

Revised: May 2019 (8th version)

Standard Commodity Classification No. of Japan
875200

- Kampo-preparation -

TEIKOKU Hangekobokuto Extract Granules

< Hangekobokuto >

Storage
Store at room temperature in a tight container. [See the "Precaution for handling" section.]

Approval No.	(61AM) 3690
Date of listing in the NHI reimbursement price	October 1987
Date of initial marketing in Japan	October 1987

Expiration date
Hangekobokuto should be used before the expiration date indicated on the label and the package.

DESCRIPTION

Composition	7.5 g of TEIKOKU Hangekobokuto Extract Granules contains 1.61 g of a dried water extract of the following mixed crude drugs.	
		JP Pinellia Tuber ----- 6.0 g JP Poria Sclerotium ----- 5.0 g JP Magnolia Bark ----- 3.0 g JP Perilla Herb ----- 2.0 g JP Ginger ----- 1.0 g (JP: The Japanese Pharmacopoeia)
Description	Inactive ingredients	JP Lactose Hydrate JP Microcrystalline Cellulose JP Magnesium Stearate
	Dosage form	Granules
	Color	Light brown
	Smell	Characteristic smell
	Taste	Bitter and slightly pungent
	ID code	TEIKOKU 16

INDICATIONS

The following symptoms of those patients who have depressed feelings and a feeling of foreign body in the throat and oesophagus and who sometimes have palpitation, dizziness, nausea, etc.:

Anxiety neurosis, nervous gastritis, hyperemesis gravidarum, coughing and hoarseness

DOSAGE AND ADMINISTRATION

For oral use, the usual adult dosage is 2.5 g of TEIKOKU Hangekobokuto Extract Granules three times daily before meals.

The dosage may be adjusted according to the patient's age, body weight, and symptoms.

PRECAUTIONS

1. Important Precautions

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

(2) When this product is coadministered with other Kampo-preparations (Japanese traditional herbal medicines), etc., attention should be paid to the duplication of the contained crude drugs.

SHO : The term "SHO" refers to a particular pathological status of a patient evaluated by the Kampo diagnosis, and is patterned according to the patient's constitution, symptoms, etc. Kampo-preparations (Japanese traditional herbal medicines) should be used after confirmation that it is suitable for the identified "SHO" of the patient.

2. Adverse Reactions

This product has not been investigated (drug use investigations, etc.) to determine the incidence of adverse reactions. Therefore, the incidence of adverse reactions is not known.

	Incidence unknown
Hypersensitivity ^{Note)}	Rash, Redness, Pruritus, etc.
Hepatic	Abnormality of hepatic function (Increased AST, ALT, etc. levels)

Note) If such symptoms are observed, administration should be discontinued.

3. Use in the Elderly

Because elderly patients often have reduced physiological function, careful supervision and measures such as reducing the dose are recommended.

4. Use during Pregnancy, Delivery or Lactation

The safety of this product in pregnant women has not been established. Therefore, the product should be used in pregnant women, women who may possibly be pregnant only if the expected therapeutic benefits outweigh the possible risks associated with treatment.

5. Pediatric Use

The safety of this product in children has not been established. [Insufficient clinical data.]

PRECAUTIONS FOR HANDLING

- 1.This product should not be stored in direct rays and should be stored in a cool place with, if possible, little humidity.
- 2.Since it is hygroscopic property, this product should not be stored in humid places after opening.

PACKAGING

- Bottles of 500 g
- Boxes of 2.5 g× 42 packets
- Boxes of 2.5 g×252 packets

REQUEST FOR LITERATURE SHOULD BE MADE TO:

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Pharmaceutical Sales & Marketing Dept.
Teikoku Seiyaku Co., Ltd.
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