EFFECTS OF PRENATAL FAMINE EXPOSURE ON ADULTHOOD METABOLIC SYNDROME AMONG SURVIVORS OF 1975-77 E.C GREAT ETHIOPIAN FAMINE IN NORTH WOLLO ZONE, NORTHEAST ETHIOPIA: A HISTORICAL COHORT STUDY



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### Abstract

Background. The mortality impacts of 1975-77 E.C Great Ethiopian famine is well documented and was clearly significant. However, the long term sequel of the assault such as its impact on priming for adulthood chronic illness among adult has never been documented. Metabolic syndrome (MetS) has been hypothesized to have an association with grave intrauterine and early childhood nutritional problems. In response, the present study is designed to explore the effects of prenatal starvation on adulthood metabolic syndrome of the survivors of the great Ethiopian famine.

Objectives: To explore effects of prenatal famine exposure on adulthood metabolic syndrome among survivors of the 1975-77 E.C Great Ethiopian famine in North Wollo, Ethiopia, 2019.

Methods: A historical cohort study was conducted from March to April/2019 among 456 adult subjects selected using multi stage sampling methods. Data on socio-demographic and economic, behavioral, dietary consumption, anthropometry and biochemical measurements were collected through face to face interview using a structured questionnaire. Data were checked, cleaned and entered in to Epidata software version 3.1, and were exported to SPSS version 23 software for analysis. The *independent T* test and chi-square were used to assess the differences between two groups. A multivariable Logistic regression was used to control the possible confounders while estimating the effect of the exposure on metabolic syndrome. Odds ratio and their 95% confidence intervals were computed and a P-value of less than 0.05 was taken to declare the level of significance.

Result: Prevalence of MetS among adults who had history prenatal famine exposure was 18.5%, while the proportion among non-exposed adults was 8.4%. After adjusting with sex, residence, dietary consumption, physical activity, alcohol consumption, wealth tertiles. Prenatal famine exposed adults were twice more likely to have MetS compared with the non-exposed groups (AOR=2.25; 95% CI: 1.28, 4.21; P=0.002). There were also higher odds of MetS among adults having low physical activity as compared to high physical activity level (AOR=1.73; 95% CI: 1.07,4.21). Similarly, significant differences were observed in waist circumference, systolic and diastolic blood pressure measurements. But test for association for the high density lipoprotein had shown non-significant relationship (p=0.33)

Conclusion: Prenatal famine exposure is found to have significant association with increased risk of metabolic syndrome. Low physical activity of adults was further strengthening the association of famine exposure for metabolic syndrome. Nutrition during fetal life remains critical in modifying the risk for adulthood chronic disease.

Key words: famine, metabolic syndrome, fetal, Ethiopia

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# List of Acronyms / Abbreviations

ATP Adult Treatment Panel
AOR Adjusted Odds Ratio
BMI Body Mass Index
BP Blood Pressure

CHD Coronary Heart Disease
CI Confidence Interval
COR Crude Odds Ratio

CVD Cardiovascular Diseases

DBP Diastolic Blood Pressure

FOAD Fetal origin of Adult Diseases

FPG Fasting Plasma Glucose

HDL-C High density lipoprotein cholesterol

HgA1c Glycosylated hemoglobin

IDF International Diabetic Federation

LDL-C Low density lipoprotein

MetS Metabolic Syndrome

NCDs Non communicable diseases

SBP Systolic Blood Pressure

SRQ Self- Reported Questionnaire

T2DM Type 2 Diabetes Mellitus

TC Total cholesterol

TG-C Triglyceride cholesterol

WHO World Health Organization

### 1. Introduction

# 1.1 Background

Metabolic syndrome (MetS) is currently an emerging public health problem in low and middle income countries (1).

MetS resulting from the interaction of genetic, hormonal, and lifestyle factors. Over the past two decades, the number of people diagnosed with the syndrome has steadily increased in tandem with the global epidemics of obesity and diabetes. (2). MetS comprises a group of disorders, including abdominal obesity, hypertension, dyslipidemia and hyperglycemia(2).

According to International Diabetes Foundation, MetS is defined as the presence of central obesity plus any two of the following markers: high triglycerides ( $\geq 150$  mg/dl), low high density lipoprotein (HDL) cholesterol < 40 mg/dl in men and < 50 mg/dl in women, hypertension (blood pressure  $\geq 130/85$  mmHg or use of antihypertensive medication), high fasting blood glucose ( $\geq 100$  mg/dl or use of treatment for diabetes mellitus (3).

MetS increases risk for a 2.5-fold increased cardiovascular mortality and a 5-fold higher risk of developing diabetes as well as cost related to health care(4).

Different famine study showed that the risks of non-communicable disease begin in fetal life and continue into later age(5,6). Nutritional insult during critical period of fetal life may alter the structural and physiologic functional development of vital organs as well as lifelong effect on later body constitution moreover the tendency to become obese (7). These effects are termed 'programming' and represent an important risk factor for non-communicable disease(NCDs) in adulthood life. Adult chronic disease, therefore, supposed to reflect cumulative differential lifetime exposures to damaging physical and social environments(8).

Developing countries with populations that are chronically undernourished in early life are undergoing a nutrition transition and are experiencing an epidemic of metabolic disease. These dual burdens are thought to be causally related(9,10).

Ethiopia has a long and troubled history of famines including prolonged droughts and frequent severe rainfall failure, for which no body had studied its chronic effect thus far. Notably, a widespread famine affected the country from 1975-77 E.C. Its epicenter is Tigray and Wollo (11–14).

The Great famine of Ethiopia from 1975-77 E.C is a catastrophe in human history and caused 1.2 million people death, 800,000 internally displaced, 400,000 refugees left the country, and almost

200,000 children were orphaned, making it one of the worst famines in recent history(11–14) .Yet, can be considered an outstanding model. The mortality impacts are clearly significant but what of the survivors and the generation to come?

