NEUTROGENA SUN RESCUE AFTER SUN MEDICATED RELIEF- camphor (synthetic) gel
Kenvue Brands LLC

Neutrogena Sun Rescue After Sun Medicated Relief Gel

Drug Facts

Active ingredient

Camphor 0.45%

Purpose

External analgesic

Use

temporarily relieves pain and itching associated with sunburn

Warnings

For external use only.

Flammable: Keep away from fire or flame

When using this product avoid contact with eyes

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occur 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 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Other information

- Store at 20° to 25°C (68° to 77°F)

Inactive ingredients

Water, Alcohol Denat., Triethanolamine, Carbomer, EDTA

Questions?

Call toll-free 800-299-4786 or 215-273-8755 (collect) or visit www.neutrogena.com

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 85 g Tube Label

NEW

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

NEUTROGENA®

SUN

RESCUE

AFTER SUN

MEDICATED RELIEF GEL

CAMPHOR EXTERNAL ANALGESIC

pain relief for sunburn

cools, calms and soothes

skin on contact

NET WT 3.0 OZ (85 g)

NEUTROGENA SUN RESCUE AFTER SUN MEDICATED RELIEF

camphor (synthetic) gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0721
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	4.5 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	

EDETIC ACID (UNII: 9G34HU7RV0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0721-3	85 g in 1 TUBE; Type 0: Not a Combination Product	10/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/01/2021	

Labeler - Kenvue Brands LLC (118772437)

Revised: 11/2024 Kenvue Brands LLC