

FOOT- menthol cream
Smith Amish, LLC

Smith Amish Foot Cream (2oz) - 72609-703-02

Drug Facts

Active Ingredient

Menthol 1.25%

Purpose

Topical analgesic

Uses

For temporary relief of minor aches and pains

Warnings

- For external use only
- Avoid contact with eyes
- If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days,, discontinue use of this product and consult a doctor.

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.

Directions

Adults and children 2 years of age and over: apply to affected area not more than 3 to 4 times daily. Children under 2 years of age consult a doctor.

Other information

Protect from excessive heat

Store at 20-25 °C (68-77F)

Inactive Ingredients

Arnica montana flower extract, cetyl palmitate, ethylhexylglycerin, glycerin, glyceryl

stearate, jojoba (*Buxus chinensis*) oil, peppermint (*mentha piperita*) oil, phenoxyethanol, sodium citrate, stearic acid, sweet almond (*Prunus amygdalus dulcis*) oil, tea tree (*Melaleuca alternifolia*) oil, tocopherol (Vitamin E), water

Questions?

Call (866) 419-3567

Monday through Friday 8:00am - 2:00pm ET

Tamper evident: Do not use if safety seal under cap is broken or missing.

Manufactured for Smith Amish LLC 16350 N Hwy 329 Reddick, FL 32634

Label

Smith Amish Arthritis Cream (72609-702-02)

Provides Penetrating Relief

- Arthritic pain
- Muscle Pain
- Joint Pain
- Back Pain

Fresh scent from Arnica & Eucalyptus

All natural and non-greasy

Net wt. 2oz (59g)

THE BRAND YOU CAN TRUST

Topical Analgesic



FOOT CREAM

PROVIDES PENETRATING PAIN RELIEF

for intense foot and leg discomfort

Fresh Scent from
Tea Tree, Arnica & Peppermint
All-Natural and Non-Greasy

Net wt. 2 oz | 59 g

Drug Facts

Active Ingredient	Purpose
Menthol 1.25 %	Topical Analgesic

Uses: For the Temporary relief of minor aches and pains.

Warnings:

- For external use only.
- Avoid contact with eyes • If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor. • Do not apply to wounds or damaged skin • Do not bandage tightly.

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.

Directions: Adults and children 2 years of age and over: apply to affected area not more than 3 or 4 times daily. Children under 2 year of age: consult a doctor.

Other Information:

- Protect from excessive heat
- Store at 20 - 25°C (68-77° F)

Inactive Ingredients: Arnica montana flower extract, cetyl palmitate, ethylhexylglycerin, glycerin, glyceryl stearate, jojoba (*Buxus chinensis*) oil, peppermint (*Mentha piperita*) oil, phenoxyethanol, sodium citrate, stearic acid, sweet almond (*Prunus amygdalus dulcis*) oil, tea tree (*Melaleuca alternifolia*) oil, tocopherol (Vitamin E), water.

Questions? Call (866) 419-3567 Monday through Friday 8:00 AM - 2:00 PM ET

Tamper evident: Do not use if safety seal under cap is broken or missing.

Manufactured for Smith Amish, LLC
16530 N Hwy 329 Reddick, FL 32634



FOOT

menthol cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72609-703
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.25 g in 100 g
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Inactive Ingredients

Ingredient Name	Strength
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ALMOND OIL (UNII: 66YXD4DKO9)	
JOJOBA OIL (UNII: 724GKU717M)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
ARNICA MONTANA WHOLE (UNII: O80TY208ZW)	
CETYL PALMITATE (UNII: 5ZA2S6B08X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TEA TREE OIL (UNII: VIF565UC2G)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72609-703-02	1 in 1 BOX	08/28/2024	
1		59 g 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 Drug	M017	08/28/2024	

Labeler - Smith Amish, LLC (081504527)

Revised: 11/2024

Smith Amish, LLC