# Assessing the use of Mapping for Health (M4H) data for immunisation programme implementation and associated impact on coverage, equity, and cost-effectiveness in the Democratic Republic of Congo

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>		
25/10/2021	No longer recruiting	[X] Protocol		
Registration date	Overall study status	Statistical analysis plan		
03/11/2021	Completed	[X] Results		
Last Edited	Condition category	[] Individual participant data		
28/01/2025	Other			

## Plain English summary of protocol

Background and study aims

There are many children who do not receive immunizations because they live in hard to reach locations that are often not included on maps. The use of maps can help to identify children who are often missed to make sure that they are included in immunization program plans. This study will measure the acceptance and use of maps as part of the planning and implementation of the immunization program in Democratic Republic of Congo (DRC) to see if they can help increase the number of children who are vaccinated, especially those who are the hardest to reach. The goal of this study is to evaluate the impact of the Mapping for Health programme, which produces and provides geospatial data with the aim of improving immunisation coverage and equity, especially in the identification of previously missed settlements.

# Who can participate?

Essential Programme for Immunisation administrators, supervisors, and vaccinators and associated individuals engaged in the generation and/or use of M4H geospatial data for immunisation microplanning, macroplanning, and service delivery.

#### What does the study involve?

EPI managers, supervisors, and vaccinators will be invited to be interviewed for the study. Those who agree to will be interviewed for 30-45 minutes about their experiences with the use of maps within the program planning and implementation.

What are the possible benefits and risks of participating? None

Where is the study run from?
Kinshasa School of Public Health (Democratic Republic of Congo)

When is the study starting and how long is it expected to run for? January 2020 to December 2023

Who is funding the study?
Gavi, the Vaccine Alliance (Switzerland)

Who is the main contact?
Dr. Patricia Mechael, patty@healthenabled.org

# Contact information

## Type(s)

Public

#### Contact name

Dr Patricia Mechael

#### **ORCID ID**

http://orcid.org/0000-0002-1034-0549

#### Contact details

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# Additional identifiers

# EudraCT/CTIS number

Nil known

#### **IRAS** number

# ClinicalTrials.gov number

Nil known

# Secondary identifying numbers

Nil known

# Study information

#### Scientific Title

Large-scale effectiveness study of the effective use of Mapping for Health Data on immunisation coverage and equity in the Democratic Republic of Congo

# Study objectives

The effective use of M4H geospatial data can increase immunisation coverage and equity through the identification of missed settlements (and zero dose children) and optimization of vaccination strategies and supply distribution as well as a gender-specific interventions in a subset of Health Zones and Health Areas in Kasai and Kinshasa.

Study aim 1: Conduct a process evaluation to understand the program and implementation context, and identify the mechanisms through which geospatial data use influences immunisation coverage

Study aim 2: Conduct a quasi-experimental design study in three provinces (two intervention-one with gender activities, one without gender activities, and one control) to determine the associated effects of the acceptance and use of M4H data by Health Zones and Health Areas on immunisation coverage and equity (especially zero-dose children).

Study aim 3: Determine the incremental cost-effectiveness of geospatial data use as compared to the status quo

# Ethics approval required

Old ethics approval format

### Ethics approval(s)

Approved 27/07/2021, Kinshasa School of Public Health Internal Review Board (BP 11850 Kinshasa, DRC; +243 817 493194; espsec\_unikin@yahoo.fr), ref: none provided

## Study design

Single-centre three-province quasi-experimental interventional study

### Primary study design

Interventional

# Secondary study design

Cluster randomised trial

## Study setting(s)

Other

# Study type(s)

Prevention

# Participant information sheet

Not available in web format, please use the contact details to request a patient information sheet.

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

Routine immunisation

#### **Interventions**

There is a control province and two intervention provinces, where sites will be stratified and then randomly selected for inclusion in the intervention strength study based on immunisation coverage and equity at baseline, urban/rural, and connectivity status.

#### Intervention:

Immunisation microplans with geospatial data, updated immunisation targets based on geospatial modeled population estimates, and population mobility data.

#### Control:

Standard practice or status quo in the control province - immunisation microplans without geospatial data and immunisation targets based on census data.

