

### **Evaluation of the effectiveness of a brief intervention based on Acceptance** and Commitment Therapy for Irritable Bowel Syndrome non-patients #1-36

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# **Background & Objective**

Irritable Bowel Syndrome (IBS) is a functional gastrointestinal disorder affecting 11.2% of the Previous study's target global population (Lovel & Ford, 2012). We must provide preventive support for IBS non-patients. Acceptance and Commitment Therapy (ACT) was effective for IBS patients (Ferreira et al., 2017). **Objective:** To examine whether a **one-day group ACT program had any benefit for IBS non-**Present study's target patients.

## Method

#### **Participants**

**IBS non-patients** (n = 26; male n = 10; mean age 19.09 years)

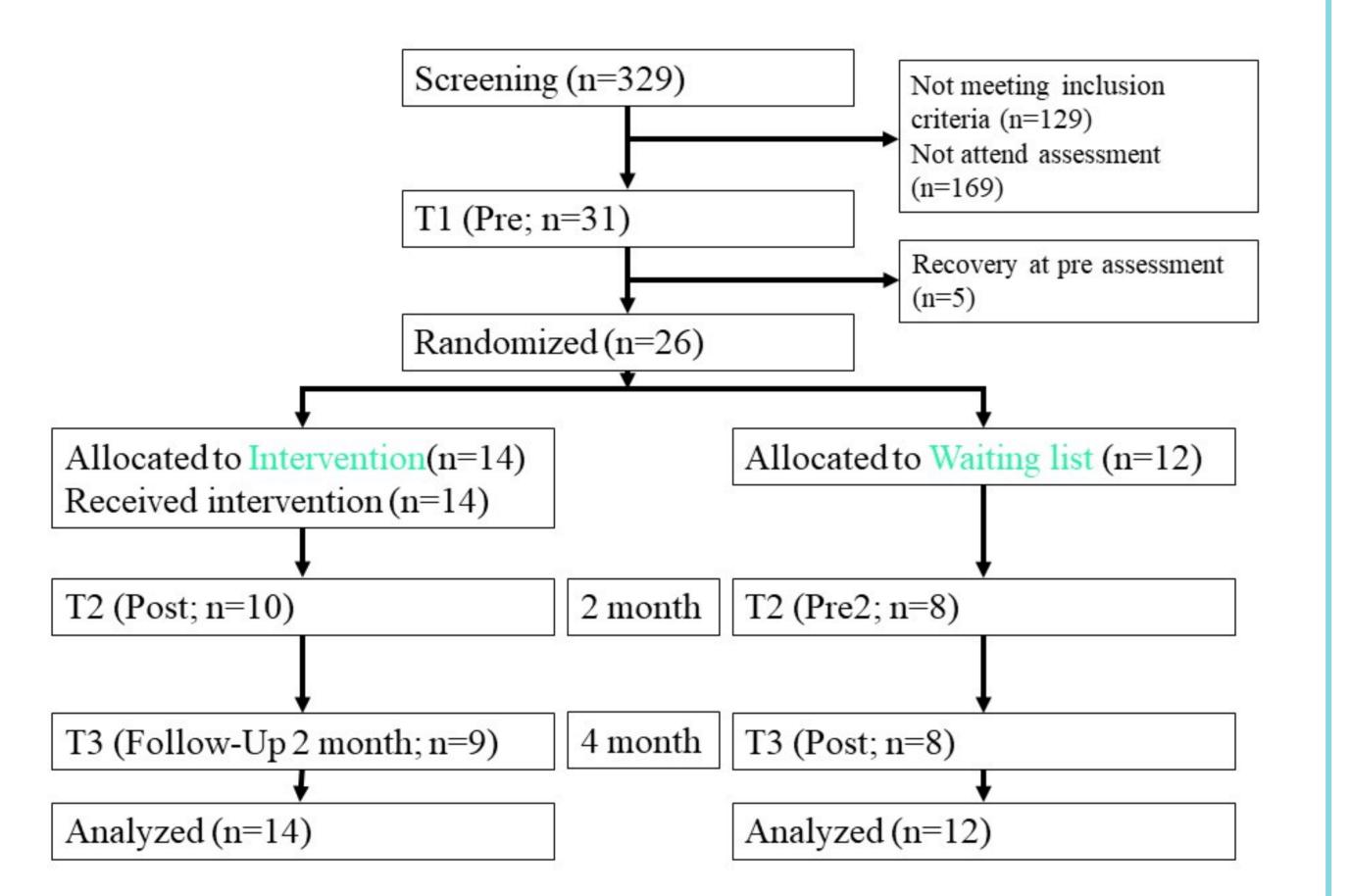
#### Results **Primary outcomes** d (pre-post) 0.95 350 <sub>I</sub> **IBSSI** This measure

