## Supplementary consent 1

#### 1. Informed Consent Form for Household Head

## Ethiopian Food and Nutrition Strategy (FNS) Baseline Survey

<u>Investigator(s)</u>: Dr.Masresha Tessema (PI), Meseret W/Yohannes, Dr. Meron Girma, Alemnesh Petros, Dr Aregash Samuel, Arnaud Laillou, Stanley Chitekwe, Kaleab Baye, Ramadhani Noor, Anne Sophie Donze and other co-authors

You are being invited to take part in this research because you are head of household. There are about [16596] households taking part in this research. We will ask you about household characteristics, and socio-economic status of your household. We will also assess dietary intake, anthropometric status, nutrition sensitive and specific indicators and micronutrient status of your household member.

# Box 1. Taking part in this research is voluntary

You can refuse to take part in this study.

You can withdraw your participation from the study at any time

## Information related to the study

The FNS baseline survey will be conducted in the 12 regions of Ethiopia. The study population will be children age 0-59 months having caregivers/mothers, school-age children (6-12 years), adolescent girls (10-19 years), reproductive-age women (15-49 years), pregnant and lactating women, and household head. The indicators that will be collected for the survey will be dietary intake, anthropometric status, nutrition sensitive and specific indicators and micronutrient status.

The expected possible adverse effects: there is no adverse effect by participating in this study

The objective of this research: to produce information on anthropometric status, dietary intakes, and micronutrient status of different population groups in Ethiopia, and assess the coverage of direct and indirect nutrition interventions.

Study design: A population-based, cross-sectional design		
The schedule of the study: The study will be conducted from July, 2021 to April, 2023		
Foreseeable risks and expected benefits arising from participation in the study		
Foreseeable risks	Expected benefits	
Risks to study participants for involvement in the	The findings of the study will help the ministry of	
l	health and other stakeholders engaged in nutrition	

with COVID pandemic. Interviewers will b minimize this risk and will use appropriate measures.	e trained to to improve and/or design appropriate health and nutrition intervention programs in the country.	
Occurrences that may take place during the	e study period	
Occurrences	How to manage	
Withdrawal of volunteers from the study	In such a case, we would respect the volunteer's decision to withdraw and also get a clear understanding of the reason for	

At the end of the study, you will not be receiving any financial benefits, but will get your results for height, weight, mid upper arm and waist circumference measurements, anemia and goiter status for time you spent and participation.

All data collected from the study will be kept confidential. If you have any questions related to the study you may contact directly Dr. Masresha Tessema who is the project PI.

## The contact persons

- 1. Dr. Masresha Tessema Tel. [+251 919782082] E-mail: [masresha88@gmail.com] or
- 2. [Mr. Ibrahim Kedir] Tel. [+251 911957161] EPHI's IRB

# **Certificate of Consent** I have read the foregoing information. I have an I confirm that the participant was given an opportunity to ask questions and all my quest opportunity to ask questions about the study and have been answered to my satisfaction. I all questions have been answered correctly. I confirm that the consent has been given volunteer to give consent to participate in this research study voluntarily Printed name of the participant Printed name of the person taking the consent Signature of the participant Signature of the person taking the consent Date Date\_ day/month/year day/month/year

# 2. Informed Consent Form for Women of Reproductive Age

## Ethiopian Food and Nutrition Strategy (FNS) Baseline Survey

<u>Investigator(s)</u>: Dr.Masresha Tessema (PI), Meseret W/Yohannes, Dr. Meron Girma, Alemnesh Petros, Dr Aregash Samuel, Arnaud Laillou, Stanley Chitekwe, Kaleab Baye, Ramadhani Noor, Anne Sophie Donze and other co-authors

You are being invited to take part in this research because you are women of reproductive age. There are [16596] households taking part in this research. We will assess your dietary intake, anthropometric status, nutrition sensitive and specific indicators and micronutrient status

## Box 1. Taking part in this research is voluntary

- You can refuse to take part in this study.
- ☐You can withdraw your participation from the study at any time

## Information related to the study

The FNS baseline survey will be conducted in the 12 regions of Ethiopia. The study population will be children age 0-59 months having caregivers/mothers, school-age children (6-12 years), adolescent girls (10-19 years), reproductive-age women (15-49 years), pregnant and lactating women, and household head. The indicators that will be collected for the survey will be dietary intake, anthropometric status, nutrition sensitive and specific indicators and micronutrient status.

The expected possible adverse effects: there is no adverse effect by participating in this study

The objective of this research: to produce information on anthropometric status, dietary intakes, and micronutrient status of different population groups in Ethiopia, and assess the coverage of direct and indirect nutrition interventions.

