HANTAI 8030 CA-D3 PREMIUM- ostreae concha, calcium tablet APEXEL CO., LTD.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

ACTIVE INGREDIENT

Ostreae Concha (Calcium 99%)

INACTIVE INGREDIENT

Magnesium, Zinc Oxide, Selenium, Germanium, Iron, Potassium, Manganese

PURPOSE

To prevent and cure the osteoporosis, cardiovascular disorders, depression, nerve stability, muscle pain, arthritis, menstrual pain, ostalgia and promote growth and healing.

KEEP OUT OF REACH OF CHILDREN

Keep out of Reach of Children

WARNINGS

- Please note it may choke during the intake tablets.
- Please check product ingredients if you have any allergies before taking.
- Please be careful during open the product package.
- Keep product out of direct sunlight, high temperature and humidity.
- Store in a cool dry place.
- Keep out of reach of children.
- Any items past the expiration date or damaged in transit can be exchanged where you originally purchased the item.

DOSAGE & ADMINISTRATION

For oral use only

USES

Take one tablet every time, two times a day with enough water.

PACKAGE LABEL



ostreae concha, calcium tablet

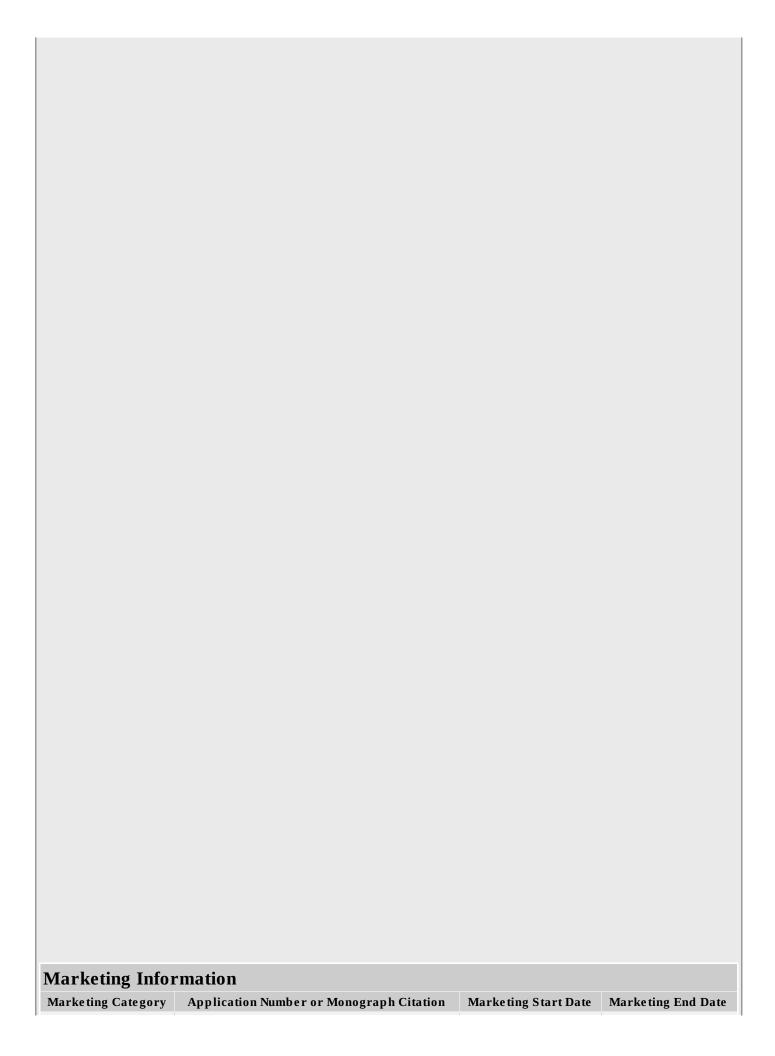
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55259-4001
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
CALCIUM (UNII: SY7Q814VUP) (CALCIUM - UNII:SY7Q814VUP)	CALCIUM	0.99		

Inactive Ingredients			
Ingredient Name	Strength		
ZINC OXIDE (UNII: SOI2LOH54Z)			
MAGNESIUM (UNII: 138 ZP9 9 9 2 A)			
SELENIUM (UNII: H6241UJ22B)			
GERMANIUM (UNII: 00072J7XWS)			
IRON (UNII: E1UOL152H7)			
POTASSIUM (UNII: RWP5GA015D)			
MANGANESE (UNII: 42Z2K6ZL8P)			

Product Characteristics			
Color	white	Score	score with uneven pieces
Shape	OVAL	Size	12mm
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code Package Description Marketing Start Date Marketing End Date			Marketing End Date
1	NDC:55259-4001-2	1 in 1 PACKAGE		
1		120 in 1 BOTTLE; Type 0: Not a Combination Product		



unapproved drug other	05/18/2015	

Labeler - APEXEL CO., LTD. (687287979)

Registrant - APEXEL CO., LTD. (687287979)

Establishment				
Name	Address	ID/FEI	Business Operations	
APEXEL CO., LTD.		687287979	manufacture (55259-4001)	

Revised: 6/2015 APEXEL CO., LTD.