

NEORELIEF FOR MENSTRUAL PAIN AND CRAMPS- homeopathic pain relief gel gel
BioLyte Laboratories, LLC

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Drug Facts Active Ingredients

All actives are HPUS* and 4X potency.

Apis, Arnica, Belladonna, Bryonia, Cimicifuga, Cyclamen, Joneosia, Kali Carb., Liliium, Rhamnus Calif., Sabina

Bellis, Cactus, Caulophyllum, Colocynthis, Gelsemium, Viburnum

Tilia

*HPUS indicates the active ingredients are in the official Homeopathic Pharmacopeia of the United States.

Purpose

Topical Analgesics

Relief of Cramping

Anti-inflammatory

Uses

For temporary relief of menstrual symptoms such as:

Menstrual pain, Cramping

Indications are based on homeopathic materia medica, not clinical tests.

Warnings

For external use only

Do not use on damaged or broken skin.

Keep out of eyes, open wounds, mucous membranes

Warnings

Stop use and ask doctor if:

Condition worsens or persists,

rash develops

Warnings

If pregnant or breastfeeding consult your doctor before use.

Warnings

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

Directions

Women ages 8 years and older, spread thin layer on lower abdomen and lower back.

Do not apply over other topical lotions; they may hinder effectiveness.

Apply as needed for relief.

Other Information

Store in a cool, dry place. Avoid direct sunlight. Tamper-evident for your protection. Use only if safety seal is intact.

Inactive Ingredients

Citric Acid, Fulvic Acid, Glycerin, Potassium Sorbate, Purified Water, Sodium Hydroxide, Ultrez-10 Polymer.

Questions or Comments?

800.538.1455

Monday through Friday, 8 a.m. - 5 p.m. EST or visit us at neorelief.com.

Reporty any serious side effects to the number above.

Additional Label Content

Packaging Recyclable

Made in the USA

Listed with the FDA

neorelief.com

Drug Facts continued peel here

FTC required disclosure: Product efficacy and claims are based on theories of homeopathy that re not supported by scientific evidence and most modern allopathic

medical professionals.

BioLyte Labs Grand Rapids, MI 49534

NEORELIEF for Menstrual Pain and Cramps

NDC 58368-003-03 Patent Pending

Neo Relief

Temporary relief of

Menstrual pain & cramps

Homeopathic pain relief gel

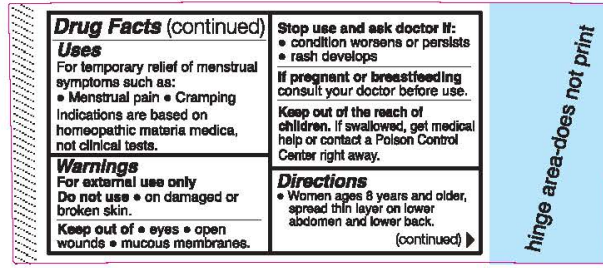
Net Wt. 0.5 oz (14.17g)

FRONT



← Dieline does not print

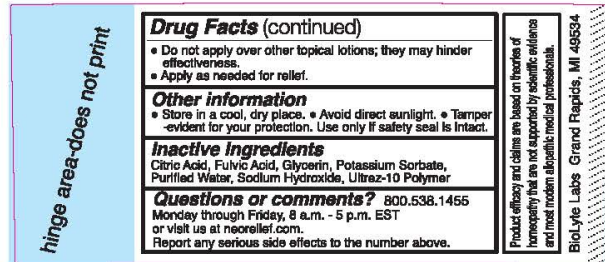
BACK



.125" (seal area does not print)

.75" (hinge area does not print)

BASE



.75" (hinge area does not print)

.125" (seal area does not print)

Job No. 129006
2/26/18



Western Shield Label Company
2146 E. Gladwick Street
Rancho Dominguez, CA 90220
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UNWIND DIRECTION →



PROOF HISTORY

2/13/17 Initial proof
2/26/18 NEW ART Proof

Client: BioLyte Labs
File Name: Neo Relief For Menstrual Pain and Cramps .5oz ECL
Dimension: 1.5711 x 3.6319"
Corner Radius: n/a **Die #:** SP 7064
Facestock: 2.6 mil Pearlescent BOPP
Coating(s): 1.0 mil MATTE BOPP Laminate

Total Plates: 9

PLEASE NOTE WELL: A signed proof is required before your project will be entered into our production schedule. Orders typically ship five (5-7) working days after receipt of an approved proof.

OK TO PRINT REVISE AND RE-PROOF



Copy Position #4

Labels on outside of roll. Left-hand side of design dispenses first.

AUTHORIZED SIGNATURE _____

DATE _____

By signing this proof, I authorize Western Shield to proceed as indicated above. I understand that this proof, when viewed on a computer or printed on a desktop printer, is not an exact representation of the color, size or quality of the actual printed label. I further understand that I am responsible to inform Western Shield of any omissions or errors on this proof, and Western Shield is not responsible for misspellings, compliance to government regulations, or barcode accuracy.

