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

HANTAI 8030 CA-D3 PREMIUM

ostreae concha, calcium tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55259-1113
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM (UNII: SY7Q814VUP) (CALCIUM - UNII:SY7Q814VUP)	CALCIUM	0.99 mg

Inactive Ingredients

Ingredient Name	Strength
ZINC OXIDE (UNII: SOI2LOH54Z)	
MAGNESIUM (UNII: I38ZP9992A)	
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
1	NDC:55259-1113-2	1 in 1 PACKAGE	03/18/2021	
1	NDC:55259-1113-1	120 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
unapproved drug other		03/18/2021	

Labeler - APEXEL CO., LTD. (687287979)

Registrant - APEXEL CO., LTD. (687287979)

Establishment

Name	Address	ID/FEI	Business Operations
APEXEL CO., LTD.		687287979	manufacture(55259-1113)

Revised: 2/2024 APEXEL CO., LTD.