**Study design:** A population-based, cross-sectional design

**The schedule of the study:** The study will be conducted from July, 2021 to April, 2023

The schedule of the study. The study will be conducted from July, 2021 to April, 2023	
Foreseeable risks and expected benefits arising from participation in the study	
Foreseeable risks	Expected benefits
Risks to study participants for involvement in the	The findings of the study will help the ministry of
coverage survey are low. There may be risks associated	health and other stakeholders engaged in nutrition
with COVID pandemic. Interviewers will be trained to	to improve and/or design appropriate health and
minimize this risk and will use appropriate prevention	nutrition intervention programs in the country.
measures.	

Occurrences that may take place during the study period	
Occurrences	How to manage
Withdrawal of volunteers from the study	In such a case, we would respect the volunteer's decision to withdraw and also get a clear understanding of the reason for their withdrawal

At the end of the study, you will not be receiving any financial benefits, but will get your results for height, weight, mid upper arm and waist circumference measurements, anemia and goiter status for time you spent and participation.

All data collected from the study will be kept confidential. If you have any questions related to the study you may contact directly Dr. Masresha Tessema who is the project Pl.

## The contact persons

- 1. Dr. Masresha Tessema Tel. [+251 919782082] E-mail: [masresha88@gmail.com] or
- 2. [Mr. Ibrahim Kedir] Tel. [+251 911957161] EPHI's IRB

Certificate of Consent	
I have read the foregoing information. I have an	I confirm that the participant was given an
opportunity to ask questions and all my quest	opportunity to ask questions about the study and
have been answered to my satisfaction. I	all questions have been answered correctly. I
volunteer to give consent to participate in this	confirm that the consent has been given
research study	voluntarily
Printed name of the participant	
	Printed name of the person taking the consent
Signature of the participant	
Date	Signature of the person taking the consent
day/month/year	Date
	day/month/year

## 3. Informed Consent Form for Pregnant Women

## Ethiopian Food and Nutrition Strategy (FNS) Baseline Survey

Investigator(s): Dr.Masresha Tessema (PI), Meseret W/Yohannes, Dr. Meron Girma, Alemnesh Petros, Dr Aregash Samuel, Arnaud Laillou, Stanley Chitekwe, Kaleab Baye, Ramadhani Noor, Anne Sophie Donze and other co-authors

You are being invited to take part in this research because you are pregnant women. There are [16596] households taking part in this research. We will assess your, anthropometric status, nutrition sensitive and specific indicators and anemia status

## Box 1. Taking part in this research is voluntary

You can refuse to take part in this study.

You can withdraw your participation from the study at any time

## Information related to the study

The FNS baseline survey will be conducted in the 12 regions of Ethiopia. The study population will be children age 0-59 months having caregivers/mothers, school-age children (6-12 years), adolescent girls (10-19 years), reproductive-age women (15-49 years), pregnant and lactating women, and household head. The indicators that will be collected for the survey will be dietary intake, anthropometric status, nutrition sensitive and specific indicators and micronutrient status.

The expected possible adverse effects: There is no adverse effect by participating in this study

**The objective of this research:** to produce information on anthropometric status, dietary intakes, and micronutrient status of different population groups in Ethiopia, and assess the coverage of direct and indirect nutrition interventions.

Study design: a population-based, cross-sectional design

Occurrences that may take place during the study period

**The schedule of the study:** The study will be conducted from July, 2021 to April, 2023

# Foreseeable risks and expected benefits arising from participation in the study

# Foreseeable risks Risks to study participants for involvement in the coverage survey are low. There may be risks associated with COVID pandemic. Interviewers will be trained to minimize this risk and will use appropriate prevention measures. Expected benefits The findings of the study will help the ministry of health and other stakeholder engaged in nutrition to improve and/or design appropriate health and nutrition intervention programs in the country.

Occurrences	How to manage
Withdrawal of volunteers from the study	In such a case, we would respect the volunteer's
	decision to withdraw and also get a clear
	understanding of the reason for their withdrawal

At the end of the study, you will not be receiving any financial benefits, but will get your results for height, weight, mid upper arm circumference measurements, anemia and goiter status for time you spent and participation.

All data collected from the study will be kept confidential. If you have any questions related to the study you may contact directly Dr. Masresha Tessema who is the PI

## The contact persons

1. Dr. Masresha Tessema

Tel. [+251 919782082] E-mail: [masresha88@gmail.com]

2. [Mr. Ibrahim Kedir] Tel. [+251 911957161]

# **Certificate of Consent** I have read the foregoing information. I have an I confirm that the participant was given an opportunity to ask questions and all my quest opportunity to ask questions about the study and have been answered to my satisfaction. I all questions have been answered correctly. I volunteer to give consent to participate in this confirm that the consent has been given voluntarily research study Printed name of the participant Printed name of the person taking the consent Signature of the participant Date Signature of the person taking the consent Date day/month/year day/month/year

