

# An mHealth Tool to improve community health agent performance for child development: a cluster-randomized controlled trial in Peru

<b>Submission date</b> 29/11/2018	<b>Recruitment status</b> No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 04/12/2018	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 10/01/2024	<b>Condition category</b> Other	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

### Background and study aims

Cultivating good child development creates long-term impact on the well-being of the individual and society. Peru has a high level of many risk factors that are associated with poor child development, including chronic malnutrition, anemia, and poor education opportunities. The use of 'community health agents' (CHA) can improve child developmental outcomes through health promotion and education, referral to health services, and surveillance. An effective CHA program can improve cognitive development, child nutrition, and prevention of diseases for populations in low-resource settings. The performance of CHAs can be enhanced by providing them information and communication technology (ICT) to use for their program activities. The current study will evaluate the impact of a tablet-based application (The Child Health Education and Surveillance Tool – The CHEST) that intends to improve the performance of CHAs during their home visits with children under 4.

### Who can participate?

Study participants will include community health agents, the primary caregiver of a child below the age of 4, and children under the age of 4. All community health agents, caregivers, and children in the intervention communities will be invited to receive the intervention. The primary caregiver of the child must be over the age of 16 and speak Spanish. The child will be excluded from the study (but not the intervention) if they have a developmental disability.

### What does the study involve?

The community health agents in the intervention group will receive a tablet pre-loaded with The CHEST Application that will guide them through the steps to execute an effective home visit and provide videos and images to help teach key health messages. The CHEST will organize their caseload and provide a mechanism to record and report child health indicators. The community health agents in the control group will not receive a tablet, and so will continue conducting their activities as community health agents as per the status quo. The outcomes in the intervention group will be compared to the outcomes in the control group. The primary outcomes include early childhood development scores, prevalence of anemia, and prevalence of chronic malnutrition.

What are the possible benefits and risks of participating?

The children in the intervention group are expected to experience improved early childhood development outcomes and improved nutrition due to their caregiver becoming more knowledgeable of healthy child-rearing practices. There are no potential risks or side effects associated with the intervention. The children in the control group will receive the stand benefits from the community health agent program without the additional ICT tool.

Where is the study run from?

The study is being conducted in communities in the country of Peru, department of Loreto, province of Maynas, and districts of Mazan, Puchana, and Las Amazonas. The study is being administered by the Peruvian research organization, Elementos. The home office is located at Jiron Domeyer 105, Barranco, Lima, Peru.

When is the study starting and how long is it expected to run for?

Recruitment for the study will begin in March, 2019 and the intervention will be implemented in May, 2019. The intervention will operate in the communities for 12 months before the follow-up survey is conducted. The study is expected to be completed in June, 2020.

Who is funding the study?

The study is funded by the grant, Saving Brains, by Grand Challenges Canada.

Who is the main contact?

Christopher Westgard  
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## Contact information

### Type(s)

Scientific

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## Additional identifiers

### Protocol serial number

15463-18

## Study information

**Scientific Title**

An mHealth Tool to improve the performance of community health workers during home visits to improve child development and nutrition. A cluster-randomized controlled trial in Peru

**Acronym**

Chest Project

**Study objectives**

1. The digital health device will improve the performance of community health agents during their home visits with children, as measured by doses, content, and inter-personal relationships.
2. Improved performance by community health agents will increase knowledge acquisition and healthy child-rearing practices by care-givers.
3. Improved knowledge and healthy practices by caregivers will improve the health and development of the child that receive the intervention.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

The Institutional Ethics Committee of the National Hospital of Mother and Children "San Bartolome", 08/11/2018, ref. 15463-18.

**Study design**

Quasi-experimental, cluster randomized controlled trial.

**Primary study design**

Interventional

**Study type(s)**

Prevention

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

Early childhood development

**Interventions**

The intervention group will receive a training workshop on how to use the digital health tool and each community health agent will receive an electronic tablet with the CHEST Application pre-loaded. The community health agents will utilize the tablet to guide them through their home visits with mothers of children less than 4 years old in their communities. The intervention group and control group will receive the baseline survey and follow-up survey, which includes early childhood development evaluations, anemia diagnostic, and a KAP survey. The control group will not receive an intervention. The community health agents in the control group will continue to conduct home visits and other activities in their community, as per the status quo. One year after the implementation of the intervention, the follow-up surveys will be conducted. For the selection of the sample populations, communities will be treated as clusters and paired 1-1 based on propensity score matching. The propensity score will be determined by the size of the community, distance from the capitol, and type of health center in the community. One cluster in each pair will be randomly assigned to the intervention group and the other cluster to the control group. The randomization will be conducted by a random number generator.

**Intervention Type**

Device

**Phase**

Not Applicable

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

Not provided at time of registration

**Primary outcome(s)**

1. Child development scores will be measured using the Caregiver Reported Early Development Instrument (CREDI) at baseline and 12 weeks.
2. Anemia prevalence in children will be evaluated by the local health center using the Hemocue at baseline and 12 weeks.
3. Chronic malnutrition prevalence in children will be assessed by the local health center during the children's growth monitoring check-ups, continuously throughout the study. The data will be gathered and analyzed at baseline and 12 weeks.

**Key secondary outcome(s)**

Caregivers:

1. Knowledge evaluation scores will be measured using a knowledge exam created by the study team, at baseline and 12 weeks.
2. The frequency of which caregivers bring their child to growth monitoring check-ups will be evaluated by data obtained from the local health center, at baseline and 12 weeks.

Community health agents:

1. The performance evaluation scores of community health agents will be measured using an observational checklist created by the author, at baseline and 12 weeks.
2. Self-efficacy scores of the community health agents will be measured using a qualitative questionnaire at baseline and 12 weeks.
3. The number of home visits made by community health agents will be measured using the CHEST Application consistently throughout the intervention.

**Completion date**

01/08/2020

**Eligibility****Key inclusion criteria**

Children:

1. Children aged 6 to 36 months.

Caregivers:

1. Primary caregiver of the child is older than 16 years.
2. The caregiver is willing to receive visits from the CHA.

**Healthy volunteers allowed**

No

**Age group**

Mixed

**Sex**

All

**Total final enrolment**

500

**Key exclusion criteria**

Caregivers:

1. Families where the main caregiver or the child does not speak Spanish.

Children:

1. Children with mental or physical disabilities.

**Date of first enrolment**

01/02/2019

**Date of final enrolment**

30/06/2019

## Locations

**Countries of recruitment**

Peru

**Study participating centre**

**Community**

Maynas, Loreto, Peru

Iquitos

Peru

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## Sponsor information

**Organisation**

Elementos

## Funder(s)

**Funder type**

Government

**Funder Name**

Grand Challenges Canada

**Alternative Name(s)**

Grands Défis Canada, gchallenges, Grand Challenges Canada / Grands Défis Canada, grandchallengescanada, GCC

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

**Location**

Canada

## Results and Publications

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

The datasets generated during and/or analysed during the current study will be stored in the publically available repository, figshare (<https://figshare.com/>). De-identified participant data, including early childhood development evaluation scores, anemia results, chronic malnutrition results, KAP survey results, and CHA performance evaluations, will be stored in the repository. The data will be available through the repository for 10 years. Data from participants will only be made available if consent was provided by the participant through the informed consent process.

**IPD sharing plan summary****Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>		22/09/2022	10/01/2024	Yes	No
<a href="#">Protocol article</a>	protocol	06/11/2019	15/01/2021	Yes	No