Because of ethical and practical concerns, common scientific approaches, such as exposing pregnant women and/or infants, adolescents to famine to investigate later effect in humans is impossible. Famine model studies can give direct evidence where early malnutrition plays a role in development of hypertension, insulin resistance, central obesity, dyslipidemia (15,16).

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# 1.2 Statement of the problem

The Great Ethiopian Famine was a historical disaster which occurred between 1975-1977 E.C (11–14). Such kind of famine studies are a natural experiment (17,18) that give a direct evidence for the hypotheses that early malnutrition plays a role in the origins of hypertension(19,20) ,central obesity (17),dyslipidemia(21,22) and impaired glucose tolerance(23–25) which are all components of MetS.

Non-Communicable Diseases (NCDs) are the leading cause of morbidity and mortality in many countries and has reached epidemic proportions globally. The 2017 WHO progress monitor showed that, NCDs kill 41 million people each year, over 85% of these premature deaths occur in low- and middle-income countries. In Ethiopia, from the total death occurred ,39% were from non-communicable diseases (26).

National survey study in Ethiopia revealed that the prevalence of hypertension, diabetes mellitus and overweight or obese were 15.6%, 6% and 6.3 % respectively(27). The prevalence of MetS in Addis Ababa (17.9%) (28) and Gondar (40.7%) (29).

The risks of adverse long-term consequences of famine exposure during critical period of growth and development is further exacerbated by increased industrialization, urbanization, sedentary life styles(30). A positive energy balance during adulthood and a nutritionally rich environment in later life (21,31,32). A good example of famine exposure is the group of Jewish migrants to Israel from Ethiopia, known as the Falashas. This group was moved from a country regularly blighted by famine to an essentially westernized nation. Within 5 years of the migration, rates of non-insulin dependent diabetes among the Falashas had risen to nearly 18%, which was 30 times greater than

the prevalence among Ethiopians living in Ethiopia and two fold greater than in the rest of the Israeli population (33).

In the view of the above, Ethiopia has double risk factors for the occurrence of non-communicable disease in which it has famine exposure history and recent increased urbanization. As of the growing burden of metabolic disorders, the comprehensive prevention and control strategies of preventing chronic diseases and their risk factors focusing on reduction of risky behaviors referred to as modifiable risk factors (physical inactivity, inadequate intake of fruits and vegetables, alcohol consumption and cigarette smoking(34) .Yet, those are the conventional approach to curb chronic diseases prevention.

An improvement in fetal, infant, and child growth which is a life course approach has the potential to reduce the incidence of metabolic disorders (10,35). Since the phenotypic effects of epigenetic modifications are long-term and potentially reversible, once the mechanistic basis of the disease is understood, intervention and strategies aimed at reversal can be devised and implemented (36). Hence this study is aimed to assess effects of prenatal famine exposure on adulthood metabolic syndrome among survivors of the Great Ethiopian famine in North Wollo, Northeast Ethiopia, March 2019

### 2. Literature Review

Fetal development is life stages that are characterized by rapid growth, development and maturation of organs and systems. Variation in the quality or quantity of nutrients consumed by mothers during pregnancy can exert permanent and powerful effects upon developing tissues. These effects are termed 'programming' and represent an important risk factor for non -communicable diseases of adulthood, including the metabolic syndrome (19).

The later development of disease triggered by the environment before and after birth is possibly explained through three kinds of process. First, they have fewer cells in key organs, such as lower nephron number in the kidney, low beta cells in the pancreases and lower cardio-myocytes in heart (37). Another process by which slow fetal growth may be linked to later disease is in the setting of hormones and metabolism.

A third link between starvation during critical period of growth and development and later disease is that people who were small at birth are more vulnerable to adverse environmental influences in later life (38).

Past and ongoing under nutrition among pregnant women may contribute to the development of metabolic syndrome as suggested by epidemiological studies from high income countries linking under nutrition in fetal life with increased burden of non-communicable diseases in later life(39). Exposure to suboptimum nutrition during crucial periods of development especially the fetal period increases the risk of MetS and non-communicable diseases in later life (40,41). The effect of maternal under nutrition during gestation on later health depends on its time during gestation. Particularly, early prenatal period is vulnerable (25).

Cross-sectional study conducted in china showed that by using ATP III criteria the prevalence of metabolic syndrome among adults in non-fetal (5.7%) and fetal exposed (7.7%) cohorts. The prevalence of the metabolic syndrome among fetal exposed with Western diet were (34.6%) and fetal famine-exposed cohort with a traditional diet in later life was only 4.2% (42).

Historical Cohort study done in Dutch showed that prenatal exposure to famine or reduced birth weight is not associated (OR: 1.2; 95% CI:0.9, 1.7) with a significantly greater prevalence of the metabolic syndrome.(38) .The mean prevalence MetS was 32% according to the widely applied national cholesterol education program definition and 49% according to IDF definition (38).

Historical cohort study done in china showed that the prevalence of MetS in the non-exposed and fetal exposed group men were 16.4% and 20.1% respectively and in women non-exposed (13.5%)

and fetal exposed (23.7%) by using IDF criteria. Famine exposure during the fetal period (OR 1.47;95% CI;1.05,2.07) had 47% higher risk of MetS as compared to non-exposed group (43). Another cross-sectional study done in china on retired workers showed that the prevalence of MetS by using IDF criteria were 25.2 %( non-exposed) and 26.9 % (fetal exposed group) and there is no significant association between fetal famine exposure and MetS (AOR=0.96, 95% CI: 0.77-1.20) (44).