## 4. Informed Consent Form for Preschool Child

## Ethiopian Food and Nutrition Strategy (FNS) Baseline Survey

Investigator(s): Dr.Masresha Tessema (PI), Meseret W/Yohannes, Dr. Meron Girma, Alemnesh Petros, Dr

Aregash Samuel, Arnaud Laillou, Stanley Chitekwe, Kaleab Baye, Ramadhani Noor, Anne Sophie Donze and other co-authors

You are being invited to take part in this research because you are either a mother or caregiver who has a child under the age of 5 years (0-59 months). There are [16596] households taking part in this research. We would collect a sample of your child's dietary information, blood, urine and stool. And, we will also measure your child's height/ length, weight, and mid upper arm circumference. Finally we will assess, your child's eye for bitot spot

## Box 1. Taking part in this research is voluntary

You can refuse to take part in this study.

You can withdraw your participation from the study at any time

## Information related to the study

The FNS baseline survey will be conducted in the 12 regions of Ethiopia. The study population will be children age 0-59 months having caregivers/mothers, school-age children 6-12 years), adolescent girls (10-19 years), reproductive-age women (15-49 years), pregnant and lactating women, and household head. The indicators that will be collected for the survey will be dietary intake, anthropometric status, nutrition sensitive and specific indicators and micronutrient status.

The expected possible adverse effects: there is no adverse effect by participating in this study

The objective of this research: to produce information on anthropometric status, dietary intakes, and micronutrient status of different population groups in Ethiopia, and assess the coverage of direct and indirect nutrition interventions.

Study design: a population-based, cross-sectional design	
The schedule of the study: The study will be conducted from July, 2021 to April, 2023	
Foreseeable risks and expected benefits arising from	participation in the study
Foreseeable risks	Expected benefits
Risks to study participants for involvement in the	The findings of the study will help the ministry of
coverage survey are low. There may be risks	health and other stakeholder engaged in nutrition to
associated with COVID pandemic. Interviewers will	improve and/or design appropriate health and
be trained to minimize this risk and will use	nutrition intervention programs in the country.
appropriate prevention measures.	
Occurrences that may take place during the study period	
Occurrences How to manage	

Withdrawal of volunteers from the study	in such a case, we would respect the volunteer's
	decision to withdraw and also get a clear
	understanding of the
	reason for their withdrawal

At the end of the study, you will not be receiving any financial benefits, but will get your results for height/length, weight, mid upper arm circumference measurements, and anemia for time you spent and participation.

All data collected from the study will be kept confidential. If you have any questions related to the study you may contact directly Dr. Masresha Tessema who is the project principal investigator

## The contact persons

1. Dr. Masresha Tessema

Tel. [+251 919782082] E-mail: [masresha88@gmail.com]

2. [Mr. Ibrahim Kedir] Tel. [+251 911957161] EPHI's IRB

Certificate of Consent		
I have read the foregoing information. I have an	I confirm that the participant was given an	
opportunity to ask questions and all my quest	opportunity to ask questions about the study and	
have been answered to my satisfaction. I	all questions have been answered correctly. I	
volunteer to give consent to participate in this	confirm that the consent has been given	
research study	voluntarily	
Printed name of the participant		
Signature of the participant's parent or guardian	Printed name of the person taking the consent	
Date		
day/month/year	Signature of the person taking the consent Date	
	day/month/year	

## 5. Informed Consent Form for School Age Children

## Ethiopian Food and Nutrition Strategy (FNS) Baseline Survey

**Investigator(s):** Dr.Masresha Tessema (PI), Meseret W/Yohannes, Dr. Meron Girma, Alemnesh Petros, Dr Aregash Samuel, Arnaud Laillou, Stanley Chitekwe, Kaleab Baye, Ramadhani Noor, Anne Sophie Donze and other co-authors

You are being invited to take part in this research because you are either a mother or caregiver who has a school-age child. Among children 6-12 years, we will collect your child's blood, urine and stool.

## Box 1. Taking part in this research is voluntary

You can refuse to take part in this study.

You **can withdraw** your participation from the study at any time

## Information related to the study

The FNS baseline survey will be conducted in the 12 regions of Ethiopia. The study population will be children age 0-59 months having caregivers/mothers, school-age children (6-12 years), adolescent girls (10-19 years), reproductive-age women (15-49 years), pregnant and lactating women, household head. The indicators that will be collected for the survey will be dietary intake, anthropometric status, nutrition sensitive and specific indicators and micronutrient status.

The expected possible adverse effects: there is no adverse effect by participating in this study

The objective of this research: to produce information on anthropometric status, dietary intakes, and micronutrient status of different population groups in Ethiopia, and assess the coverage of direct and indirect nutrition interventions.

