PHQUIROGEL- camphor, menthol gel Issislen Laboratori S.L.

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.

PH-QUIROGEL+ Pain Relieving Gel

Drug Facts

Active Ingredients

Camphor (3.01%)

Menthol (2.10%)

Purpose

Pain Relieving

Uses

• For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains bruises and sprains.

Warnings

For external use only

Do not use

- on wounds or damaged skin
- with a bandage to cover the surface

When using this product

• avoid contact with the eyes

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days
- symptoms 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 6 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 6 years of age: Consult a doctor.

Inactive Ingredients

ALCOHOL DENAT., BENZYL ALCOHOL, CARBOMER, DISODIUM EDTA, FD&C BLUE NO.1, FD&C YELLOW NO.5, GLYCERIN, GINSENG, PEG-7 GLYCERYL COCOATE,

PHENOXYETHANOL, POTASSIUM SORBATE, SODIUM BENZOATE, SODIUM CHLORIDE, TOCOPHEROL, TRIETHANOLAMINE, WATER.

Package Insert:



Pain Relieving Gel / Gel para Aliviar el Dolor

ENGLISH

Uses: Indicated for the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises and sprains.

Mode of employment: Adults and children 6 years of age and older: Apply to affected area not more than 3 to 4 times daily. Children under 6 years of age: Consult a doctor. After application do not expose treated skin to sunlight.

Warnigs: For external use only. DO NOT USE: on wounds or damaged skin or with a bandage to cover the surface. WHEN USING THIS PRODUCT: avoid contact with the eyes. STOP USE AND ASK A DOCTOR IF: condition worsens, symptoms persist for more than 7 days or symptoms clear up and occur again within a few days. 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. Avoid the solar exhibition after the application. If pregnant or breast-feeding, ask a health professional before use. DO NOT USE: if carton is open or if transparent plastic seal on tube cap is missing or broken.

Active Ingredients: CAMPHOR (3.01%), MENTHOL (2.10%).

Inactive Ingredients: ALCOHOL DENAT., BENZYL ALCOHOL, CARBOMER, DISO-DIUM EDTA, FD&C BLUE NO.1, FD&C YELLOW NO.5, GLYCERIN, GINSENG, PEG-7 GLYCERYL COCOATE, PHENOXYETHANOL, POTASSIUM, SORBATE, SODIUM BENZO-ATE, SODIUM CHLORIDE, TOCOPHEROL, TRIETHANOLAMINE, WATER.

ESPAÑOL

Usos: Indicado para el alivio de dolores y molestias leves de músculos y articulaciones asociados con el dolor de espalda, artritis, torceduras, hematomas y esguinces.

Modo de empleo: Adultos y niños mayores de 6 años: Aplicar sobre el área afectada de 3 a 4 veces al día. Niños menores de 6 años: consultar con un médico.

Después de su aplicación no exponer al sol.

Advertencias: Uso externo. NO USAR: en heridas o sobre piel dañada o con vendajes que cubran la superficie. DURANTE EL USO DEL PRODUCTO: Evitar el contacto con los ojos. DEJAR DE USAR Y CONSULTAR CON MÉDICO SI: el estado empeora, los síntomas persisten más de 7 días o los síntomas mejoran y reaparecen en pocos días. Durante el embarazo o el periodo de lactancia, consultar a su médico antes de usar. MANTENER ALEJADO DEL ALCANCE DE LOS NIÑOS. Si es ingerido, buscar atención médica o contactar con un centro de toxicología inmediatamente. Evitar la exposición solar después de su aplicación. NO USAR SI: la caja está abierta o si el sello de plástico transparente del tapón del tubo está roto.

Ingredientes Activos: ALCANFOR (3,01%), MENTOL (2,10%).

Ingredients Inactivos: AGUA, ALCOHOL BENCÍLICO, ALCOHOL DESNATURALIZADO, BENZOATO DE SODIO, CARBOMER, CLORURO DE SODIO, EDTA DISÓDICO, FD&C AZUL NO.1, FD&C AMARILLO NO.5, FENOXIETANOL, GINSENG, GLICERINA, PEG-7 GLICERIL COCOATE, SORBATO DE POTASIO, TOCOFEROL, TRIETANOLAMINA.

MANUFACTURED FOR / FABRICADO POR: ISSISLEN LABORATORI
JOSEP ANSELM CLAVÉ 47,ESPLUGUES DE LLOBREGAT, BARCELONA (SPAIN)
DISTRIBUTED BY / DISTRIBUÍDO POR: INICIATIC DIGITAL SERVICES
845 N. RAINBOW BLVD, #3102 LAS VEGAS, NV 89107, U.S. Tel: +1 (702) 6050471

MADE IN SPAIN / HECHO EN ESPAÑA

Package Labeling:

PAIN RELIEVING GEL / GEL PARA ALIVIAR EL DOLOR

20 2 Ri Neto eso Ď

Drug Facts

Active Ingredients

Purpose Pain Relieving

Camphor (3.01%). Menthol (2.10%). Pain Relieving

Uses

For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains bruises and sprains

Warnings

For external use only

Do not use

on wounds or damaged skin with a bandage to cover the surface

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days symptoms 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 6 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 6 years of age: Consult a