Cross-sectional study conducted in urban residence of china population showed that prevalence of metabolic syndrome in Women with fetal exposed group (7.3%) and non-exposed group(4%). The odds of MetS is 1.89 times (AOR=1.87:95%CI: 1.15–3.04) higher in fetal exposed group as compared to non-exposed cohort (45). This study also showed that the prevalence of metabolic syndrome among men in control and fatally exposed groups were 20.1% and 22.5% respectively, but there was no significant difference of prevalence among the two groups (45).

Meta-analysis showed that fetal exposure group had 11% increase with risk of MetS Compared with the unexposed group.(46).

Cross-sectional study conducted on Holocaust famine survivors showed that prevalence of MetS among a non- exposed and exposed group were 9 % and 17% respectively. Fetal exposed group were 2 times (AOR = 2.14, 95% CI):1.48-3.47) more likely to have MetS as compared to non-exposed group (47).

Retrospective cohort study done in China in 2010 reported that SBP/DBP was about 0.7/ 0.7 mmHg higher in subjects exposed to famine during fetal development as compared to the non-exposed cohort (20).

Historical Cohort Study in china showed that fetal exposed group had a significantly higher SBP (2.2 mmHg) and DBP (0.9mmHg) difference as compared to non-exposed group (48).

Study conducted in Western Holland among Dutch famine survivors reported that famine exposure of at least 10 weeks duration was associated with elevated systolic (2.77 mmHg and diastolic (1.27 mmHg) blood pressure as compared to non-exposed (49).

Historical Cohort Study conducted in Nigeria showed that Fetal-infant exposure to famine was associated with elevated systolic (+7 mmHg) and diastolic (+5 mmHg) blood pressure.(39).

Cross-sectional study conducted in Israel (Holocaust famine) showed that diabetes among exposed group were 15. % greater than non-exposed group (50).

Cross-Sectional Survey conducted in Northeastern China showed that increased risk of diabetes in the fetal exposure cohort was 2.7% higher than non-exposed group (51).

Another study conducted in china in 2010 showed that prevalence of hyperglycemia among adults in non-exposed and fetal exposed were 2.4%, 5.7% respectively (23).

Individuals who had been exposed to famine during the fetal period and who had lived in a severely affected area had 0.31% higher glycosylated hemoglobin (HbA1c) as compared with unexposed individuals (52).

Other cohort study conducted in china in 2016 reported that the prevalence of diabetes in the exposed group was 12.8% higher as compared with unexposed individuals (53).

# 2.1 conceptual frame work

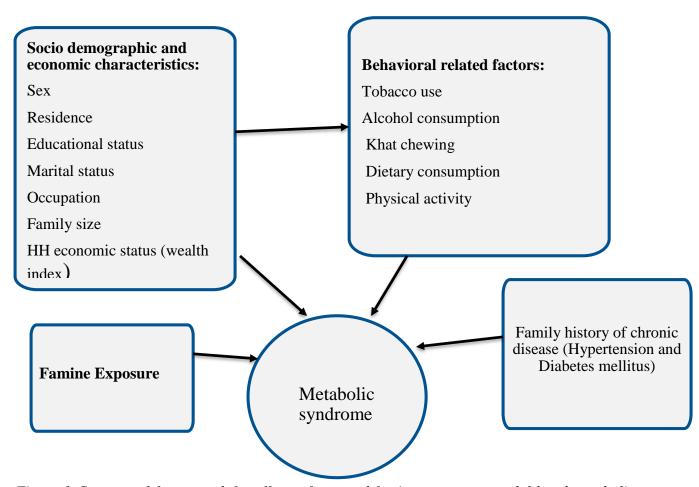


Figure 1: Conceptual frame work for effects of prenatal famine exposure on adulthood metabolic syndrome, based on Agent-Based Model (Jeff, 2010)

# 2.2. Significance of the study

Because of ethical and practical reasons, such as exposing pregnant women and/or infants, adolescents to famine to investigate later effect in humans is impossible. Now it is a previous chance for us to study the long-term effects of malnutrition in the womb. As a result, the study will provide a clue of what is behind on alarmingly increasing non-communicable chronic diseases in our country, Ethiopia. Understanding the role of early life nutrition is essential for the implementation of initiatives aimed at curbing the current metabolic disorders which is particularly relevant to Ethiopian.

The study will enhance our knowledge of fetal origin of adult's disease. The study is also help to establish the first African cohort of starvation study.

Besides providing evidence on the impact of famine on chronic non-communicable disease, the study will contribute to the broader empirical literature on the importance of maternal and early childhood nutrition. The potential of the finding is great with respect to several implications for policy, health service and self-care practice, preventing the disease that occur later in life and transgenerational effect of famine. The study will provide peculiar evidence to formulate setting specific prevention strategies for non-communicable diseases.

# 3. Objectives

# 3.1 General objective

To assess the effects of prenatal famine exposure on metabolic syndrome among survivor's adulthood of the Great Ethiopian famine in North Wollo, Northeast Ethiopia, March 2019

# 3.2 specific objectives

- ✓ To examine the effects of prenatal famine exposure on risk of metabolic syndrome among survivor's adulthood.
- ✓ To identify factors associated with the risk of metabolic syndrome among survivor's adulthood.

### 4. Methods and Materials

# 4.1 Study Area and Period

The study was conducted in North Wollo Zone, Ray kobo town and Rural Woreda, located in the northeastern part of Amhara regional state from March 15 to April 30, 2019. The two Woreda are located at 49.7km form Woldya capital city of north wollo zone and 408 km from Bahirdar the capital city of Amhara regional state.

