ZPOL ULTRA- menthol, methyl salicylate cream
Laboratorios Zepol S.A.

Zpol Ultra

Drug Facts

Active ingredients

Menthol 4%Methyl Salicylate 10%

Purpose

Active Ingredients	Purpose
Menthol 2.84%	Topical Analgesic
Methyl salicylate 18.24%	Topical Analgesic

Uses

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

Warnings

For external use only.

When using this product

Do not apply to wounds or damaged skin

Do not bandage tightly

Avoid contact with eyes

Do not use with a heating pad

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.

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 older: Apply to affected area not more than 3 to 4 times daily.

Children under 2 years of age: Do not use, consult a doctor.

Other information

Store between 20-25°C (68-77°F).

Inactive ingredients

Allantoin, benzoic acid, cetearyl alcohol, cetearyl glucoside, pheonoxyethanol, stearic acid, water.

Questions?

Call toll free +506 (800)-937-6572 laboratorioszepol@zepolab.com

zpol ULTRA 1.02 Oz

NDC 55715-008-01

zpol ULTRA

MUSCULAR Topical analgesic cream

Net wt. 1.02 Oz

ZepoLAB

Manufactured by

Laboratorios Zepol, S.A.

Curridabat, San José,

Costa Rica, 11801

Product of Costa Rica

IN COSTA RICA CALL FRO FREE 800-937-6572

(800-SZEPOLSA)

Regional Customer Service:

laboratorioszepol@zepolab.com

www.zepolab.com

Tamper Evident: Do not use if seal undercap is broken or missing

EXTERNAL USE/OVER THE COUNTER Zpol ULTRA Cream

is a medicated cream, due to its components and texture it is recommended for the

relief of muscle pain, bumps, sprains and sports injuries. It also relieves arthritic and rheumatic pain.

zpol ULTRA 2.08 Oz

NDC 55715-008-02

zpol ULTRA

Topical analgesic cream

2.08 Oz

ZepoLAB

Zpol ULTRA 4.02 Oz

NDC 55715-008-03

zpol ULTRA

Topical analgesic cream

4.02 Oz

ZepoLAB

ZPOL ULTRA

menthol, methyl salicylate cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55715-008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.84 g in 100 g
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	18.24 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SORBITAN TRISTEARATE (UNII: 6LUM696811)	
CARBOMER (UNII: 0A5MM307FC)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
PEG-40 STEARATE (UNII: ECU18C66Q7)	
CETEARETH-20 (UNII: YRC528SWJY)	
EUCALYPTOL (UNII: RV6J6604TK)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
TRIETHANOLAMINE (UNII: 9O3K93S3TK)	
THYMOL (UNII: 3J50XA376E)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55715-008-01	29 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2025	
2	NDC:55715-008-02	59 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2025	
3	NDC:55715-008-03	119 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/01/2025	

Labeler - Laboratorios Zepol S.A. (853070985)

Establishment

Name	Address	ID/FEI	Business Operations
Laboratorios Zepol S.A.		853070985	manufacture(55715-008)

Revised: 3/2025 Laboratorios Zepol S.A.