

Revised: August 2017 (6th version)

Standard Commodity Classification No. of Japan
875200

- Kampo-preparation -

TEIKOKU Shohangekabukuryoto Extract Granules

< Shohangekabukuryoto >

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

Approval No.	(62AM) 0067
Date of listing in the NHI reimbursement price	October 1987
Date of initial marketing in Japan	October 1987

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

DESCRIPTION

Composition	7.5 g of TEIKOKU Shohangekabukuryoto Extract Granules contains 1.73 g of a dried water extract of the following mixed crude drugs.	
	JP Pinellia Tuber -----	8.0 g
	JP Ginger -----	1.5 g
	JP Poria Sclerotium -----	5.0 g
	(JP: The Japanese Pharmacopoeia)	
Inactive ingredients	JP Lactose Hydrate	
	JP Microcrystalline Cellulose JP Magnesium Stearate	
Description	Dosage form	Granules
	Color	Light brown
	Smell	Slightly characteristic smell
	Taste	Slightly pungent
	ID code	TEIKOKU 21

(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. Use in the Elderly

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

3. 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.

4. Pediatric Use

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

INDICATIONS

Hyperemesis gravidarum, vomiting, and nausea

DOSAGE AND ADMINISTRATION

For oral use, the usual adult dosage is 2.5 g of TEIKOKU Shohangekabukuryoto 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.

PRECAUTIONS FOR HANDLING

- This product should not be stored in direct rays and should be stored in a cool place with, if possible, little humidity.
- 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:

Product Information Unit,
Pharmaceutical Sales & Marketing Dept.
Teikoku Seiyaku Co., Ltd.
6-6 Nihonbashi-kobunacho, Chuo-Ku, Tokyo
103-0024, Japan
Tel 0120-189-567

Manufactured by:

Teikoku Kampo Seiyaku Co., Ltd.
80-11 Kitahara, Donari, Donari-cho, Awa, Tokushima
771-1506, Japan

Distributed by:

Teikoku Seiyaku Co., Ltd.
567, Sanbonmatsu, Higashikagawa, Kagawa
769-2695, Japan