The land in the area is mostly mountainous and not really suitable for cultivation. There are five agro-ecological zones namely, Kolla (low land and warm), Woinga-dega (moderate), Dega (cool), Kur Wurch (cold) and Berha desert with high temperatures. The two Woreda has 38 kebeles of which 4 were urban and the other 34 were rural. Currently about 289, 877populations are living in the Woreda, out of which 147,837 are females (North Wollo population, EFY,2011) This area had troubled history of famines occurred between 1975-1977 E.C where its epicenter is Tigray and Wollo. ''Waja'' a small town between kobo and Alamata was place where famine first began. The 1975-1977 E.C crisis is referred to by some as the "time men ate grass": many peasants had eaten wild plants to save themselves from a slow, agonizing death. It was the worst disaster in living memory, in human history and caused over half a million deaths, making it one of the worst famines in recent history, yet can be considered an outstanding model (11,14).

Severe rainfall failure, military offense, aerial bombardment of markets, destruction of cattle and grain stores, burning of crops, and tight controls on movements of migrants combined to prevent the normal redistribution of grain and livestock surpluses in northern Ethiopia (13,14).

# 4.2 Study Design

A historical cohort study design was employed.

#### 4.2.1 Famine cohorts and exposure age categories

Subjects were categorized into two exposure cohorts according to the subject's birthday and corresponding exposure period (54). The prenatal famine exposure periods were taken, using 1 April ,1975. E.C., the start of the famine, as reference to assess age at famine exposure, Subjects who were born between April 1,1975 to August 30, 1977.E.C. categorized as exposed group whereas those born between 1 September 1979 and 30 September 1981.E.C. categorized as non-exposed group based famine condition lasted in the area(11). However, to reduce effect of famine on subsequent group of control, a period of one-year transition time ( washout time of famine) (39)

considered and hence, adults born between 1 September 1978 to 30 August 1978 were excluded (55). Accordingly, exposed group were adults of current age range from 34 to 36 and non-exposed group was those age range from 30 to 32.

Table 1::Window of exposure to the 1975-77 E,C Great Ethiopian Famine cohort, North Wollo zone, 2019.

Date of birth	Exposure to famine (September	Age in 2011
(dd/mm/yyyy)	1975-August 1981)	(years)
01/08/1975-30/12/1977	Prenatally exposed individuals	34-36
01/01/1978-30/12/1978	Transition (Washout period)	33
01/01/1979-30/01/1981	No exposure (reference group)	30-32

# 4.3 Population

# 4.3.1 Source population

All adults born during the 1975-77. E.C. Great Ethiopian famine and adult born after the famine (1979-1981E.C).

# 4.3.2 Study population

All randomly selected adult born during the 1975-77. E.C. Great Ethiopian famine and adult born after the famine September 1979-1981 E.C.

# 4.4 Sample Size Determination and Sampling Technique

# 4.4.1 Sample size determination

The sample size was calculated by using Epi-Info version 7 by considering 5% margin of error 80% power, design effect 1.5, 5% non-response rate and 1:1 ratio (r=1). Assuming a prevalence of type 2 diabetes mellitus in fetal exposed to be (22.6%) and non-exposed group (9.8% (53) Hence the final sample size was 456.

# 4.4.2 Sampling procedures

### 4.4.2.1 Selection of famine exposed and non-exposed groups

Multistage stage sampling technique was used to select the study subjects across North wollo Zone. First, Raya kobo Woreda and Raya kobo town was purposely selected then 30% of Raya kobo Woreda (10 kebeles) and Raya kobo town was selected.

Second, baseline survey was conducted on selected kebeles to identify the cohort of adult who were exposed for prenatal famine during the great Ethiopia famine period with the guidance of developmental army and health extension workers. Registration were done for all adults born during the famine season and after famine season and assigned a unique number to each of them in order to prepare sampling frame. Then number of adults were allocated for selected kebeles based on population proportion to size.

Adults were selected from each kebeles by using simple random sampling method from registration. In households with more than one adult was found by using lottery method one of the adult were taken.

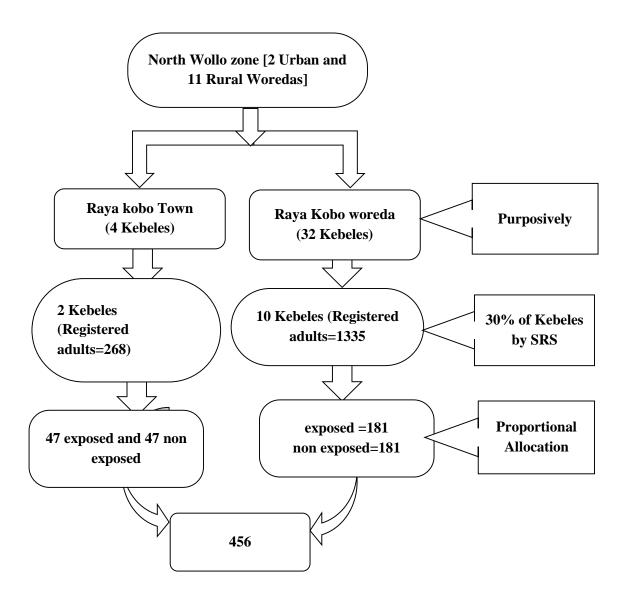


Figure 2:Sampling procedure in the section of eligible subjects for 1975-77.E.C Great Ethiopian Famine, North Wollo zone, Ethiopia

### 4.5 Study Variables

### 4.5.1 Dependent variable

Metabolic syndrome

### 4.5.2 Independent variable

- ✓ Prenatal exposure to famine
- ✓ Socio demographic characteristics: Sex, residence, level of education, marital status, occupation, family size and household economic status (wealth index).
- ✓ Behavioral related factors: Tobacco use, alcohol consumption, Khat chewing, physical activity, history of dietary consumption
- ✓ History of chronic illness (Hypertension, Diabetes mellitus)

# 4.6 Eligibility Criteria

#### 4.6.1 Inclusion criteria

Adults with current age 34-36 who were born during the 1975-77. E.C. Great Ethiopian famine and adults with current age 30-32 who were born after the famine (1979-1981 E.C).

