

TALL G U- calcium tablet
coexleaders 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

Oystershell powder

INACTIVE INGREDIENT

Sweet potato powder, chitosan powder, black soybean powder

PURPOSE

Supplements to treat height growth

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

Tall G U



Drug Facts

Active Ingredients

Oystershell powder

Uses

- Supplements to treat height growth

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- consult your doctor if any abnormal symptoms occur

Directions

- take two capsules once, two times a day

Other Information

- read the directions and warnings before use
- avoid freezing and excessive heat above 40 degree C (104 degree F)

Inactive Ingredients

Seaweed powder, anchovy powder, stevia

TALL G U

calcium tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81445-0008	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
CALCIUM (UNII: SY7Q814VUP) (CALCIUM - UNII:SY7Q814VUP)		CALCIUM	0.85	
Inactive Ingredients				
Ingredient Name			Strength	
STEVIA LEAF (UNII: 6TC6NN0876)				
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:81445-0008-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/25/2021	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			01/25/2021	

Labeler - coexleaders co.,ltd. (695097730)

Registrant - coexleaders co.,ltd. (695097730)

Establishment			
Name	Address	ID/FEI	Business Operations
coexleaders co.,ltd.		695097730	manufacture(81445-0008)