OSTEODORON PM- osteodoron pm 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 PM

Directions: FOR ORAL USE ONLY.

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

Active Ingredients: 100 gm contains: 5gm Conchae (Oyster shells) 1X; Quercus (Oak) 4X, Fluorite (Calcium fluoride) 6X, Quartz (Rock crystal) 6X

Inactive Ingredient: Lactose

"prepared using rhythmical processes"

Use: 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 PM
osteodoron pm powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-7191
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	6 [hp_X] in 1 g
OSTREA EDULIS SHELL (UNII: 49OY13BE7Z) (OSTREA EDULIS SHELL - UNII:49OY13BE7Z)	OSTREA EDULIS SHELL	1 [hp_X] in 1 g
QUERCUS ROBUR WHOLE (UNII: R7QMG0BT2W) (QUERCUS ROBUR WHOLE - UNII:R7QMG0BT2W)	QUERCUS ROBUR WHOLE	4 [hp_X] in 1 g
CALCIUM FLUORIDE (UNII: O3B55K4YKI) (FLUORIDE ION - UNII:Q80VPU408O)	CALCIUM FLUORIDE	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:48951-7191-4	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(48951-7191)

Revised: 3/2024 Uriel Pharmacy Inc.