Brand name: Kansei

RISK CATEGORIES

Second-class OTC drug

CHARACTERISTICS

I This is a medicine to support liver function.

Kansei is an herbal medicine originally

formulated with 12 types of crude drugs including those to improve liver function, those to enhance bile secretion, and those with both effects.

This medicine is recommended for persons who are diagnosed as having mild abnormal hepatic function at health examination, complete medical checkup, etc.







ACTIVE INGREDIENTS

3 pouches of Kansei (6 g) contain 2 g of dried Kansei extract prepared from the following crude drugs.

Jujube Seed...1.46 g,

Ginseng...1.46 g

Gardenia Fruit...1.46 g,

Plantago Seed...1.46 g

Mulberry Bark…1.46 g,

Immature Orange…1.46 g

Amomum Seed...1.46 g,

Cinnamon Bark…0.37 g

Rhubarb…0.37 g,

Mume Fructus...1.46 g

Tachibana Pericarpium...1.46 g,

Artemisia Leaf…1.46 g

Inactive ingredients: Lactose, Carmellose

Sodium

<Precautions>

As this medicine is manufactured from crude drugs, slight difference in color and taste may occur between each product, but will not affect the quality.

For further information:

Distributed by:	TAIHO PHARMACEUTICAL CO., LTD.		
Manufactured and Distributed by:	Tatebayashi Shokakudo Co., Ltd.		
Dosage form	Granules		
Packaging unit	2 g x 1 pouches	2 g x 21 pouches	2 g x 60 pouches
Manufacturer's suggested retail price	JPY 220 (JPY 200 excluding tax)	JPY 2,750 (JPY 2,500 excluding tax)	JPY 6,600 (JPY 6,000 excluding tax)
JAN Code	49684360	4987117219104	4987117219005
Expiration	3 years		

DOSAGE AND ADMINISTRATION

Take the medicine with water or warm water between meals.

Age	Dose	Daily dose
Adults (15 years and over)	1 pouch (2 g)	3 times
Under 15 years	Do not use.	

Between meals mean 2 to 3 hours after meals.

INDICATIONS

Abnormal hepatic function, hepatic hypertrophy, acute/chronic hepatitis, jaundice, cholecystitis

PRECAUTIONS FOR USE

The following precaution should be observed.

(If not observed, it may aggravate the current symptoms or increase the risk of adverse reactions.) Lactating mothers should avoid using the medicine or stop breastfeeding.

Consultation

- 1. The following persons should consult a physician, pharmacist or registered salesperson before using this medicine.
 - (1) Persons under treatment by a physician
 - (2) Pregnant or possibly pregnant women
 - (3) Persons who are physically weak (declining of strength or constitutional weakness)
 - (4) Persons who have gastrointestinal weakness and are prone to diarrhea
 - (5) Persons with a history of rash, redness, itching, etc. due to any drug
 - (6) Persons who are taking the following drugs Cathartic drugs (laxatives)
- 2. The following symptoms may be adverse reactions to the medicine. If any of these symptoms occur, discontinue the use of the medicine immediately, and consult a physician, pharmacist or registered salesperson with this leaflet.

Affected body system	Symptoms
Skin	Rash, redness, itching
Gastrointestinal	Diarrhea accompanied by severe abdominal pain, abdominal pain

^{*} The following serious symptoms occur in rare cases. In such cases, consult a physician immediately.

Name of symptom	Symptoms
Mesenteric vein sclerosis	Abdominal pain, diarrhoea, constipation, abdominal distension, etc. occur repeatedly by long-term taking.

3. The following symptoms may be adverse reactions to the medicine. If any of these symptoms persist or get worse, discontinue the use of the medicine immediately, and consult a physician, pharmacist or registered salesperson with this leaflet.

Soft stool, diarrhea

- 4. If symptoms do not improve after using the medicine for about one month, discontinue the use of this medicine, and consult a physician, pharmacist or registered salesperson with this leaflet.
- 5. When the medicine is to be used continuously for a long term, consult a physician, pharmacist or registered salesperson.