- A total of 26 undergraduates who scored above the clinical cutoff on the IBS-SI at screening.
- This group was not under medical care and did not have organic disease as suggested by the presence of warning symptoms.

### **Treatment Protocol**

#### **ACT for Irritable Bowel Syndrome**

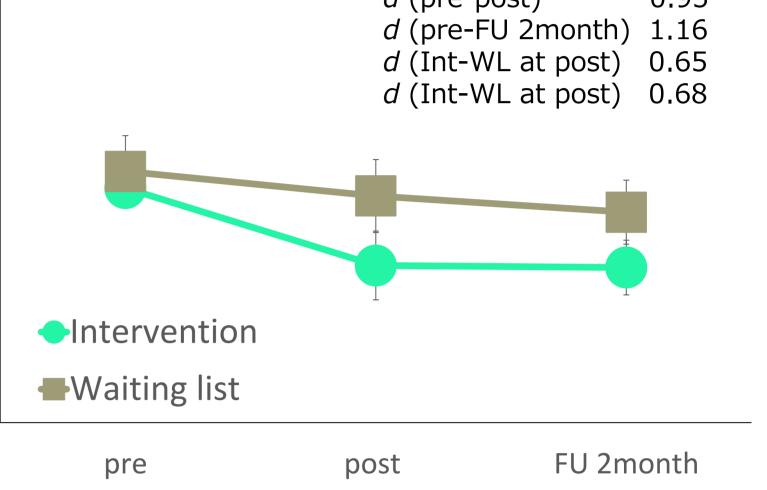
We used "ACT for IBS" (Ferreira & Gillanders, 2012). We translated this protocol into Japanese. We used the workshop protocol and worksheet and the workbook "Get out your mind".



q 005 Da( showed significant time main **0** 250 effect (*F*[2,48] = **4.45**), with no significant group main effect 200 (F[1,24] = 1.94) and interaction **p** 150 0 6 100 (F[2,48] = 1.01). Cohen's d within-group was large from SI 50 IBS pre to post (d = 0.95) and pre to FU (d = 1.16)

