Revised: Jan. 2024（st Version）

Storage：Store at room temperature
Shelf Life：3 years

Standard Commodity Classification No. of Japan	
875200	
Approval No.	16100AMZ04764000
Date of Initial Marketing in Japan	Oct. 1987

Kampo product

JPS Hangekobokuto Extract Granules for Ethical Use

3. COMPOSITION AND PRODUCT DESCRIPTION

3.1 Composition

Product Name	JPS Hangekobokuto Extract Granules for Ethical Use
Active Ingredients	JP Pinellia Tuber 6 g JP Poria Sclerotium 5 g JP Magnolia Bark 3 g JP Perilla Herb 3 g JP Ginger 1 g 7.5 g/day of this product contains 2.2 g of a dried JP Hangekobokuto extract of the above mixed crude drugs.
Inactive Ingredients	Mg Stearate, Sucrose Esters of Fatty Acids, Lactose Hydrate

3.2 Product Description

Dosage Form	Granules
Color	Light brown
Smell	Characteristic smell
Taste	Slightly bitter
ID Code	J-16

4. INDICATIONS

The following symptoms in patients with depressed feeling, feeling of a foreign body in the throat or esophagus, sometimes accompanied by palpitations, dizziness, nausea, etc.:
Anxiety neurosis, neurotic gastritis, mild hyperemesis gravidarum, coughing, hoarseness

6. DOSAGE AND ADMINISTRATION

The usual adult dose is 7.5 g/day orally in 2 or 3 divided doses before or between meals. The dosage may be adjusted according to the patient's age and body weight, and symptoms.

8. IMPORTANT PRECAUTIONS

8.1 When this product is used, the patient's "SHO" (constitution/symptoms) should be taken into consideration. The patient's progress should be carefully monitored, and if no improvement in symptoms or findings is observed, continuous administration should be avoided.

8.2 When this product is used in combination with other Kampo products, etc., attention should be paid to the duplication of the contained crude drugs.

9. PRECAUTIONS CONCERNING PATIENTS WITH SPECIFIC BACK GROUNDS

9.5 Pregnant Women
This product should be used in pregnant women or women who may possibly be pregnant only if the expected therapeutic benefits outweigh the possible risks associated with treatment.

9.6 Brest-Feeding Women
Considering the therapeutic benefits and the benefits of breastfeeding, continuation or discontinuation of breastfeeding should be considered.

9.7 Pediatric Use
No clinical studies have been conducted in children.

9.8 Geriatric Use
Since the physiological functions are generally decreased in elderly patients, careful supervision is recommended; measures such as reducing the dose may be considered.

11. ADVERSE REACTIONS

The following adverse reactions may occur. Patients should be carefully monitored, and if any abnormalities are observed, appropriate measures such as discontinuation of administration should be taken.

11.2 Other Adverse Reactions

	Frequency unknown
Hypersensitivity	Rash, Redness, Pruritus, etc.
Liver	Hepatic function abnormal (increased AST, ALT, etc.)

20. PRECAUTIONS FOR HANDLING

20.1 To maintain the quality of the product, avoid moisture as much as possible and store in a cool place, away from direct sunlight.

20.2 Avoid moisture, especially after opening, and handle with care.

20.3 Since this product is made from crude drugs, the color of the product may vary.

22. PACKAGING

750g（2.5g×300 packets）
105g（2.5g×42 packets）

24. REFERENCE REQUEST AND CONTACT INFORMATION

JPS Pharmaceutical CO., LTD
4-42-22 Higashiyamata, Tsuzukiku, Yokohama City, Kanagawa Prefecture 224-0023, Japan
TEL：045-593-2060

26. MARKETING AUTHORIZATION HOLDER, etc.

26.1 Manufactured and Distributed by:
JPS Pharmaceutical CO., LTD
196-1 Hagadai, Hagamachi, Hagagun, Tochigi Prefecture 321-3325, Japan