

**GINSENGGO- camphor, menthol, methyl salicylate patch**  
**Sinsin Pharmaceutical Co., Ltd.**

*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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**Drug Facts**

**Active ingredients**

**DL-Camphor 0.70 %**

**L-Menthol 1.40 %**

**Methyl Salicylate 2.80 %**

**Purpose**

**Topical Analgesic**

**Uses**

for temporary relief of minor pain associated with:

- Arthritis
- Simple backache
- Tendonitis
- Muscle strains
- Muscle sprains
- Bruises

**Warnings**

**For external use only.**

- **If pregnant or breast feeding**, consult a healthcare professional.
- **Allergy Alert:** If you are allergic to any ingredients of this product, ask a doctor before use.

**Do not use**

- on wounds or damaged skin
- with, or at the same time as, other external analgesic products.

**When using this product**

- use only as directed
- do not bandage tightly or use with heating pad.
- avoid contact with the eyes and mucous membranes.

**Stop use and ask a doctor if**

- conditions worsens.
- symptoms persist more than 7 days or clear up and occur again within a few days.
- rash, itching, or excessive irritation develops.

**Keep out of reach of children.**

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

### **Directions**

Adults and children 12 years of age and older

- clean and dry affected area
- remove patch from film.
- apply to affected area not more than 3 to 4 times daily for 7 days.

***Children under 12 years of age consult a doctor***

***Reseal pouch containing used portion patched***

### **Other information**

- avoid storing product in direct sunlight
- protect product from excessive moisture

### **Inactive ingredients**

Butylated Hydroxy Toluene, Butyl Paraben, Carboxymethylcellulose sodium, Castor Oil, Concentrated Glycerin, D-Sorbitol Solution, Disodium Edetate, Gelatin, Kaolin, Magnesium Aluminometasilicate, Methyl paraben, Phellodendron Bark Extract, Polyethylene glycol Monostearate, Polyvinyl Alcohol, Purified Water, Sodium Polyacrylate, Tartaric acid

### **How To Apply**

1. Pull off printed paper
2. Peel off protective film
3. Gently apply medicated pad in the affected area
4. Pull off the remaining paper

### **Manufactured by:**

**SINSIN PHARM. CO., LTD.**

<http://www.sinsin.com> tel: 82-31-776-1140~3

776-6, Wonsi-Dong, Danwon-Gu, Ansan City, Kyunggi-Do, Korea

Made in Korea

### **Principal Display Panel**



# GINSENGGO

camphor, menthol, methyl salicylate patch

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55264-005
Route of Administration	TOPICAL, CUTANEOUS, TRANSDERMAL		

Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)		CAMPHOR (SYNTHETIC)	0.7 mg
LEVOMENTHOL (UNII: BZ1R15MTK7) (LEVOMENTHOL - UNII:BZ1R15MTK7)		LEVOMENTHOL	1.4 mg
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)		METHYL SALICYLATE	2.8 mg

Inactive Ingredients	
Ingredient Name	Strength
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
CASTOR OIL (UNII: D5340Y2I9G)	
GLYCERIN (UNII: PDC6A3C0OX)	
SORBITOL (UNII: 506T60A25R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GELATIN (UNII: 2G86QN327L)	
KAOLIN (UNII: 24H4NWX5CO)	
SILODRATE (UNII: 9T3UU8T0QK)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PHELLODENDRON AMURENSE BARK (UNII: PBG27B754G)	
PEG-8 STEARATE (UNII: 2P9L47VI5E)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
WATER (UNII: 059QF0KO0R)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JN12J)	
TARTARIC ACID (UNII: W4888I119H)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55264-005-06	1 in 1 CARTON	11/13/2020	
1	NDC:55264-005-01	6 in 1 POUCH; 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	08/22/2013	

**Labeler** - Sinsin Pharmaceutical Co., Ltd. (823149161)**Registrant** - Sinsin Pharmaceutical Co., Ltd. (823149161)**Establishment**

Name	Address	ID/FEI	Business Operations
Sinsin Pharmaceutical Co., Ltd.		687867143	manufacture(55264-005)

Revised: 11/2020

Sinsin Pharmaceutical Co., Ltd.