

RECOVERY ROLL-ON- menthol, camphor gel
PROXIMITY CAPITAL PARTNERS LLC DBA ASUTRA

Recovery Roll-On (Asutra)

Active Ingredients

Menthol 5%

Camphor 5%

Purpose

Menthol 5%.....Topical Analgesic

Camphor 5%.....Topical Analgesic

Uses

For the temporary relief of minor aches and pains of muscles and joints associated with:

- simple backache
- arthritis
- strains
- bruises
- sprains

Warnings

Use only as directed.

For external use only.

When using this product:

- avoid contact with eyes and mucous membranes
- do not apply to wounds or damaged skin
- do not bandage tightly or use with a heating pad, other ointments, creams, sprays, or liniments.

Stop use and ask doctor if:

- condition worsens
- symptoms persist for more than 7 days, or clear up and reoccur again within a few days.

If pregnant or breastfeeding,ask a health professional before use.

Keep out of reach of children.If accidentally ingested, get medical help or contact Poison Control Center immediately.

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:

- Consult a doctor.

Other Information

- Store in a cool, dry place.
- Avoid direct sunlight.
- Tamper-evident for your protection. Use only if safety seal is intact.
- Keep away from excessive heat or flame.

Inactive Ingredients

Water, Prunus Amygdalus Dulcis (Sweet Almond) Oil, Dimethyl Sulfone, Magnesium Chloride, Cetearyl Alcohol, Glyceryl Stearate, Polyglyceryl-6 Palmitate/Succinate, Glycerin, Hydroxyacetophenone, Tocopheryl Acetate, Xanthan Gum, Caprylyl Glycol, 1,2-Hexanediol, Acacia Senegal Gum, Pentaerythrityl Tetra-Di-T-Butyl Hydroxyhydrocinnamate, Magnesium Sulfate, Potassium Chloride, Sodium Chloride, Calcium Chloride

Questions or Comments?

888-819-6472; Monday-Friday, 9am-5pm CT; Report any serious side effects to number above.

NDC: 72683-008-01

Non-toxic

Paraben Free

Asutra*

Recovery Roll-On

5% Menthol + 5% Camphor

Pain Relief Gel

With Revitalizing Magnesium

1.7 fl. oz. | 50 ml.

NDC:
79687-008-01

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Drug Facts (continued)

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Learn more at asutra.com



Pool

Glue area — does not print

Drug Facts (continued)

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Inactive Ingredients

Water, Prunus Amygdalus Dulcis (Sweet Almond) Oil, Dimethyl Sulfoxide, Magnesium Chloride, Cetyl Alcohol, Glyceryl Stearate, Polyglyceryl-6 Palmirate/Succinate, Glycerin, ▶

Drug Facts (continued)

Hydroxyacetophenone, Tocopheryl Acetate, Xanthan Gum, Caprylyl Glycol, 1,2- Hexanediol, Acacia Senegal Gum, Perlaerythryl Tetra-Di-T-Butyl Hydroxyhydrocinnamate, Magnesium Sulfate, Potassium Chloride, Sodium Chloride, Calcium Chloride

Questions or Comments?

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Listed with the FDA
Distributed by Asutra
4158 W. Montrose Ave.
Chicago, IL 60641
Manufactured by Wanadis Kft
Budapest, Hungary

RECOVERY ROLL-ON

menthol, camphor gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72683-008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	5 g in 100 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
POLYGLYCERYL-6 STEARATE (UNII: ETY9Q81E2T)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
ALMOND OIL (UNII: 66YXD4DKO9)	
ACACIA (UNII: 5C5403N26O)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
WATER (UNII: 059QF0KO0R)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72683-008-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/17/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	10/17/2022	

Labeler - PROXIMITY CAPITAL PARTNERS LLC DBA ASUTRA (081214985)

Establishment

Name	Address	ID/FEI	Business Operations
BioLyte Laboratories, LLC		015560564	manufacture(72683-008)

Revised: 6/2025

PROXIMITY CAPITAL PARTNERS LLC DBA ASUTRA