Study design: a population-based, cross-sectional design

The schedule of the study: The study will be conducted from July, 2021 to April, 2023

Foreseeable risks and expected benefits arising from participation in the study	
Foreseeable risks	Expected benefits
Risks to study participants for involvement in the	The findings of the study will help the ministry of
coverage survey are low. There may be risks	health and other stakeholder engaged in nutrition
associated with COVID pandemic. Interviewers	to improve and/or design appropriate health and
will be trained to minimize this risk and will use	nutrition intervention programs in the country.
appropriate prevention measures.	
Occurrences that may take place during the study period	
Occurrences	How to manage

Withdrawal of volunteers from the study	In such a case, we would respect the volunteer's
	decision to withdraw and also get a clear
	understanding of the reason for their withdrawal

At the end of the study, you will not be receiving any financial benefits, but you will get your **anemia** and **goiter** status for time you spent and participation.

All data collected from the study will be kept confidential. If you have any questions related to the study you may contact directly Dr. Masresha Tessema who is the project's PI

## The contact persons

1. Dr. Masresha Tessema

Tel. [+251 919782082] E-mail: [masresha88@gmail.com]

2. [Mr. Ibrahim Kedir] Tel. [+251 911957161] EPHI's IRB

Certificate of Consent		
I have read the foregoing information. I have an	I confirm that the participant was given an	
opportunity to ask questions and all my quest	opportunity to ask questions about the study and	
have been answered to my satisfaction. I	all questions have been answered correctly. I	
volunteer to give consent to participate in this	confirm that the consent has been given	
research study	voluntarily	
Printed name of the participant		
Signature of the participant's parent or guardian	Printed name of the person taking the consent	
Date	Signature of the person taking the consent Date	
day/month/year	·	
	day/month/year	

# 6. Assent form for Adolescent Girls (10-19 years)

## Ethiopian Food and Nutrition Strategy (FNS) Baseline Survey

**Investigator(s):** Dr.Masresha Tessema (PI), Meseret W/Yohannes, Dr. Meron Girma, Alemnesh Petros, Dr Aregash Samuel, Arnaud Laillou, Stanley Chitekwe, Kaleab Baye, Ramadhani Noor, Anne Sophie Donze and other co-authors

You are being invited to take part in this research because you are Adolescent girl. There are [16596] households taking part in this research. We will measure your dietary information (for those adolescent girls aged 15-17 years), information related to nutrition -sensitive and nutrition-specific practices, blood, and stool, we will also measure your height, weight, and mid upper arm and waist circumference and your goiter status

## Box 1. Taking part in this research is voluntary

- You can refuse to take part in this study.
- You **can withdraw** your participation from the study at any time

## Information related to the study

The FNS baseline survey will be conducted in the 12 regions of Ethiopia. The study population will be children age 0-59 months having caregivers/mothers, school-age children (6-12 years), adolescent girls (10-19 years), reproductive-age women (15-49 years), pregnant and lactating women, and household head. The indicators that will be collected for the survey will be dietary intake, anthropometric status, nutrition sensitive and specific indicators and micronutrient status

The expected possible adverse effects: There is no adverse effect by participating in this study

The objective of this research: to produce information on anthropometric status, dietary intakes, and micronutrient status of different population groups in Ethiopia, and assess the coverage of direct and indirect nutrition interventions.

Study design: a population-based, cross-sectional design  The schedule of the study: The study will be conducted from July, 2021 to April, 2023		
Foreseeable risks	Expected benefits	
Risks to study participants for involvement in the coverage survey are low. There may be risks associated with COVID pandemic. Interviewers will be trained to minimize this risk and will use appropriate prevention measures.	of health and other stakeholder engaged in nutrition to improve and/or design appropriate	

Occurrences that may take place during the study period	
Occurrences	How to manage
Withdrawal of volunteers from the study	In such a case, we would respect the volunteer's
	decision to withdraw and also get a clear
	understanding of the reason for their withdrawal

At the end of the study, you will not be receiving any financial benefits, but will get your results for height, weight, mid upper arm and waist circumference measurements, anemia and goiter status for time you spent and participation.

All data collected from the study will be kept confidential. If you have any questions related to the study you may contact directly Dr. Masresha Tessema who is the project's PI

# The contact persons

1. Dr. Masresha Tessema

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2. [Mr. Ibrahim Kedir] Tel. [+251 911957161] EPHI's IRB

Certificate of Assent		
I have read the foregoing information. I have an	I confirm that the participant was given an	
opportunity to ask questions and all my quest	opportunity to ask questions about the study and	
have been answered to my satisfaction. I	all questions have been answered correctly. I	
volunteer to give assent to participate in this	confirm that the assent has been given voluntarily	
research study		
Printed name of the participant	<del></del>	
	Printed name of the person taking the assent	
Signature of the participant		
Date	Signature of the person taking the assent Date	
day/month/year		
	day/month/year	