Argan) Oil, Arnica Montana Flower Extract, Ascorbyl Palmitate (Vitamin C), Calendula Officinalis Extract, Caprylyl Glycol, Centaurea Cyanus (Cornflower) Extract, Chamomilla Recutita (Chamomile) Extract, Coriandrum Sativum (Coriander) Oil, Emu Oil, Glycerin, Hamamelis Virginiana (Witch Hazel), Lamium Album (White Nettle) Extract, Methylsulfonylmethane (MSM), Phenoxyethanol, Salix Nigra (Willow) Bark Extract, Sodium Cocoyl Isethionate, Sodium Hydroxide, Sorbitol, Tanacetum Parthenium (Feverfew) Extract, Tetrasodium EDTA, Tilia Cordata (Linden) Extract, Tocopheryl Acetate (Vitamin E), Vitis Vinifera (Grape) Seed Extract.

Package Labeling:



SUPER WHITE STUFF PAIN RELIEF

menthol cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:14448-445
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	14 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARGAN OIL (UNII: 4V59G5UW9X)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CHAMOMILE (UNII: FGL3685T2X)	
CORIANDER (UNII: 1OV56052IK)	

EMU OIL (UNII: 344821WD61)
GLYCERIN (UNII: PDC6A3C0OX)
WITCH HAZEL (UNII: 101I4J0U34)
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
SALIX NIGRA BARK (UNII: QU52J3A5B3)
SODIUM COCOYL ISETHIONATE (UNII: 518XTE8493)
SODIUM HYDROXIDE (UNII: 55X04QC32I)
SORBITOL (UNII: 506T60A25R)
FEVERFEW (UNII: Z64FK7P217)
EDETATE SODIUM (UNII: MP1J8420LU)
TILIA CORDATA WHOLE (UNII: W5E5UB44GD)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
VITIS VINIFERA SEED (UNII: C34U15ICXA)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:14448-445-00	118 mL in 1 JAR; Type 0: Not a Combination Product	03/04/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/04/2024	

Labeler - BLUE SPRING WELLNESS (182950118)

Revised: 1/2024

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