#### 4.6.2 Exclusion criteria

Adult who have physical disability including deformity (Kyphosis, Scoliosis, limb deformity), pregnant women or lactating mother of less than 3 months and using hormonal contraceptives and seriously ill. Those who displaced to other area of the country and those were in other location during the famine adults were excluded

### 4.7 Data Collection Tools and Techniques

Data of socio-demographic characteristics, history of famine condition, history of chronic illness and behavioral characteristics of the participants such as dietary history, smoking status, alcohol consumption, khat chewing, physical inactivity were collected using structured questionnaire. Household wealth status was assessed by asking about household assets, utilities and housing characteristics were used finally to generate household wealth index based on EDHS (56).

### Assessing famine condition

The famine related questions was asked to obtain some information on exposure status to the famine and determine extent of mortality and morbidity consequences using recommended WHO verbal autopsy tool supplementing specific area and period of exposure to the famine.(57)

### Dietary assessment

The dietary consumption over the past 12 months was assessed based on recommended seven food group by FAO, using 38 food items that highly consumed in the area. Participants were asked about the usual frequency of each food consumed during the past 12 months. According to the frequency of food intake, each food item was classified into nine intervals. Before actual data collection, it was validated in the study area (Cronbach alpha,0.8) conducting pretest (58).

Physical activity: Physical activity was assessed based on intensity, duration and frequency of physical activity at work, in recreational settings and involving transportation (journeys), using a set of 16 questions. Data were collected on the number of days, hours and minutes of physical activity performed at work, involving transportation and in recreational settings for at least 10 minutes or more continuously each day. The three levels of physical activity suggested in the WHO recommendations for classifying populations were used as low, moderate, and high level physical activity (27).

#### **History of Hypertension and Diabetes mellitus**

Individuals and family history of diabetes and raised blood pressure were assessed by asking whether specific measurements for these purposes previously performed and any confirmed of the case by a doctor or health worker. Study participants were also asked about if any medication was taken for aforementioned purposes.

### Anthropometric measurement

The anthropometric data were collected by using the procedure stipulated by the WHO for taking anthropometric measurement. All anthropometric measurements were done in two times and the average value was used for further analyses (59).

Body weight: Weight was measured using portable battery operated Seca digital scale (Seca Germany). The weighing scale was checked for zero reading before the adult was asked to stand on it. In addition, the proper performance of each scale was checked every day by known 3kg sand filled plastic bottle before the fieldwork. A standardization exercise was performed during the training to capture technical error of measurement (TEM). During the procedure the subjects has been worn light clothes and taken off their shoes. The weight was recorded to the nearest 0.1 kg.

Height: Height was measured using portable stadiometer (Seca Germany). All participants have been measured against the wall in upright position, without foot wear and with heels together and their heads positioned and eyes looking straight ahead (Frankfurt plane) so that the line of sight

perpendicular to the body. knees straight, the heels, buttocks and the shoulders blades touching the vertical surface of the stadiometer. The height was recorded to the nearest 0.1cm. The same measurer had employed for a given anthropometric measurement to avoid variability.

Waist circumference: Waist circumferences was measured using figure finder tape measure (SECA, Germany, Model 200) recommended by the WHO. The measurement was taken in the mid-axillary line midway between the last rib (10th rib/lower margin of lowest palpable rib) and the superior iliac crest with the client wearing no or light clothing at the end of a normal expiration/at the end of exhalation. The measurement was taken to the nearest 0.1 cm over the skin.

#### **Blood pressure measurements**

Blood pressure (BP) was measured in the left arm with sphygmomanometer. Three seated BP measurements were taken for each subject spaced five minutes apart. To improve the reliability of measurement three readings was taken with 5 min interval and the average of the three readings was recorded as the final BP of the patient. But if the difference between the two readings was greater than 5 mmHg, a third measurement was taken and the average of the three readings was recorded as the final BP of the patient.

#### **Biochemical measurements**

Biochemical measures include fasting total cholesterol; plasma glucose; triglycerides, and high-density lipoproteins were measured. The collection of venous blood was carried out the day after collection of behavioral and physical measurements. Blood was taken from the survey participant after overnight fasting (8-12 h) in the morning (07:00–09:00). Five milliliter of venous blood was collected from each patient in plane test tubes and serum was separated immediately. The serum was separated and put into ice bag and safely transported to Dessie regional laboratory. The extracted serum was investigated for High density lipoprotein-cholesterol(HDL-c), triglyceride and fasting glucose using A25 bio-system clinical chemistry analyzer.

#### Data collectors

In the data collection process, 12 developmental army and 12 health extension workers were deployed over 12 Kebeles to conduct base line cohort survey. Four team composed of eight BSc nurses, four laboratory technologist and four Public Health officers who are fluent in the local language were participated as a data collector and supervisor respectively. The interview and

physical measurements was taken at household level and repeat at nearby facility appointing them for subsequent day of visit since those measurement was expected taken at their fasting state.

## 4.8 Data Quality Control

The questionnaire was prepared in English and then translated to local language (Amharic). Validation and calibration of the instrument after each measurement and after moving the instrument from one place to another was performed. Data collectors were recruited based on their qualification and prior experience of data collection. An intensive 4 days training were provided for the selected data collectors and field supervisors. A pretest was done on 5% of the sample in Woldya Zuria a one week prior to the actual data collection.

Standardize operated procedures were follow to collect blood samples and to perform laboratory analysis.

# 4.9 Data Processing and Analysis

The data were cleaned, coded and entered into Epidata version 3.1 and analyzed using SPSS version 23. Data were summarized and presented using descriptive statistics. The comparison of baseline characteristics between famine exposed and non-exposed participants were carried out using the independent t-test for continuous variables with normal distribution and the chi-square test for discrete variables.

