GREEN COMPASS REDUCE TOPICAL PAIN- camphor, menthol, methyl salicylate cream Green Compass, Inc.

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.

Green Compass Reduce Topical Pain Cream

Drug Facts

Active ingredients

Camphor 3%

Menthol 10%

Methyl Salicylate 10%

Purpose

Topical Analgesic

Topical Analgesic

Topical Analgesic

Uses

For temporary relief of minor aches and pains of muscles and joints.

Warnings

For external use only.

Do not use

- on wounds or damaged skin
- with a heating pad

When using this product

- avoid contact with the eyes
- do not bandage tightly

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days
- condition clears up and occurs again within a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 12 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 12 years of age: ask a doctor

Inactive ingredients

butyrospermum parkii (shea) butter, caprylhydroxamid acid, caprylic capric triglyceride, CBD hemp extract, cetearyl alcohol, cetearyl glucoside, citric acid, glycerin, glyceryl caprylate, glyceryl stearate citrate, organic aloe barbadensis (aloe vera) leaf juice, organic arnica extract, organic cinnamon leaf essential oil, organic hemp oil, organic horsetail oil, organic rosmarinum officinalis (rosemary) leaf extract, organic tapioca starch modified, sodium ascorbyl phosphate, turmeric extract, water (aqua), xanthan gum.

LEARN MORE AT

GreenCompassGlobal.com

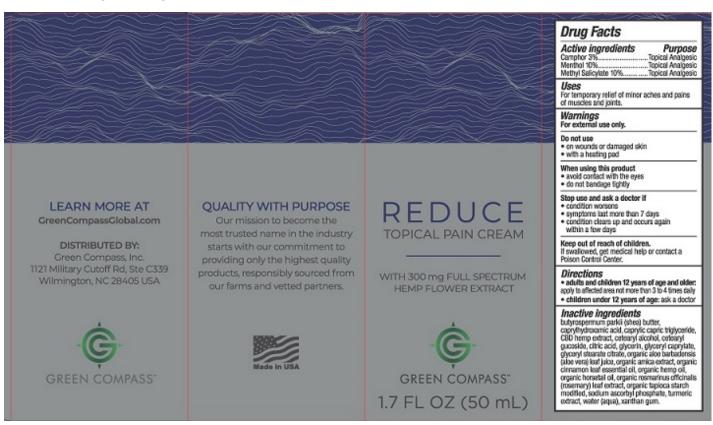
DISTRIBUTED BY:

Green Compass, Inc. 1121 Military Cutoff Rd., Ste C339 Wilmington, NC 28405 USA

REDUCE

TOPICAL PAIN CREAM
WITH 300 mg FULL SPECTRUM HEMP FLOWER EXTRACT
GREEN COMPASS

1.7 FL OZ (50 mL)



GREEN COMPASS REDUCE TOPICAL PAIN

camphor, menthol, methyl salicylate cream

Product	Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73132-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	3 g in 100 mL			
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	10 g in 100 mL			
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:0414PZ4LPZ)	METHYL SALICYLATE	10 g in 100 mL			

Ingredient Name ANTHAN GUM (UNII: TTV12P4NEE) RNICA MONTANA (UNII: O80TY208ZW) ANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S) LYCERYL STEARATE CITRATE (UNII: WH8T92A065) INNAMON LEAF OIL (UNII: S92U8SQ71V) QUISETUM ARVENSE WHOLE (UNII: 73DM367W4P) URMERIC (UNII: 856Y01Z64F) EDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) LYCERIN (UNII: PDC6A3C0OX)	gth
RNICA MONTANA (UNII: O80TY208ZW) ANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S) LYCERYL STEARATE CITRATE (UNII: WH8T92A065) INNAMON LEAF OIL (UNII: S92U8SQ71V) QUISETUM ARVENSE WHOLE (UNII: 73DM367W4P) URMERIC (UNII: 856Y01Z64F) EDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S) LYCERYL STEARATE CITRATE (UNII: WH8T92A065) INNAMON LEAF OIL (UNII: S92U8SQ71V) QUISETUM ARVENSE WHOLE (UNII: 73DM367W4P) URMERIC (UNII: 856YO1Z64F) EDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
LYCERYL STEARATE CITRATE (UNII: WH8T92A065) INNAMON LEAF OIL (UNII: S92U8SQ71V) QUISETUM ARVENSE WHOLE (UNII: 73DM367W4P) URMERIC (UNII: 856YO1Z64F) IEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
INNAMON LEAF OIL (UNII: S92U8SQ71V) QUISETUM ARVENSE WHOLE (UNII: 73DM367W4P) URMERIC (UNII: 856YO1Z64F) IEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
QUISETUM ARVENSE WHOLE (UNII: 73DM367W4P) URMERIC (UNII: 856YO1Z64F) IEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
URMERIC (UNII: 856YO1Z64F) EDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EMP (UNII: TD1MUT01Q7) ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ETEARYL GLUCOSIDE (UNII: 09FUA47KNA) ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
, , ,	
LYCERIN (UNII: PDC6A3C0OX)	
LYCERYL CAPRYLATE (UNII: TM2TZ D4G4A)	
HEA BUTTER (UNII: K49155WL9Y)	
APRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
LOE VERA LEAF (UNII: ZY81Z83H0X)	
OSEMARY (UNII: IJ67X351P9)	
TARCH, TAPIOCA (UNII: 24SC3U704I)	
ODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	
VATER (UNII: 059QF0KO0R)	

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:73132-001- 50	1 in 1 CARTON	07/24/2019			
1		50 mL in 1 BOTTLE; 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	part348	07/24/2019			

Labeler - Green Compass, Inc. (105963322)

Revised: 1/2023 Green Compass, Inc.