RELIEVEIT FOOTSPRAY TOPICAL ANALGESIC- menthol 8% liquid Cosmetic Specialty labs, 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.

RelieveIt Footspray

Drug Facts

Active Ingredient

Menthol 8.0%

Purpose

External analgesic

Uses

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

Warnings

for external use only

When using this product: use only as directed• avoid contact with eyes or mucous membrane• do not apply to open wounds, damaged or very sensitive skin • Do not use if you are allergic to any of this product's Active or Inactive ingredients

If pregnant or breast-feeding, ask a health professional before use.

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

Directions

- Spray balls of feet. arches, heels, toes and legs on bare skin before starting your day or activity:
- 6 8 sprays each foot Apply generously when wearing stockings; 8+ sprays each foot
- Let dry completely Apply again as needed for extended relief

Children under 2: consult a doctor before use

Inactive ingredients

Arnica Montona, Cyclomethicone, Ethyl Alcohol. Ethoxydiglycol. Eucalyptus Oil. Horpogophytum. lsopropyl Palmitate, Lemon Oil, Lemongrass Oil, Pinus Resin, Propylene Glycol, Peppermint Oil, Teo Tree Oil, Wintergreen Oil

Manufactured for: SprainGo, LLC

3802 Spectrum Blvd, Suite 112E

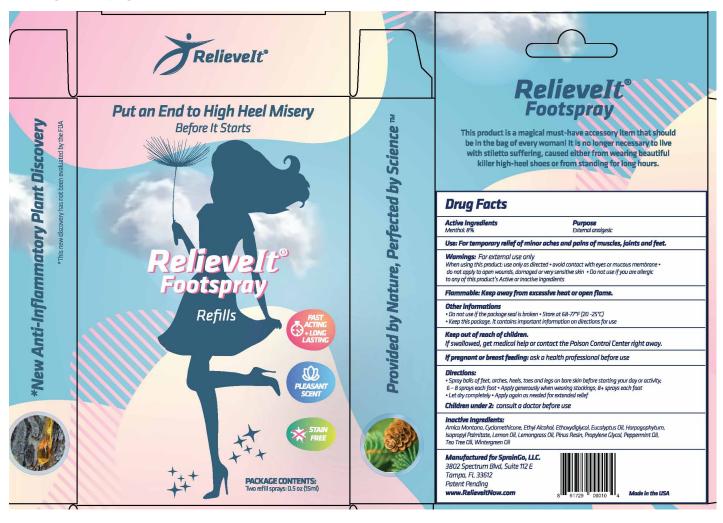
Tampa, FL 33612

Made in the USA

Other Information

- Do not use if the package seal is broken• Store at 68-77°F (20 -25°)
- Keep this package. It contains important information on directions for use

PRINCIPAL DISPLAY PANEL





RELIEVEIT FOOTSPRAY TOPICAL ANALGESIC

menthol 8% liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:58133-956

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.2 mg in 15 mL

Inactive Ingredients	
Ingredient Name	Strength
METHYL SALICYLATE (UNII: LAV5U5022Y)	
CYCLOMETHICONE (UNII: NMQ347994Z)	
BIS-ETHO XYDIGLYCOL CYCLO HEXANE 1,4-DICARBO XYLATE (UNII: R6 MU0 U9 L45)	
HARPAGOPHYTUM PROCUMBENS ROOT (UNII: 10 YM338 E89)	

ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
LEMON OIL (UNII: 19 GRO 8 2 4 LL)	
EAST INDIAN LEMONGRASS OIL (UNII: UP0 M8 M3 VZW)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
TEA TREE OIL (UNII: VIF565UC2G)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
1-(3-AMINO-4-HYDROXYPHENYL)-2-((2-(4-METHOXYPHENYL)-1-METHYLETHYL)AMINO)ETHANOL~(UNII: GJY78AX70Q)	
PINUS RESINOSA WHOLE (UNII: AS8B4M6X9N)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
PROPYLENE GLYCOL 1-ALLYL ETHER (UNII: QRB8092KPK)	

	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:58133-956-	30 mL in 1 CARTON; Type 0: Not a Combination Product	03/13/2020	
ı	2 NDC:58133-956-	2 30 mL in 1 CARTON; Type 0: Not a Combination Product	03/13/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	03/13/2020	

Labeler - Cosmetic Specialty labs, Inc. (032973000)

Registrant - Cosmetic Specialty labs, Inc. (032973000)

Establishment				
Name	Address	ID/FEI	Business Operations	
Cosmetic Specialty labs, Inc.		032973000	manufacture(58133-956)	

Revised: 3/2020 Cosmetic Specialty labs, Inc.