Bivariate analysis was used to check each independent variables having association with the dependent variable then those variables found to have p-value of less than 0.25 was entered in to multivariable logistic regression for controlling the possible confounders. Odds ratio with their 95% CI was computed and variables having p - value less than 0.05 in the multivariate logistic regression model was considered as significantly associated with the dependent variable.

Hosmer and Lemeshow test used for assessing of the goodness-of fit of the model with chi-square of 5.9 and p value of 0.55. All tests are two sided, and p values less than 0.05 were considered statistically significant.

#### 4.10 Ethical Consideration

Ethical Approval was obtained from Jimma University Institute of Health, Ethical Review Committee. Permission letters was secured from Amhara regional health bureau and North Wollo Zone Health Department as well as from the two Woreda health offices. The purpose and importance of the study was explained to the participants. Written consent was sought from participants following and explanation of the research study, clarification that participants can be

free to withdraw from the study at any time. The names and address of the participants was not recorded in the questionnaire to assure confidentiality.

After identifying of adult with metabolic syndrome during taking of the laboratory result. All adults with risk of metabolic disease were get counseling and linked to the health facility as well advice given for strengthen further screening habit for chronic illness.

#### 4.12 Dissemination Plan

The results of the study will be presented and submitted to Department of Population and Family Health, Institute of Health of Jimma University, Amhara Regional State Health Office, North Wollo Zone Health Administration and Woreda Health Offices.

Study results will be disseminated to the scientific public by presentation at national and international conferences and by publication in peer-reviewed journals. we aim to also disseminate the study findings among nutritionists, midwives, pediatricians and gynecologists who are in a position to advise pregnant women, lactating mothers and children about nutrition and the public in general by publishing results in national newspapers.

# 4.13 Operational Definitions

Metabolic syndrome: According to international Diabetes Federation (IDF) metabolic syndrome can be defined as the presence of abdominal obesity (waist circumference (male  $\geq 83.7$  cm, female  $\geq 78$  cm) plus any of the four parameters (3)

- ➤ Raised TGs >1.7 mmol/L (>150mg/dl) or treatment for this dyslipidemia (3)
- ➤ Reduced HDL-C <1.03 mmol/L (40mg/dl) in men or <1.29 mmol/L (<50mg/dl) in women, or treatment for dyslipidemia (3)
- Raised BP: systolic BP  $\geq$ 130 or diastolic BP $\geq$ 85mmHg, or treatment of hypertension
- Raised FPG  $\geq$ 5.6 mmol/L ( $\geq$ 100 mg/dL) or a history of type 2 diabetes (3)

Body max index (BMI): Body mass index a person's weight in kilograms (kg) divided by his or her height in meters squared.

Waist circumference (WC): Waist circumference is body composition measurement and an important predictor of health outcomes in adult men and women of all age groups cut-off for obesity was 83.7 cm(males) and 78.0 cm (females (60).

Famine exposure: famine exposure was defined as prenatal exposure to famine period of 1975-77. E.C. (time exposure) and born in the area of widespread scarcity of food, caused by several factors

including war, inflation, crop failure, or government policies. This phenomenon is accompanied verbal autopsy regarding food scarcity and increased mortality in the area (61).

Wealth index: is a composite measure of a household's cumulative living standard. Households were given scores based on the number and kinds goods they own ranking each person in to high, medium and low socio economic status after performing principal component analysis.

Assessing tobacco use: all current smoker and past smoker of more than one cigarette per day while nonsmoker if they found never smoker. (27).

A standard drink is any drink containing about 10g of alcohol. Standard drinks per drinking occasion. A standard drink contains approximately 10g of pure alcohol. One bottle of factory beer or one tin of local alcoholic drink like tella and tejj considering common local measurement were considered as standard drink. standard alcoholic drinks during one occasion was asked among current (past 30 days) drinkers (27).

Consumption of  $\geq$ 60gm of pure alcohol for men and  $\geq$ 40gm of pure alcohol for women on an average day in the past 30 days were considered high level use of alcohol (62).

Khat use: all current khat chewer and past chewer of more than one bundle of khat per day/ week while non chewer if they found never chewer (27).

Unhealthy dietary consumption: consumed low fruit and vegetables and high salt and animal source food.

High-level physical activity involves a person reaching any of the following criteria: moderate-intensity activity at least three days per week, achieving at least 1500 MET minutes per week Moderate level physical activity involves a person not meeting the criteria for the high-level category, but meeting, achieved 600 to 1500 MET-minutes per week.

Low level physical activity involves a person not meeting any of the above-mentioned criteria less than 600 MET-minutes per week (27).

### 5. Result

# 5.1 Sociodemographic-economic characteristics of adults

Out of the total 456 adults, initially planned for the study, 447 of them were included in this analysis with response rate of 98%. The mean (SD) age of adults was 33.19(2.09). The study consisted 258 (57.7%) of females while the rest were males. Majority (92.6%) of respondents were orthodox by religion and the remaining of the respondents were Muslim followers. Higher proportion (82.1%) were living in rural area while 18.8% of respondents were living in urban areas. Two third (64.9%) of the respondents were married followed by single marital status. One third (30.6%) of the respondents cannot read and write. More than half (51.7%) respondents had medium wealth tertile while 31.5% were from low socio economic status (Table2). According to table 2 below except marital status, educational status and family size all socio demographic characteristics were not different among the famine exposed and non-exposed groups.

