Assessing the effectiveness of front-of-pack food labels in Kenya

Submission date	Recruitment status	Prospectively registered		
28/11/2023	No longer recruiting	[_] Protocol		
Registration date	Overall study status	[] Statistical analysis plan		
06/12/2023	Completed	[X] Results		
Last Edited 24/02/2025	Condition category Other	Individual participant data		

Plain English summary of protocol

Background and study aims

Kenya has a high rate of unhealthy food consumption because there are no comprehensive policies in place to regulate the food environment. The main objective of the study is to evaluate the effect of different front-of-pack labels on participants' identification of products high in nutrients of concern, identification of unhealthy foods, and intention to purchase unhealthy foods and beverages.

Who can participate?

Adults over the age of eighteen years who are not employed by marketing firms, food and beverage companies, health promotion organizations, the tobacco industry, or the food and beverage sector (grocery stores, eateries, etc.)

What does the study involve?

This research involves the comparison and identification of a front-of-pack label that most effectively improves nutrient comprehension among the Kenyan population.

What are the possible benefits and risks of participating? No individual benefits are anticipated from participation in this study. The proposed study is expected to provide evidence for public policy, and participants will be duly informed of this. The study poses minimal risks to the participants.

Where is the study run from? African Population and Health Research Center (APHRC) (Kenya)

When is the study starting and how long is it expected to run for? March 2022 to January 2024

Who is funding the study?

- 1. International Development Research Centre (Canada)
- 2. Rockefeller Foundation (USA)

Who is the main contact? Dr Gershim Asiki, gasiki@aphrc.org Shukri Mohammed, smohamed@aphrc.org

Contact information

Type(s) Principal Investigator

Contact name Dr Gershim Asiki

ORCID ID http://orcid.org/0000-0002-9966-1153

Contact details

African Population and Health Research Center (APHRC) P.O BOX, 10787 Nairobi Kenya 00100 +254 714712834 gasiki@aphrc.org

Type(s) Public, Scientific

Contact name Dr Shukri Mohammed

ORCID ID http://orcid.org/0000-0002-8693-1943

Contact details

African Population and Health Research Center (APHRC) P.O BOX, 10787 Nairobi Kenya 00100 +254 713089457 smohamed@aphrc.org

Additional identifiers

EudraCT/CTIS number Nil known

IRAS number

ClinicalTrials.gov number Nil known

Secondary identifying numbers ERC/P1323/2022

Study information

Scientific Title

Assessing the effectiveness of front-of-pack food labels in Kenya: A randomized controlled trial

Acronym

FEP-ACTION

Study objectives

The presence of a Front-of-Pack Label will lead to a higher rate of correct identification of unhealthy foods by consumers compared to a condition where no label is present.

Ethics approval required

Ethics approval required

Ethics approval(s)

Approved 24/04/2023, AMREF Ethics and Scientific Review Commitee (Nairobi, Nairobi, 00100, Kenya; +254(0) 206994000; erc.kenya@amreg.org), ref: ERC/P1323/2022

Study design Three-arm randomized controlled trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) Community, Retail/food outlet

Study type(s) Prevention, Efficacy

Participant information sheet See outputs table

Health condition(s) or problem(s) studied

Identification of unhealthy foods using different front-of-pack labels in urban and rural settings in Kenya

Interventions

The intervention involves two phases: a control (placebo) phase and an experimental phase. Each participant will be exposed to both the control and experimental phases, with the aim of assessing the within- and between-subject effects.

During the control (placebo) phase, all participants will view product images displayed on mock packages without any front-of-pack label. Subsequently, they will respond to a set of questions. In the experimental phase, the same participants will be randomly assigned to one of three label conditions (intervention or experiment). They will view the same product images, but this time the images will be presented with a front-of-pack label (the intervention), and they will be asked to respond to the same identical set of questions as in the control phase.

The study duration is approximately is one month, and there will be no subsequent follow-up for the study arms.

