THREEFOLD BALANCE- threefold balance tablet
Uriel Pharmacy Inc.

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.

Threefold Balance

Directions: FOR ORAL USE ONLY

Ages 12 and older: 1-2 tablets 3-4 times daily. Ages 2-11: 1 tablet. Under age 2: Consult a doctor.

Active Ingredients: Olibanum (Rubber resin from wood of incense tree) 2X, Myrrha (Myrrh resin) 2X, Aurum metallicum (Metallic gold) 6X

Inactive Ingredients: Microcrystalline cellulose, Stearic acid, Magnesium stearate

Uses: For overall well-being. Temporary relief of environmental sensitivity.

KEEP OUT OF REACH OF CHILDREN.

Allergen Statement: This product was produced on equipment that also processes products that contain lactose.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858

Made with care by Uriel, East Troy, WI 53120

www.urielpharmacy.com

THREEFOLD BALANCE

threefold balance tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-1198
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
FRANKINCENSE (UNII: R9XLF1R1WM) (FRANKINCENSE - UNII:R9XLF1R1WM)	FRANKINCENSE	2 [hp_X]

MYRRH (UNII: JC71GJ1F3L) (MYRRH - UNII:JC71GJ1F3L)	MYRRH	2 [hp_X]
GOLD (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO)	GOLD	6 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	brown	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-1198-1	200 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	09/01/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-1198)

Revised: 5/2018 Uriel Pharmacy Inc.