The overall structure of the research study includes three provinces - one province selected for inclusion to assess M4H geospatial data interventions with gender-specific activities in a subset of Health Zones and Health Areas, namely Kasai; another M4H province selected to assess effects of M4H geospatial data interventions without gender, namely Lomami; and a control site to compare both M4H sites, namely Kasai Central. In addition, the protocol includes prioritization and segmentation of primary research participants, Essential Programme for Immunisation (EPI) staff and affiliates, for the intervention strength survey and qualitative indepth interviews (where possible engaging with equal numbers of male and female EPI managers, supervisors, and vaccinators) as well as in the secondary analyses of immunisation coverage and equity surveys (gender-disaggregated analyses of birth cohorts). Each study arm will be segmented into the following categories (urban, rural, per-urban, conflict, and high/low /moderate connectivity) and a random selection of Health Areas will be chosen for inclusion in the intervention strength survey and qualitative in-depth interviews. Complementary qualitative data collection will be conducted in the form of direct observation and in-depth interviews at the National, Province, and Health Zone levels. Additional rapid ethnographic field work will be conducted to assess the gender -specific interventions.

#### Intervention Type

Mixed

#### Primary outcome measure

Measured at the end of the study:

- 1. Acceptance of M4H geospatial data in micro planning and macro planning- measured as the percent of Provinces and Health Zones EPI Managers who state that they intend to use M4H geospatial data in EPI programme planning and implementation
- 2. Use of M4H geospatial data in micro planning and macro-planning measured as the percent of Health Zone EPI Managers who include the M4H geospatial data in their micro plans
- 3. Use of M4H data in immunisation programme delivery measured as the percent of Health Area EPI supervisors and vaccinators who use the M4H geospatial data to manage the implementation of the routine immunisation programme/ all eligible Health Area EPI supervisors and vaccinators
- 4. Effects of M4H data on immunisation programme outcomes measured as change in immunisation coverage and equity through secondary analyses of immunisation survey data from baseline to end-line (pre/ post-intervention) in Health Areas as correlated to the percentages of acceptance and use of M4H geospatial data
- 5. Cost-effectiveness of M4H data on immunisation programme outcomes measured in Lives Saved using the Lives Saved Tool and Disability-Adjusted Life-years alongside change in immunisation coverage and equity through secondary analyses of immunisation survey data from baseline to end-line (pre/ post-intervention) in Health Areas as correlated to the percentages of acceptance and use of M4H geospatial data

#### Secondary outcome measures

Gender-associated affects in the creation, acceptance, and use of M4H geospatial datameasured through a Rapid Ethnographic Study of the gender intervention and through genderrelated questions and disaggregation of respondents by gender as correlated to the percentages of acceptance and use of M4H geospatial data at the end of the study

# Overall study start date

01/01/2020

# Completion date

31/12/2023

# **Eligibility**

#### Key inclusion criteria

Essential Programme for Immunisation administrators, supervisors, and vaccinators and associated individuals engaged in the generation and/or use of M4H geospatial data for immunisation microplanning, macroplanning, and service delivery.

### Participant type(s)

Health professional

#### Age group

Adult

#### Sex

Both

# Target number of participants

~500

#### Total final enrolment

111

#### Key exclusion criteria

Those who do not provide consent to participate in the study.

#### Date of first enrolment

01/11/2021

#### Date of final enrolment

31/12/2023

# Locations

#### Countries of recruitment

Congo, Democratic Republic

# Study participating centre Kinshasa School of Public Health

H8Q3+2HV Kinshasa Congo, Democratic Republic

# Sponsor information

# Organisation

Gavi

#### Sponsor details

Global Health Campus Chemin du Pommier 40 Le Grand-Saconnex Switzerland 1218 +41 22 909 6500 info@gavi.org

#### Sponsor type

Charity

#### Website

http://www.gavi.org/

#### **ROR**

https://ror.org/0141yg674

# Funder(s)

# Funder type

Charity

#### **Funder Name**

**GAVI Alliance** 

#### Alternative Name(s)

Gavi, Gavi The Vaccine Alliance

# **Funding Body Type**

Government organisation

# Funding Body Subtype

Trusts, charities, foundations (both public and private)

#### Location

Switzerland

# **Results and Publications**

#### Publication and dissemination plan

The aim will be to create one comprehensive report with all research findings and submit a summary paper to a peer-reviewed journal.

# Intention to publish date

31/12/2023

# Individual participant data (IPD) sharing plan

All data generated or analysed during this study will be included in the subsequent results publication

#### IPD sharing plan summary

Published as a supplement to the results publication, Other

# **Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<u>Protocol file</u>		21/10/2021	29/10/2021	No	No
Funder report results		01/11/2023	17/06/2024	No	No
Results article		24/01/2025	28/01/2025	Yes	No