

**SHELO NABEL THERAPYGEL- menthol gel**

Corporativo Serysi S de R L de C V

*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.*

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**Sheló NABEL Therapygel Topical Analgesic Gel**

**Active Ingredient U061**

<b>Active Ingredient</b>	<b>Purpose</b>
Menthol 2.6% .....	Topical Analgesic
Camphor 2 % .....	Topical Analgesic

**Inactive Ingredient U061**

<b>Inactive ingredients</b>
Distilled Water, Ethanol, Polysorbate 20, Carbomer 940, Triethanolamine, FD&C Blue No 1.

**Warnings U061**

<b>Warnings</b>
<b>For external use only</b>
<b>Avoid contact with eyes</b>
<b>Stop use and ask a doctor if</b> Condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.
<b>Do not apply to wounds or damaged skin</b>
<b>Do not bandage tightly</b>
<b>If pregnant or breast-feeding, ask a health professional before use</b>
<b>Keep out of reach of children</b>
<b>If swallowed, get medical help or contact a Poison Control Center right away.</b>

Uses U061

**Use:** For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises, and sprains. Cooling pain relief.

KROC U061

**Keep out of reach of children**

**If swallowed, get medical help or contact a Poison Control Center right away.**

Admon U061

**Directions** ■ Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. Gently massage the area until absorbed. ■ Children under 2 years of age: Consult a physician.

Indications U061

**COOLING PAIN RELIEF**

Display Panel





# THERAPYGEL

TOPICAL ANALGESIC GEL



**COOLING PAIN RELIEF**

**MENTHOL 2.6%**

**CAMPBOR 2%**

CAMPHOR 2%

Made in Mexico

NET CONTENTS 8.4 fl oz / 250 ml

## SHELO NABEL THERAPYGEL

menthol gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71424-1061
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	26 mg in 1 g
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	20 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
POLYSORBATE 20 (UNII: 7T1F30V5YH)	

### Packaging

		Marketing Start	Marketing End
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71424-1061-1	250 g in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/18/2017	
<b>Marketing Information</b>				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	OTC monograph not final	part348	05/18/2017	

**Labeler** - Corporativo Serysi S de RL de C V (816628390)

**Registrant** - Corporativo Serysi S de RL de C V (816628390)

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