

# Comparing compliance and efficacy of isocaloric oral supplementation using 1.5 Kcal/mL or 1 Kcal/mL sip feeds in mild to moderate malnutrition in children

<b>Submission date</b> 28/04/2011	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 09/05/2011	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 09/05/2011	<b>Condition category</b> Nutritional, Metabolic, Endocrine	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

**Contact name**  
Dr Yoga Devaera

**Contact details**  
Departemen Ilmu Kesehatan Anak Fkui  
Jl Salemba 6  
Jakarta  
Indonesia  
10430

## Additional identifiers

**Protocol serial number**  
N/A

## Study information

**Scientific Title**

Comparing compliance and efficacy of isocaloric oral supplementation using 1.5 Kcal/mL or 1 Kcal/mL sip feeds in mild to moderate malnutrition in children: an open label, parallel, randomised controlled trial

### **Study objectives**

Is compliance and efficacy of high calorie supplementation given orally better than standard one in mild to moderate malnourished children?

### **Ethics approval required**

Old ethics approval format

### **Ethics approval(s)**

Ethical Committee Faculty of Medicine of University of Indonesia approved on April 25th 2011 (224/PT02.FK/43/N/2011)

### **Study design**

Open label parallel randomised controlled trial

### **Primary study design**

Interventional

### **Study type(s)**

Treatment

### **Health condition(s) or problem(s) studied**

Mild to moderate malnutrition

### **Interventions**

Ready to drink oral supplementation 600 kcal for 28 days. Study group will receive 2 bottles each 200 mL, 1.5 kcal/mL product. Controlled group will receive 3 bottles each 200 mL 1kcal/mL product.

The randomisation was done by a statistician not included in this study. He then sealed in the envelopes with consecutive number at front page. If a patient enters the study, we pick number accordingly then check the code inside. The code is A or B. The investigator knows what the codes A and B are for but the statistician doesn't. After completion of the study, the statistician will analyse the data and at the end, we all meet to hear the result from him and disclose the code to the statistician.

### **Intervention Type**

Supplement

### **Phase**

Not Applicable

### **Drug/device/biological/vaccine name(s)**

Isocaloric oral supplementation

### **Primary outcome(s)**

1. Weight gain: measure body weight every week of intervention
2. Product consumption: measure the remainings in bottle(s)
3. Gastrointestinal complaints: questionnaire and bristol stool chart

### **Key secondary outcome(s)**

Calorie intake from solid food before and after intervention: 24 recall diet before (day 1) and after intervention (day 29)

### **Completion date**

30/06/2011

## **Eligibility**

### **Key inclusion criteria**

1. Boys and girls aged 3-5 years
2. Z-score for weight-for-height is -1 to -3 according to the World Health Organisation (WHO 2006) growth standards
3. Child is used to drinking milk
4. Stable health status (in the opinion of the Health Care Professional)
5. Written informed consent from parents/caretakers

### **Participant type(s)**

Patient

### **Healthy volunteers allowed**

No

### **Age group**

Child

### **Lower age limit**

3 years

### **Upper age limit**

5 years

### **Sex**

All

### **Key exclusion criteria**

1. Taking medications which increase appetite (appetite stimulants), such as steroids, cyproheptadine
2. Conditions which need a special diet like major renal and hepatic dysfunction
3. Known cows milk allergy, galactosaemia, major gastrointestinal intolerance (e.g. severe vomiting, severe diarrhoea)
4. Children requiring a fiber-free diet
5. Children with oedema
6. Use of parenteral feeding and/or enteral tube-feeding
7. Investigator's uncertainty about the willingness or ability of the child/caretaker to comply

with the protocol requirements

8. Participation in any other study involving investigational or marketed products concomitantly or within two weeks prior to entry into the study

**Date of first enrolment**

02/05/2011

**Date of final enrolment**

30/06/2011

## **Locations**

**Countries of recruitment**

Indonesia

**Study participating centre**

Departemen Ilmu Kesehatan Anak Fkui

Jakarta

Indonesia

10430

## **Sponsor information**

**Organisation**

University of Indonesia (Indonesia)

**ROR**

<https://ror.org/0116zj450>

## **Funder(s)**

**Funder type**

Industry

**Funder Name**

Nutricia (Indonesia)

## **Results and Publications**

**Individual participant data (IPD) sharing plan**

**IPD sharing plan summary**

Not provided at time of registration