

**SUPER WHITE STUFF PAIN RELIEF- menthol cream**  
**BLUE SPRING WELLNESS, L.L.C.**

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**SUPER WHITE Stuff Pain Relief Cream**

***Drug Facts***

***Active Ingredient***

Menthol, USP 1.4%

***Purpose***

Topical Analgesic

***Uses***

For the temporary relief of minor aches and 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 clears up and occur again within a few days.

**If Pregnant or Breast-Feeding**

ask a health professional before use.

**Consult a doctor**

before use on children under 12 years of age if arthritis conditions are present. For all

other listed uses, consult a doctor before use on children under 2 years of age.

**Keep out of reach of children.**

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

***Other information***

Store at room temperature.

***Directions***

Apply to affected area not more than 3 to 4 times a days. before using for children under 12 for arthritic conditions and for all children under 2 years of age for all other uses. **Consult a doctor**

***Inactive Ingredients***

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

***Questions or Comments?***

Call or visit [www.bluestuff.com](http://www.bluestuff.com) **1-800-452-3700**

**Package Labeling:355**

BLUESPRING

# Super White Stuff<sup>OTC</sup> pain relief cream

No Added Color or Fragrance

Natural Pain Relief

- Arthritis
- Joint Pain
- Backaches
- Muscle Pain
- Sprains & Strains
- Bruises



Made in the USA

12 fl. oz. (355 ml.)

NDC # 14448-307-50



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Oklahoma City, OK 73157

LB-003-200619

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## SUPER WHITE STUFF PAIN RELIEF

menthol cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:14448-307
<b>Route of Administration</b>	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
WATER (UNII: 059QF0KO0R)	

<b>ARNICA MONTANA FLOWER</b> (UNII: OZ0E5Y15PZ)
<b>ASCORBYL PALMITATE</b> (UNII: QN83US2B0N)
<b>CALENDULA OFFICINALIS FLOWER</b> (UNII: P0M7O4Y7YD)
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)
<b>CHAMOMILE</b> (UNII: FGL3685T2X)
<b>TANACETUM PARTHENIUM WHOLE</b> (UNII: 6GE7Z0761K)
<b>CORIANDER</b> (UNII: 10V56052IK)
<b>EMU OIL</b> (UNII: 344821WD61)
<b>HAMAMELIS VIRGINIANA ROOT BARK/STEM BARK</b> (UNII: T7S323PKJS)
<b>LAMIUM ALBUM WHOLE</b> (UNII: 046Y1357I6)
<b>DIMETHYL SULFONE</b> (UNII: 9H4PO4Z4FT)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)
<b>SALIX NIGRA BARK</b> (UNII: QU52J3A5B3)
<b>SODIUM COCOYL ISETHIONATE</b> (UNII: 518XTE8493)
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)
<b>SORBITOL</b> (UNII: 506T60A25R)
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)
<b>TILIA CORDATA WHOLE</b> (UNII: W5E5UB44GD)
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)
<b>VITIS VINIFERA SEED</b> (UNII: C34U15ICXA)
<b>CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)</b> (UNII: 59TL3WG5CO)
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:14448-307-50	335 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/01/2020	

**Labeler** - BLUE SPRING WELLNESS, L.L.C. (182950118)

Revised: 11/2023

BLUE SPRING WELLNESS, L.L.C.