

RESTO L- 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

Protein powder, chaga mushroom powder, ginger powder

PURPOSE

Lung disease treatment supplements

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

Resto L

Drug Facts

Active Ingredients
Oystershell powder

Uses
■ Lung disease treatment supplements

Warning

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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

Protein powder, chaga mushroom powder, ginger powder

RESTO L

calcium tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81445-0006	
Route of Administration	ORAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CALCIUM (UNII: SY7Q814VUP) (CALCIUM - UNII:SY7Q814VUP)	CALCIUM	0.71

Inactive Ingredients	
Ingredient Name	Strength
GINGER (UNII: C5529G5JPQ)	

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:81445-0006-1	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(81445-0006)