Responsive Complementary Feeding in Bangladesh

Submission date Recruitment status Prospectively registered 10/07/2007 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 04/09/2007 Completed [X] Results Individual participant data **Last Edited** Condition category 17/05/2019 Nutritional, Metabolic, Endocrine

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

ICDDR,B #2006-007

Study information

Scientific Title

Responsive Complementary Feeding in Bangladesh

Acronym

CRF

Study objectives

- 1. Mothers in the intervention will show more responsive feeding than the comparison group
- 2. Children in the intervention will show more self-feeding, take more mouthfuls of food, and gain more weight than comparison children

Ethics approval required

Old ethics approval format

Ethics approval(s)

Received from the International Centre for Diarrhoeal Diseases Research Bangladesh ICDDR,B) in February 2006 (ref: #2006-007).

Study design

Cluster randomised controlled trial, with villages being the clusters randomly assigned to intervention or control. Mothers are not blind; research assistants are blind.

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Quality of life

Participant information sheet

Health condition(s) or problem(s) studied

Malnutrition

Interventions

Five weekly group sessions and booster session given by trained village peer educators to mothers and their children using a Manual for Responsive Feeding. Mothers observe a demonstration and then practice the behaviour with their child. Group discussions on how to handle feeding problems, such as refusals, without forceful feeding. Control mothers will receive a reminder of their nutrition education on foods to feed.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

- 1. Mother responsive feeding
- 2. Child weight, self-feeding, and number of mouthfuls eaten

The time points for measurements are as follows: baseline, 2 weeks post intervention, 4 months post intervention.

Secondary outcome measures

- 1. Foods fed and messages recalled
- 2. Possibly other maternal feeding behaviours such as non-responsive encouragement and speech

The time points for measurements are as follows: baseline, 2 weeks post intervention, 4 months post intervention.

Overall study start date

01/05/2007

Completion date

30/11/2007

Eligibility

Key inclusion criteria

- 1. Living in three unions in Jaldhaka, in the north of Bangladesh
- 2. Mothers and their children 8 to 20 months of age
- 3. Must have started complementary food

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

200

Total final enrolment

202

Key exclusion criteria

Children too ill or disabled to feed themselves.

Date of first enrolment

01/05/2007

Date of final enrolment

Locations

Countries of recruitment

Bangladesh

Canada

Study participating centre Department of Psychology Montreal Canada H3A 1B1

Sponsor information

Organisation

International Centre for Diarrhoeal Diseases Research (Bangladesh)

Sponsor details

Mohakhali (or GPO 128) Dhaka Bangladesh 1212

Sponsor type

Research organisation

Website

http://www.icddrb.org/

ROR

https://ror.org/04vsvr128

Funder(s)

Funder type

Government

Funder Name

Department for International Development in Bangladesh (Bangladesh) (grant no.: 00479)

Funder Name

Social Science and Humanities Research Council of Canada (Canada) (grant no.: 861-2006-0033)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/10/2008	17/05/2019	Yes	No