LITHIUM- lithium bromatum liquid Marco Pharma International LLC.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Drug Facts

Active Ingredients

Lithium Bromatum 3xHPUS

The letters HPUS indicates that the components in this product are officially monographed in the Homeopathic Pharmacopoeia of United Sates.

Purpose

For the temporary relief of symptoms due to depression, anxiety, and vertigo.

Keep out of reach of children.

Dosage

Adults: Take 20 drops on the toungue or in water two to three times daily. Children receive 1/2 of the adult amount.

Warnings

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

(Read Suggested Use Section)

Inactive Ingredients

Ethyl alcohol (18% by vol.) and water.



Dosage: Adults take 20 drops on the tongue or in water two to three times daily. For children, please consult a healthcare professional.

Warning: If pregnant of breast-feeding, consult a health professional before use. Keep out of the reach of children.

Tamper Evident: Use this product only if tamper evident ring has not broken away from base of cap.

INGREDIENTS: 100 mL contains: Lithium Bromatum 3x HPUS. Contains Ethyl Alcohol (18% by vol.) and Water.

HPUS indicates that the active ingredients are in the official Homeopathic Pharmacopoeia of the United States.

Combination
Medicine
Indications: For temporary relief

of symptoms due to depression, anoiety, brain fog, forgetfulness and vertigo.

3.38 FI Oz (100 mL)

Distributed by: Marco Pharma 851 NW Highland St., Roseburg, 0R 97470 Made in Germany

LITHIUM

lithium bromatum liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:60986-1019 Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength LITHIUM BROMIDE (UNII: 864G646184) (LITHIUM CATION - UNII:8H8Z5UER66) LITHIUM BROMIDE 3 [hp_X] in 1 mL

Inactive Ingredients					
Ingredient Name	Strength				
WATER (UNII: 059QF0KO0R)					
ALCOHOL (UNII: 3K9958V90M)					

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:60986- 1019-3	50 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	02/28/2018		
2	NDC:60986- 1019-4	100 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	02/28/2018		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		03/01/1992		

Labeler - Marco Pharma International LLC. (161994277)

Registrant - Marco Pharma International LLC. (161994277)

Establishment							
Name	Address	ID/FEI	Business Operations				
Nestmann Pharma GmbH		323426262	manufacture(60986-1019)				

Revised: 12/2021 Marco Pharma International LLC.