

**SOMTHAWIN ANGKI THAI AROMA HERB YELLOW OIL- somthawin angki thai
aroma herb yellow oil solution
AMBIENCE FAMILY INC**

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

FOR EXTERNAL USE ONLY.

If swallowed, get medical attention or contact a Poison Center immediately Directions Apply generously of affected areas and massage gently until cream is absorbed in to the skin.

Stop use and ask doctor if Condition worsens.

Severe skin irritation occurs.

Symptoms persist for more than 7 days.

If pregnant or breast-feeding, or if you have sensitive skin, ask a healthcare professional before use.

Champhor 25%

Menthol 11%

External Analgesic

Temporarily relieve body aches and pains of muscles and joints associated with psimple backache parthritis astrains abruises asprains

For adults and children over 12 apply on the affected area not more than 3-4 times daily.

For children under 12 years of age,

consult a doctor before using.

Stop use and ask doctor if Condition worsens.

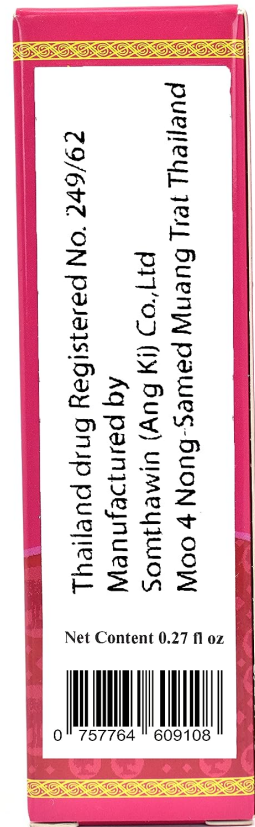
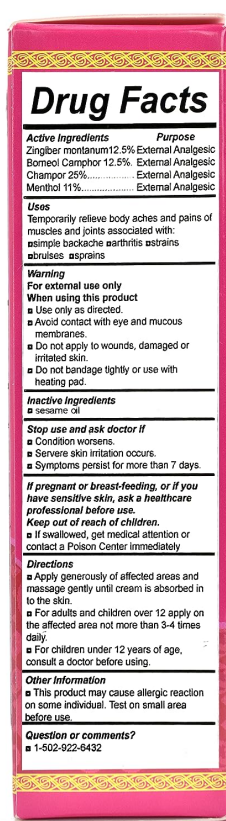
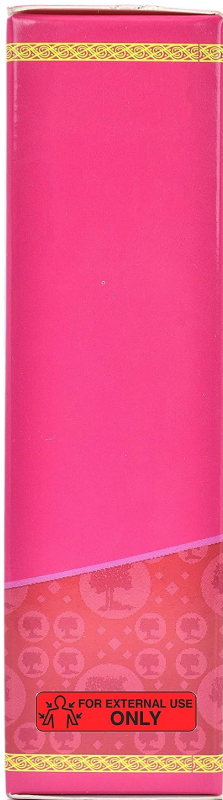
Severe skin irritation occurs.

Symptoms persist for more than 7 days.

External Analgesic

Zingiber montanum12.5% External Analgesic

Borneol Camphor 12.5%. External Analgesic





Yellow Oil
Somthawin (Ang Ki)

SOMTHAWIN ANGKI THAI AROMA HERB YELLOW OIL

somthawin angki thai aroma herb yellow oil solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84212-334
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BORNEOL (UNII: M89NIB437X) (BORNEOL - UNII:M89NIB437X)	BORNEOL	1 mg in 8 mL
ZINGIBER MONTANUM ROOT (UNII: 3NG517NPD8) (ZINGIBER MONTANUM ROOT - UNII:3NG517NPD8)	ZINGIBER MONTANUM ROOT	1 mg in 8 mL

Inactive Ingredients

Ingredient Name	Strength
MENTHOL (UNII: L7T10EIP3A)	
CAMPHOR OIL (UNII: 75IZZ8Y727)	
SESAME OIL (UNII: QX10HYY4QV)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84212-334-01	8 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/01/2024	

Labeler - AMBIENCE FAMILY INC (084561491)**Registrant** - Venture Go LLC (118882840)

Revised: 6/2024

AMBIENCE FAMILY INC