AG-X- 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

calcium

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

AG-X

Drug Facts

Active ingredients
calcium (ostreae concha)

Uses
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Growth and development
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Direction
take two capsules once, two times a day

Other Information
keep in a room temperature
keep out of the direct sunlight

Inactive Ingredient
magnesium, vitamin D, red ginseng powder

AG-X
calcium tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55259-8001
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CALCIUM (UNII: SY7Q814VUP) (CALCIUM - UNII:SY7Q814VUP)	CALCIUM	0.76

Inactive Ingredients	
Ingredient Name	**Strength**
MAGNESIUM (UNII: I38ZP9992A)	
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			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55259-8001-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	08/02/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/02/2019	

Labeler - APEXEL CO., LTD. (687287979)

Registrant - APEXEL CO., LTD. (687287979)

Establishment

Name	Address	ID/FEI	Business Operations
APEXEL CO., LTD.		687287979	manufacture(55259-8001)

Revised: 8/2019 APEXEL CO., LTD.