POWER GEMSTONE- gins eng pill Myeong SA

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.

ginseng

Magnolia officinalis, cinnamon, Gleditsia japonica, barberry root, Asteris Radix, ballon flower, Zingiber officinale, japanese aconite root

prevention and treatment of Atopy, allergy, rhinitis

keep out or reach of the children

6 times a day (Every 4 hours)

Dose: 4g

Take the appropriate amount.

If any of symptoms occur, consult with your doctor or pharmacist.

6 times a day

POWER GEMSTONE

Drug Facts

Active Ingredient

Ginseng

Use

prevention and treatment of Atopy, allergy, rhinitis

Warning

- Take the appropriate amount.
- If any of symptoms occur, consult with your doctor or pharmacist.
- Precaution for Storage and Handling
 - 1) Close the lid after use
 - 2) Keep out of reach of infants and children
- Do not to store in a place with high/low temperature and exposed to direct sunlight

Direction

6 times a day (Every 4 hours)

Inactive Ingredient

Magnolia officinalis, cinnamon, Gleditsia japonica, barberry root, Asteris Radix, ballon flower, Zingiber officinale, japanese aconite root

Other Information

store between 20-25 °C (68-77 °F) avoid freezing and excessive heat above 40 °C (104 °F)

POWER GEMSTONE

ginseng pill

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71960-0003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
PANAX GINSENG WHOLE (UNII: 9L5JEP7MES) (PANAX GINSENG WHOLE - UNII: 9L5JEP7MES)	PANAX GINSENG WHO LE	20 g in 100 g		

Inactive Ingredients			
Ingredient Name	Strength		
MAGNOLIA OFFICINALIS WHOLE (UNII: 9OB542Y2EX)			

Product Characteristics			
Color	green	Score	no score
Shape	OVAL	Size	5mm
Flavor	PEPPERMINT	Imprint Code	None
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:71960- 0003-1	300 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/13/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/13/2018	

Labeler - Myeong SA (689605005)

Registrant - Myeong SA (689605005)

Establishment			
Name	Address	ID/FEI	Business Operations
Myeong SA		689605005	manufacture(71960-0003)

Revised: 1/2020 Myeong SA