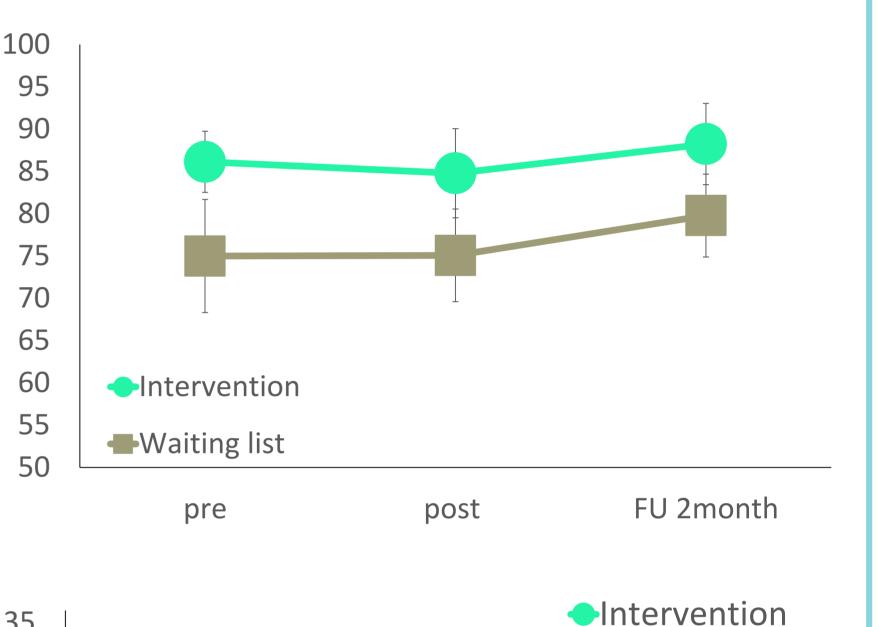


(po IBSQOL This measure did not show significant change g (time main effect; F[2,48] = 1.02: 00 group main effect; F[1,24] = 2.38: interaction; F[2,48] = 0.39). (ba SF-36 This measure did SQOL not show significant interaction in all three summary scores (PCS, F[2,48]



IBS patients

**IBS** non-patients



35

Intervention

-Waiting list

post

FU 2month

**Fig.1** Flow chart of this trial

### **Outcomes**

#### Primary outcome

- **IBSSI** (Irritable Bowel Syndrome Severity Index) Secondary outcomes
- **IBSQOL** (Irritable Bowel Syndrome Quality of Life)
- **SF-36** (Medical Outcomes Study Short Form 36)
- **BDI** (Beck Depression Inventory II)
- **STAI** (State-Trait Anxiety Inventory)

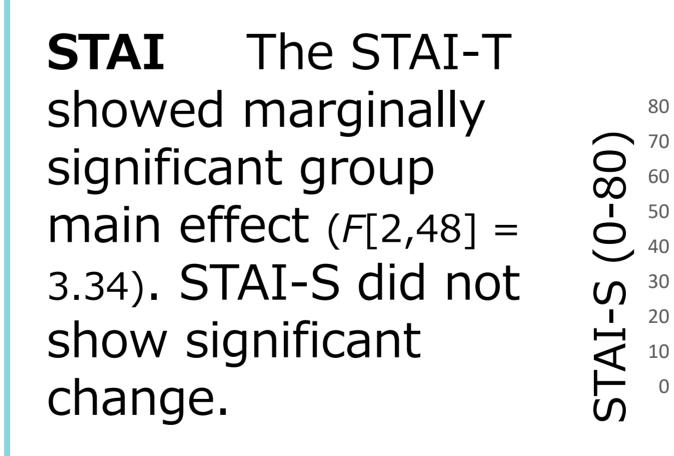
#### **Process measures**

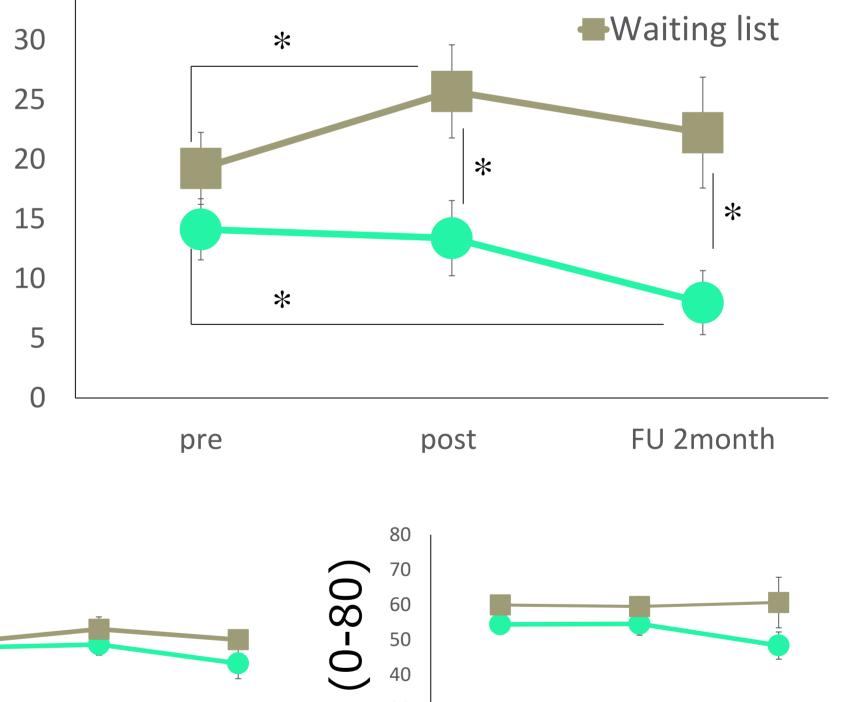
- **AAQ** (Acceptance and Action Questionnaire II)
- **CFQ** (Cognitive Fusion Questionnaire) **FFMQ** (Five-Facet Mindfulness Questionnaire)