Table 2:Socio demographic characteristics of the study participants by famine exposure Status, in North Wollo zone, Northeast Ethiopia, March to April 2019 (n = 447)

Variables	Total N (%)	Famine Non- exposed	famine Exposed group	Pa
Sex Female Male	258 (57.7) 189 (42.3)	123 (47.7) 102 (54)	135 (52.3) 87 (46)	0.189
Residence Urban Rural	84 (18.8) 363 (81.2)	42 (50 183 (50.4)	42 (50) 180 (49.6)	0.946
Religion Orthodox Muslim	414 (92.6) 33 (7.4)	211 (51) 14 (42.4)	203 (49) 19 (57.6)	0.345
Educational status cannot read and write primary school secondary school above secondary school	137 (30.6) 97 (21.7) 113 (25.3) 100 (22.4)	53 (38.7) 42 (43.3) 67 (59.3) 63 (63)	84 (61.3) 55 (56.7) 46 (40.7) 37 (37)	0.02
Marital status Single Married divorced/widowed	92 (20.6) 290 (64.9) 65 (14.5)	59 (64.1) 145 (50) 21 (32.3)	33 (35.9) 145 (50) 44 (67.7)	0.0004
Occupational status				0.06
Government employee	99 (22.1)	55(55.6)	44(44.4)	
Non- government employee	38 (8.5)	23(60.5)	15(39.5)	
Farmer	88 (19.7)	36(40.9)	52(59.1)	
House wife	122 (27.3)	56(45.9)	66(54.1)	
Merchant	66 (14.8)	32(48.5)	34(51.5)	
Others <sup>1</sup>	34 (7.6)	23(67.6)	11(32.4)	
Family size =<4 =>5	322 (72) 125( 28)	174 (54) 51(40.8)	148(46) 74(59.2)	0.012
Wealth index Low Medium High	141 ()31.5 231(51.7) 75 (16.8)	67(47.5) 121(52.4) 37(49.3)	74(52.5) 110(47.6) 38(50.7)	0.649

1= students, daily laborer, Drivers

# 5.3 Behavioral related characteristics of adults

More than half (54.1%) of the respondents were had high physical activity while small portion (10.1%) were had low physical activity practice. One hundred ninety-one (42.7%) were consumed unhealthy diet and the rest were consumed healthy diet. Two third (63.5%) were drink alcohol greater than one standard per one occasion. larger proportion of study participant do not chew khat and use cigarrete. There is no difference in behavioral related factors between groups.

Table 3:Behavioral characteristics of the study participants by famine exposure Status, in North Wollo zone, Northeast Ethiopia, March to April 2019 (n = 447)

Variables	Total N (%)	Famine Non- exposed	famine Exposed group	P a
Physical activity level				0.472
Low	45 (10.1)	20 (44.4)	25 (55.6)	
Moderate	160 (35.8)	86 (53.8)	74 (46.3)	
High	242 (54.1)	119 (49.2)	123 (50.8)	
Dietary consumption				
Healthy	256 (57.3)	120(46.9)	136 (53.1)	
Unhealthy	191 (42.7)	105(55)	86 (45)	
Alcohol consumption				0.851
Low level drinker	163 (36.5)	83 (50.9)	80 (49.1)	
High level drinker	284 (63.5)	142 (50)	142 (50)	
Khat chewing				0.864
Yes	25 (5.6)	13 (52)	12 (48)	
No	422 (94.4)	212 (50.2)	210 (49.8)	
Smoking				0.65
Yes	6 (1.3)	3(50)	3 (50)	
No	441 (98.7)	222(50.3)	219 (49.7)	

P a p- value for chi-square and fisher exact test

### 5.2 Famine Intensity Assessment of 1975-77 E.C Ethiopian Great Famine

The level of great famine condition that had been 30 years back in area were assessed through verbal autopsy and based on available demographic data. Participants were asked and responded to the issues related to feelings toward state of famine by remembering parents telling about issue, availability of food items to cope with famine and presence of death in households to know severity

of famine in complement with area exposure and time exposure. The result shows 441(98.1%) felt hardly while the rest were felt little worry of famine as explained from by their family, none of participants had sufficient foods or resource during famine period. Nineteen (4.3%) of respondents were reported deaths from family member due to extreme famine

### 5.4 Differences among metabolic parameters based on famine exposure

There existed significance difference in waist circumference P=0.025), triglyceride(p=0.01), systolic and diastolic blood pressure (p=0.052, p=0.003), fasting plasma glucose(p=0.001) and metabolic syndrome (p=0.002) between famine exposed and non-exposed groups. However, there is no significance difference in high density lipoprotein (HDL)(p=0.336) among famine exposed and non-exposed adults. (Table 4)

Table 4:Association of famine exposure status with metabolic components of adults in North Wollo zone, Northeast Ethiopia, March to April 2019 (n = 447)

Metabolic components	Non-e	exposed	Exposed		P value	Mean difference 95% CI
Waist circumference, cm	81.33	± 11.1	83.66	5 ±10.42	0.025	2.27 (0.28, 4.26)
HDL, mg/dl	45.3 ±	$.3 \pm 11.8$ 44.2		±12.27	0.336	-1.09 (-3.34, 1.14)
Triglycerides, mg/dl	76.69	(60.99)	87.65 (78.24)		0.010	10.96**
Systolic BP, mmHg	112.2	± 11.9	114.6	0±12.91	0.052	2.23 (-0.02,4.46)
Diastolic BP, mmHg	74.5 ±	9.4	76.9±9		0.003	2.47 (0.84, 4.11)
Raised FPG n (%) (yes)	26 (1)	1.6)	52 (23.4)		0.001	11.8*
MetS n (%) (yes)	19	(8.4)	41	(18.5)	0.002	10.1*

Data presented are mean ± SD for continuous variables, and n (%) for frequency variables.

#### **5.4 Factors associated with metabolic syndrome**

Bivariate and multivariable logistic regression analysis was done using enter method to analyze factors associated with metabolic syndrome On the Bivariate analysis variable like famine exposure, sex, residence, wealth tertiles, physical activity, dietary consumption, alcohol consumption became a candidate for multivariate analysis.

