OSTEODORON AM- osteodoron am powder
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.

Osteodoron AM

Directions: FOR ORAL USE ONLY.

Take in the morning. Use in combination with Osteodoron PM Powder. Ages 12 and older: 1/8 teaspoon. Ages 2-11: 1/16 teaspoon. Under age 2: Consult a doctor.

Active Ingredients: Cucurbita (Squash) 3X, Apatite (Nat. calcium fluorophosphate) 6X, Fluorite (Calcium fluoride) 6X, Quartz (Rock crystal) 6X

Inactive Ingredient: Lactose

prepared using rhythmical processes

Uses: For healthy bone and teeth development.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Contains lactose. 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
Uriel, East Troy, WI 53120
shopuriel.com Lot:

OSTEODORON AM
osteodoron am powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-7190
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CUCURBITA PEPO FLOWER (UNII: 413MGP37HQ) (CUCURBITA PEPO FLOWER - UNII:413MGP37HQ)	CUCURBITA PEPO FLOWER	3 [hp_X] in 1 g
CALCIUM FLUORIDE (UNII: O3B55K4YKI) (FLUORIDE ION - UNII:Q80VPU408O)	CALCIUM FLUORIDE	6 [hp_X] in 1 g
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	6 [hp_X] in 1 g
FLUORAPATITE (UNII: M4CM1H238J) (FLUORAPATITE - UNII:M4CM1H238J)	FLUORAPATITE	6 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-7190-4	50 g 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(48951-7190)

Revised: 4/2024 Uriel Pharmacy Inc.