In terms of randomization, a manual process utilizing an Excel sheet will be used to randomly allocate study participants to one of three front-of-pack labels. The initial step involves using the sample allocation for counties and sub-counties to ensure an equal distribution of labels among participants in those specific counties. Participants will then be assigned labels randomly based on their specific unique IDs. This process will be imported into the data collection equipment, and random symbols will appear for each participant during the interviews.

Intervention Type

Behavioural

Primary outcome measure

Measured during the control and intervention phases:

1. Identification of foods high in nutrients of concern measured by presenting products and asking the participant to identify the products that are high in salt, sugar, and fat (yes, no, or don' t know) and identify the products as unhealthy (healthy or unhealthy)

2. Identification of unhealthy foods measured by presenting paired images of products and asking the participant to identify the product higher in salt, sugar, or fat and determine the unhealthier product.

3. Consumers intention to purchase unhealthy foods measured using a four-point Likert scale.

Secondary outcome measures

There are no secondary outcome measures

Overall study start date 01/03/2022

Completion date 31/01/2024

Eligibility

Key inclusion criteria Participants >18 years and those who purchase packaged products.

Participant type(s) Resident, Population

Age group

Adult

Lower age limit 18 Years

Upper age limit 69 Years

Sex Both

Target number of participants 2,185

Total final enrolment 2196

Key exclusion criteria

1. Participants aged less than 18 years 2. Participants working in the marketing industry, food and beverage companies, health promotion, tobacco industry, and food and beverage industry (e.g., supermarkets, restaurants, retail companies)

Date of first enrolment 29/11/2023

Date of final enrolment 22/12/2023

Locations

Countries of recruitment Kenya

Study participating centre

Nairobi County Embakasi North and South counties, and Langata sub counties Nairobi Kenya 00100

Study participating centre Mombasa County Mvita and Kisauni Sub Counties Mombasa Kenya 80100

Study participating centre Garrisa County Garrisa township and Fafi sub counties. Garrisa Kenya 7 0100

Study participating centre Kisumu county Nyando and Kisumu central sub counties Kisumu Kenya 4 0100

Sponsor information

Organisation

International Development Research Centre

Sponsor details

45 O'Connor St, Ottawa, ON K1P 1A4 Ottawa Canada K1G 3H9 +1 (613) 236-6163 info@idrc.ca

Sponsor type

Government

Website

http://www.idrc.ca/EN/Pages/default.aspx

Organisation Rockefeller Foundation

Sponsor details

420 Fifth Avenue New York United States of America NY 10018 +1 20 4987500 info@rockfound.org

Sponsor type Research organisation

Website https://www.rockefellerfoundation.org/

ROR https://ror.org/03sfkwk85

Funder(s)

Funder type Government

Funder Name International Development Research Centre

Alternative Name(s) Centre de recherches pour le développement international, IDRC, CRDI

Funding Body Type Government organisation

Funding Body Subtype Local government

Location Canada

Funder Name Rockefeller Foundation

Alternative Name(s) Centre de recherches pour le développement international, IDRC, CRDI

Funding Body Type Government organisation

Funding Body Subtype

Local government

Location Canada

Results and Publications

Publication and dissemination plan

We plan to publish the findings of this study in a peer-reviewed journal. The findings will also be disseminated to various audiences, including government stakeholders, civil society, communities where the data will be collected, scientific conferences, etc.

Intention to publish date

30/12/2024

Individual participant data (IPD) sharing plan

The datasets generated during and/or analyzed during the current study will be available upon request from gasiki@aphrc.org. The shared data will be anonymized and have no identifiers linking it back to the study participants. Broad consent will be sought from the study participants thus allowing for future analysis.

IPD sharing plan summary

Available on request

Study outputs					
Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet			01/12/2023	No	Yes
Basic results		24/10/2024	28/10/2024	No	No
Results article		10/11/2024	24/02/2025	Yes	No