Phone: 310.527.6212 • Fax: 310.327.3871

NEORELIEF FOR MENSTRUAL PAIN AND CRAMPS

homeopathic pain relief gel gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58368-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION - UNII:7UJQ5OPE7D)	POTASSIUM CARBONATE	4 [hp_X] in 1 g
FRANGULA CALIFORNICA BARK (UNII: 1LZ13MQR0S) (FRANGULA CALIFORNICA BARK - UNII:1LZ13MQR0S)	FRANGULA CALIFORNICA BARK	4 [hp_X] in 1 g
ARNICA MONTANA (UNII: O80TY208Z W) (ARNICA MONTANA - UNII:O80TY208Z W)	ARNICA MONTANA	4 [hp_X] in 1 g
ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H)	ATROPA BELLADONNA	4 [hp_X] in 1 g
BRYONIA ALBA ROOT (UNII: T7J046Y12B) (BRYONIA ALBA ROOT - UNII:T7J046Y12B)	BRYONIA ALBA ROOT	4 [hp_X] in 1 g
BLACK COHOSH (UNII: K73E24S6X9) (BLACK COHOSH - UNII:K73E24S6X9)	BLACK COHOSH	4 [hp_X] in 1 g
CYCLAMEN PURPURASCENS TUBER (UNII: G728143D8Q) (CYCLAMEN PURPURASCENS TUBER - UNII:G728143D8Q)	CYCLAMEN PURPURASCENS TUBER	4 [hp_X] in 1 g
SARACA INDICA BARK (UNII: 341ALE870W) (SARACA INDICA BARK - UNII:341ALE870W)	SARACA INDICA BARK	4 [hp_X] in 1 g
LILIUM LANCIFOLIUM WHOLE FLOWERING (UNII: X67Z2963PI) (LILIUM LANCIFOLIUM WHOLE FLOWERING - UNII:X67Z2963PI)	LILIUM LANCIFOLIUM WHOLE FLOWERING	4 [hp_X] in 1 g
JUNIPERUS SABINA LEAFY TWIG (UNII: Z5BEX9K2G1) (JUNIPERUS SABINA LEAFY TWIG - UNII:Z5BEX9K2G1)	JUNIPERUS SABINA LEAFY TWIG	4 [hp_X] in 1 g
BELLIS PERENNIS (UNII: 2HU33I03UY) (BELLIS PERENNIS - UNII:2HU33I03UY)	BELLIS PERENNIS	4 [hp_X] in 1 g
SELENICEREUS GRANDIFLORUS STEM (UNII: 7114SV0MYK) (SELENICEREUS GRANDIFLORUS STEM - UNII:7114SV0MYK)	SELENICEREUS GRANDIFLORUS STEM	4 [hp_X] in 1 g
CAULOPHYLLUM THALICTROIDES ROOT (UNII: JTJ6HH6YEH) (CAULOPHYLLUM THALICTROIDES ROOT - UNII:JTJ6HH6YEH)	CAULOPHYLLUM THALICTROIDES ROOT	4 [hp_X] in 1 g
CITRULLUS COLOCYNTHIS FRUIT PULP (UNII: 23H32AOH17) (CITRULLUS COLOCYNTHIS FRUIT PULP - UNII:23H32AOH17)	CITRULLUS COLOCYNTHIS FRUIT PULP	4 [hp_X] in 1 g
GELSEMIUM SEMPERVIRENS ROOT (UNII: 639KR60Q1Q) (GELSEMIUM SEMPERVIRENS ROOT - UNII:639KR60Q1Q)	GELSEMIUM SEMPERVIRENS ROOT	4 [hp_X] in 1 g
VIBURNUM OPULUS BARK (UNII: T1UG6H6805) (VIBURNUM OPULUS BARK - UNII:T1UG6H6805)	VIBURNUM OPULUS BARK	4 [hp_X] in 1 g
TILIA X EUROPAEA FLOWER (UNII: NHV2K1OUDH) (TILIA X EUROPAEA FLOWER - UNII:NHV2K1OUDH)	TILIA X EUROPAEA FLOWER	4 [hp_X] in 1 g
APIS MELLIFERA (UNII: 7S82P3R43Z) (APIS MELLIFERA - UNII:7S82P3R43Z)	APIS MELLIFERA	4 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM SORBATE (UNII: 1VPU26JZ4)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	

FULVIC ACID (UNII: XII14C5FXV)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58368-003-03	1 in 1 CARTON	04/27/2018	
1		14.17 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		04/27/2018	

Labeler - BioLyte Laboratories, LLC (015560564)

Establishment

Name	Address	ID/FEI	Business Operations
BioLyte Laboratories, LLC		015560564	manufacture(58368-003)

Revised: 1/2024

BioLyte Laboratories, LLC