Inactive Ingredients

ALCOHOL DENAT., BENZYL ALCOHOL. CARBOMER, DISODIUM EDTA, FD&C BLUE NO.1, FD&C YELLOW NO.5, GLYCERIN, GINSENG, PEG-7 GLYCERYL COCOATE, PHENOXYETHANOL, POTASSIUM SORBATE, SODIUM BENZOATE, SODIUM CHLORIDE, TOCOPHEROL, TRIETHA-NOI AMINE WATER

Recommendations:

- Avoid sun exposure on treated areas If pregnant or breast-feeding, ask a health professional before use.
- Do not use if carton is open or if transparent plastic seal on tube cap is missing or broken.

www.phquirogel.com

MANUFACTURED BY / **FABRICADO POR: ISSISLEN LABORATORI** JOSEP ANSELM CLAVÉ 47, 5 ESPLUGUES DE LLOBREGAT BARCELONA 08950 SPAIN

DISTRIBUTED BY / DISTRIBUIDO POR: **INICIATIC DIGITAL SERVICES** 848 N. RAINBOW BLVD. #3102 LAS VEGAS, NV 89107, U.S.

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MADE IN SPAIN HECHO EN ESPAÑA





Datos del Medicamento

Ingredientes Activos Acción .Alivio del Dolor Alcanfor (3,01%). Mentol (2,10%). Alivio del Dolor

Indicaciones

■ Alivio temporal de dolores y molestias leves de músculos y articulaciones asociados con el dolor de espalda, artritis, torceduras, hematomas y esguinces

Advertencias

Para uso externo solamente

No usar

- en heridas o sobre piel dañada con vendajes que cubran la superficie

Durante el uso del producto

evitar el contacto con los ojos

Dejar de usar y consultar con un médico

- si el estado empeora los síntomas persisten más de 7 días los síntomas mejoran y reaparecen en pocos días

Mantener alejado del alcance de los niños. Si es ingerido, buscar atención médica o contactar con un centro de toxicología inmediatamente

Instrucciones

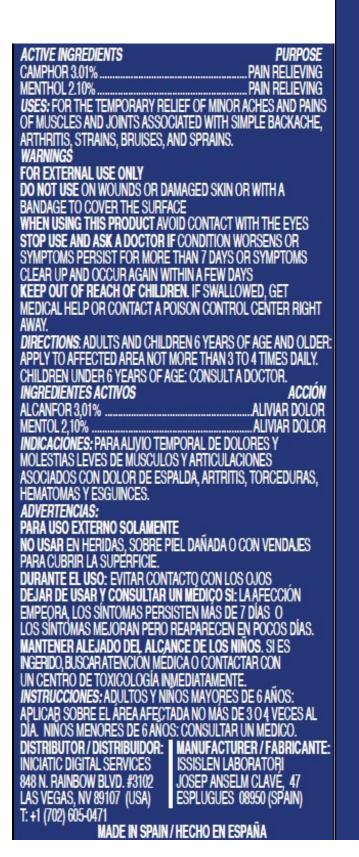
- Adultos y niños de 6 años o más: Aplicar en la zona afectada hasta 3 o 4 veces al día.
- Niños menores de 6 años: Consultar con un médico

Ingredientes Inactivos

AGUA, ALCOHOL BENCÍLICO, ALCOHOL DESNATURALIZADO, BENZOATO DE SODIO, CARBOMER, CLORURO DE SODIO, EDTA DISÓDICO, FD&C AZUL NO.1, FD&C AMARILLO NO.5, FENOXI-ETANOL, GINSENG, GLICERINA, PEG-7 GLICERIL COCOATE, SORBATO DE POTASIO, TOCOFEROL, TRIETANOLAMINA.

Recomendaciones:

- Evitar exponer al sol el area afectada
- Embarazadas o en periodo de lactancia, consultar a su médico antes de usar
- No utilizar si la caja está abierta o si el sello de plástico transparente del tapón está roto.



GEL PARA ALIVIAR EL DOLOR Net Wt. / Peso Neto 2.5 oz (70.1 g) PH-QUIROGE GEL PAIN RELIEVING

PHQUIROGEL

camphor, menthol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70544-001	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	30.1 mg in 1 mL		
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	21 mg in 1 mL		

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		
CARBOXYPOLYMETHYLENE (UNII: 0 A5MM307FC)		
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		
GLYCERIN (UNII: PDC6A3C0OX)		
ASIAN GINSENG (UNII: CUQ3A77YXI)		
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
TOCOPHEROL (UNII: R0ZB2556P8)		
TROLAMINE (UNII: 9O3K93S3TK)		
WATER (UNII: 059QF0KO0R)		

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:70544-001-01	1 in 1 BOX	04/05/2016		
1	75 mL in 1 TUBE; 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	04/04/2016		

Labeler - Issislen Laboratori S.L. (463538806)

Revised: 4/2016 Issislen Laboratori S.L.