

SHELO NABEL GEL PARA GOLPES- 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.

Sheló NABEL Gel Para Golpes

Active Ingredient

Active Ingredient	Purpose
Menthol 1.26%	Topical Analgesic

Inactive ingredient

Inactive Ingredients Demineralized Water, Carbopol Ultrez 10, White Willow Extract, Harpagophytum Extract, Mint Essential Oil, Eucalyptus Essential Oil, Aloe Vera Extract.

warnings

Warnings For external use only

Avoid contact with eyes

Stop use and ask a doctor if Condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Do not apply to wounds or damaged skin

Do not bandage tightly

If pregnant or breast-feeding, ask a health professional before use

Uses

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

KROC

Keep out of reach of children

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

ADMINISTRATION

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

FOR THE TEMPORARY RELIEF OF MINOR ACHES
AND PAINS OF MUSCLES AND JOINTS
ASSOCIATED WITH SIMPLE BACKACHE,
ARTHRITIS, STRAINS, BRUISES AND SPRAINS.

Display Panel



GEL PARA GOLPES

**TOPICAL
ANALGESIC GEL
MENTHOL 1.26%**

FOR THE TEMPORARY RELIEF OF MINOR ACHES
AND PAINS OF MUSCLES AND JOINTS
ASSOCIATED WITH SIMPLE BACKACHE,
ARTHRITIS, STRAINS, BRUISES AND SPRAINS.

Made in Mexico

NET CONTENTS 8.7 fl oz / 260 ml

SHELO NABEL GEL PARA GOLPES

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71424-1347
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	12.6 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
MINT (UNII: FV98Z8G1TP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71424-1347-1	260 g in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2017	

Marketing Information

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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