ID Code	SG-18R

June 2009
(2nd version)

Standard Commodity Classification No. of Japan
875200

- Kampo product -
Powerful Drug (only Large Package)

OHSUGI Keishikaryojutsubuto Extract Granules G

(Keishikaryojutsubuto)

Storage: Store at room temperature.
See the section "PRECAUTIONS FOR HANDLING"
Expiration date: The expiration date is specified on the container or the outer package

Approval No.	(61AM) 3894
Date of listing in the NHI reimbursement price	October 1986
Date of initial marketing in Japan	October 1986

DESCRIPTION

(1) The daily dose of this product, 9.0g, contains 4.6g of the dried extract (Keishikaryojutsubuto extract) from the following mixed crude drugs.
JP Cinnamon Bark ---------- 4 g
JP Glycyrrhiza ---------- 2 g
JP Ginger ---------- 1 g
JP Poria Sclerotium ---------- 4 g
JP Jujube ---------- 4 g
JP Atractylodes Lancea Rhizome ---------- 4 g
JP Peony Root ---------- 4 g
JP Powdered Processed Aconite Root ---------- 1 g
(JP: The Japanese Pharmacopeia)
The inactive ingredients contained are Lactose Hydrate, Corn Starch and Magnesium Stearate.

(2) This product is light grayish dark brown-colored granules, smells uniquely, and tastes sweet lingeringly.
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INDICATIONS

Arthralgia and neuralgia

DOSAGE AND ADMINISTRATION

The usual adult dose is 9.0 g/day orally in 2 or 3 times before or between meals. The dosage may be adjusted according to the patient's age, body weight, and symptoms.

PRECAUTIONS

(1) Careful Administration (Keishikaryojutsubuto should be administered with care in the following patients.)

1) Patients with strong constitution [Adverse reactions are likely to occur, and the symptoms may be aggravated.]
2) Patients with sensitivity to heat, a tendency towards hot flushes and red face.

(2) 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) Since this product contains Glycyrrhiza, careful attention should be paid to the serum potassium level, blood pressure, etc., and if any abnormality is observed, administration should be discontinued.
3) When this product is coadministered with other Kampo-products (Japanese traditional herbal medicines), etc., attention should be paid to the duplication of the contained crude drugs. Special caution should be exercised when this product is coadministered with preparation containing Aconite Root.

(3) Drug Interactions

Precautions for coadministration (Keishikaryojutsubuto should be administered with care when coadministered with the following drugs.)

Drugs	Signs, Symptoms, and Treatment	Mechanism and Risk Factors
(1) Preparations containing Glycyrrhiza (2) Preparations containing glycyrrhizinic acid or glycyrrhizinates	Pseudoaldosteronism is likely to occur, Besides, myopathy is likely to occur as a result of hypokalemia. (Refer to the section "Clinically significant adverse reactions")	Since glycyrrhizinic acid has an accelerating action on the potassium excretion at the renal tubules, an acceleration of decrease in the serum potassium level has been suggested.

(4) 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.

1) Clinically significant adverse reactions

① **Pseudoaldosteronism**: Pseudoaldosteronism such as hypokalemia, increased blood pressure, retention of sodium/body fluid, edema, increased body weight, etc. may occur. The patient should be carefully monitored (measurement of serum potassium level, etc.), and if any abnormality is observed, administration should be discontinued and appropriate measures such as administration of potassium preparations should be taken.

② **Myopathy**: Myopathy may occur as a result of hypokalemia. The patient should be carefully monitored, and if any abnormality such as weakness, convulsion/paralysis of limbs, etc. are observed, administration should be discontinued and appropriate measures such as administration of potassium preparations should be taken.

2) Other adverse reactions

	Incidence Unknown
Hypersensitivity Note 1)	Rash, Redness, Pruritus, etc.
Other	Palpitations, Flush, Numbness of the tongue, Nausea, etc.

Note 1) If such symptoms are observed, administration should be discontinued.

(5) Use in the Elderly

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

(6) Use during Pregnancy, Delivery or Lactation

Use of this product in pregnant woman, women who may possibly be pregnant is not recommended. [Adverse reactions due to Powdered Processed Aconite Root contained in this product are likely to occur.]

(7) Pediatric Use

This product should be administered with care in children. [This product contains Powdered Processed Aconite Root.]

PRECAUTIONS FOR HANDLING

- Store at dry and cool place, protected from direct sunlight.
- Since this product contains natural crude drugs, some differences may be noted in the color or taste, etc. However, there is no change in the effect.

PACKAGING

500g
882g (3.0g x 294 packets)
252g (3.0g x 84 packets)

REQUEST FOR LITERATURE SHOULD BE MADE TO:

Dep. of PMS Information,
Ohsugi Pharmaceutical Co., Ltd.
1-8-6, Yamasaka, Higashisumiyoshi-ku, Osaka 546-0035
050-3776-0358