#### F[2,48] = 0.88).

= 0.18: MCS, *F*[2,48] = 0.37: RCS,

This measure showed  $\_$ BDI 9 significant interaction  $\mathbf{M}$ (interaction; F[2,48] = 10.17). Post hoc analyses showed **H**<sup>15</sup> intervention group improved **I**0 10 Ш pre to FU and WL group worsened pre to post.





#### **Process measures**

These measures did not show significant interaction. (AAQ, F[2,48] = 1.67: CFQ, F[2,48] = 0.50: FFMQ, F[2,48] = 1.05)

60

50

10

pre

AI

S

Intervention

-Waiting list

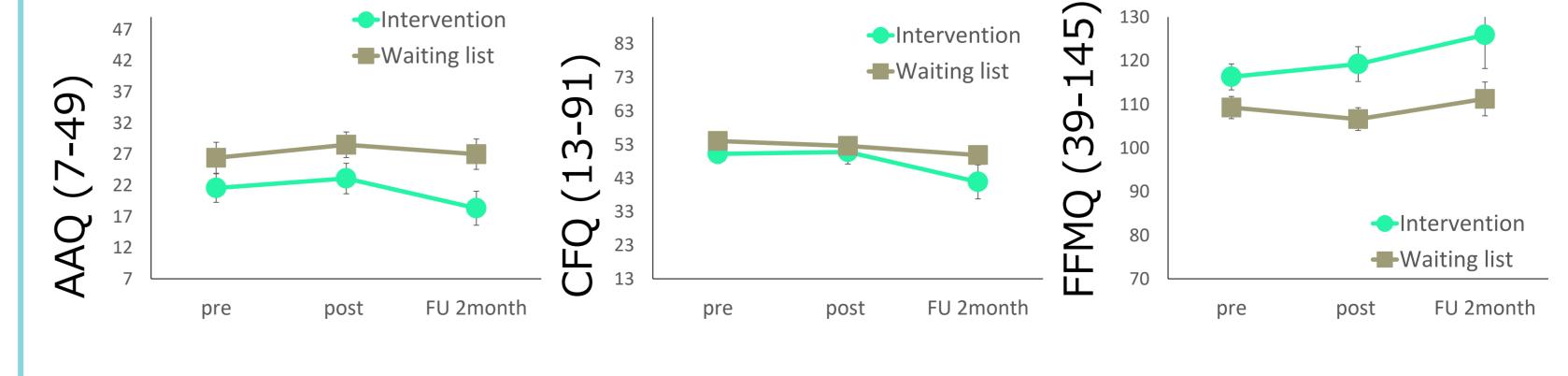
post

pre

FU 2month

### Analysis

We used **repeated ANOVA for pre, post and follow up** data. For missing data, we used the Last Observation Carried Forward imputation method. Data from pre-FU were used to calculate effect size in the intervention group.



## Discussion

This study showed that a one-day group ACT program for IBS non-patients had preventive effects. The intervention group showed improvement of depressed mood and medium to large effect size in IBS symptom severity. There was concern that the treatment process was unclear because the intervention group did not show improvement in process measures. We suggested that a one-day session was not enough to promote continuous exercise to improve psychological flexibility.