GREAT CALCIUM- ostreae concha 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

calcium (ostreae concha)

INACTIVE INGREDIENT

magnesium, vitamin D, red ginseng powder

PURPOSE

Treatment and prevention of osteoporosis

Growth and development

Treatment and prediction of cardiovascular disease due to calcification of blood
Attention deficit in children, depression, nervous symptoms, nervous stability to anxiety
Myalgia, Bone Pain, Arthritis, Dysmenorrhea Treatment

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children.

WARNING

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.

Any items past the expiration date or damaged in transit can be exchanged where you originally purchased the item.

consult your doctor if any abnormal symptoms occur

USES

for oral use only

INDICATION & USAGE SECTION

take two capsules once, two times a day

GREAT CALCIUM

Drug Facts

Active ingredients

calcium (ostreae concha)

Uses

Treatment and prevention of osteoporosis

Growth and development

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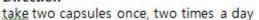
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Direction



Other Information

keep in a room temperature keep out of the direct sunlight

Inactive Ingredient

magnesium, vitamin D, red ginseng powder

GREAT CALCIUM

ostreae concha tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55259-6001	
Route of Administration	ORAL			

of Strength Streng	th
0.76	
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Inactive Ingredients		
Ingredient Name	Strength	
MAGNESIUM (UNII: 138ZP9992A)		
PANAX GINSENG WHOLE (UNII: 9L5JEP7MES)		
VITAMIN D (UNII: 9VU1KI44GP)		

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

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:55259- 6001-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	11/18/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/18/2018	

Labeler - APEXEL CO., LTD. (687287979)

Registrant - APEXEL CO., LTD. (687287979)

Establishment				
Name	Address	ID/FEI	Business Operations	
APEXEL CO., LTD.		687287979	manufacture(55259-6001)	

Revised: 3/2021 APEXEL CO., LTD.