P values in independent T test for differences in means or Chi-square tests for differences in proportions between non-exposed and exposed group \*\* median (interquartile range) the data were not normally distributed and p-value from logarithmic transformation \* proportion

Adults having prenatal famine exposure were 2.25 times more likely to have metabolic syndrome as compared to non-exposed group (AOR=2.25; 95% CI: 1.28, 4.21). The odds of metabolic syndrome were nearly two times among adults with low physical activity level as compared to high level physical activity (AOR=1.73; 95% CI: 1.07,4.21).

Table 5: :Multi variable logistic regression analysis of metabolic syndrome among adults in NorthWollozone,NortheastEthiopia,2019(n=447)

Variables	Metabolic syndrome		COR(95%CI)	AOR(95%CI)	
	Yes	No			
	n (%)	n (%)			
Famine exposure status					
Exposed	41(18.5)	181(81.5)	2.46 (1.37, 4.38)	2.25(1.28, 4.21) *	
Non-exposed	19(8.4)	206(91.6)	1	1	
Sex					
Female	34(13.2)	224(86.8)	1.05 (0.88, 1.87)	1.02 (0.55,1.65)	
Male	26 (13.8)	163(86.2)	1	1	
Residence					
Urban	13(15.5)	71(84.5)	1.23 (0.63, 2.39)	1.22 (0.61,2.23)	
Rural	47(12.9)	316(87.1)	1	1	
Wealth tertiles					
Low	19(13.5)	122(86.5)	1	1	
Medium	30(13)	201(87)	0.96 (0.52, 1.77)	0.85 (0.45,1.67)	
High	11(14.7)	64(85.3)	1.11 (0.49, 2.46)	1.09 (0.48,2.49)	
Physical activity level					
Low	8(17.8)	37(82.2)	1.79 (1.16, 4.26)	1.73 (1.07,4.21)*	
Moderate	26(16.3)	134(83.8)	1.61 (0.89,2.89)	1.56 (0.84,2.77)	
High	26(10.7)	216(89.3)	1.01 (0.07,2.07)	1.50 (0.04,2.77)	
Dietary consumption	20(10.7)	210(0).3)	1	1	
_ = ===================================					
Unhealthy diet	26(13.6)	165(86.4)	1.03 (0.59,1.78)	1.01 (0.54,1.71)	
Healthy diet	34(13.3)	222(86.7)	1	1	
	34(13.3)	222(00.7)			
Alcohol consumption					
Low level drinker	21(12.9)	142(87.1)	1	1	
High level drinker	39(13.7)	245(86.3)	1.07 (0.61,1.90)	1.03 (0.57,1.87)	

<sup>\*=</sup> variables having statistically significant association (p-value <0.05) and 1=reference group.

### 6: Discussion

Our results demonstrated significant difference in waist circumference, MetS, systolic and diastolic blood pressure and diabetes mellitus between prenatal famine exposed and non-exposed group in general in the study setting. The overall prevalence of metabolic syndrome was 13.4% which is lower compared to Dutch study (32%) (38). This difference could be existing socioeconomic disparity, criteria used to diagnose metabolic syndrome; the Dutch study used the National Cholesterol Education Program definition whereas the current study use IDF. Age and racial difference of the study participants might be another possible justification.

The prevalence of metabolic syndrome among adults in prenatal famine exposed group was 18.5% and non-exposed group was 8.4%. our finding is comparable with study done on holocaust survivors in which MetS were 17% in fetal exposed adult and 9% of non –exposed adults. But this finding is higher compared to the study done in china in which MetS among adults in prenatal famine exposed group (7.7%) and non-exposed (5.7%) groups (42). This inconsistency could be the difference in methodological factors where the present study had smaller sample, use one to one ratio (exposed non-exposed ratio) comparison for the outcome, racial difference, age difference of study participants and time difference where chines study done 8 years back (2011) from current study (2019). Another reason could be difference in using diagnostic criteria for MetS, china study use ATPIII in which it needs higher cut off for central obesity and fasting plasma glucose for MetS definition whereas present study use IDF for the diagnosis of MetS.

Contrarily the present prevalence of MetS was lower compared to two other china study where prevalence of MetS in the non-exposed (16.4%) and fetal exposed was 20.1%. (43) and report from another china study in which MetS in non-exposed (25.2%) and fetal exposed (26.9%) group (44). This discrepancy could be explained by methodological difference where the china study had larger sample, studied participants at different age and racial difference and chines study had relatively had longer duration (about 3 years). Developmental difference in which china were developed country in which the nutritional transition, use of technology and others favors in high metabolic syndrome as compared to our country.

The present study reported significant association between prenatal famine exposed adults and MetS. Adult who had prenatal famine exposure history were 2.25 times more likely to have a risk of metabolic syndrome as compared to non-exposed group. This report is supported with other similar studies conducted earlier (43,45,47). This finding explained by barker hypothesis in which

adults of the MetS may originate from adverse conditions during gestation and may depend on the timing and nature of the insult in utero. Organs and tissues are more vulnerable during periods of rapid growth and development, the so-called critical periods. Thus, exposure to famine during a specific period of gestation may lead to problems associated with the organs or physiologic systems that are undergoing development at that particular phase of gestation (37,38).

Prenatal famine exposure impact on adult chronic disease were explained in the different ways, by lower number of nephrons in kidney, lower number of  $\beta$  pancreatic cells in pancreas, lower number of cardio myocyte in heart and optimizing growth of the central nervous system.

Second, Fetal under nourished baby may also establish "thrifty" way of handling food. For example, Insulin resistance may be viewed as persistence of a fetal response by which blood glucose concentrations were maintained for the benefit of the brain, but at the expense of glucose transport into the muscles and muscle growth and others organs.(5).

Prior study on the Ethiopian migrants to Israel were strengthen our finding in which the Felisha who prior living in repeatedly famine attacked area and get adult hood post-natal mismatch result in high prevalence of diabetes mellitus (33,63).

unlike to the china famine study this study find significant association between the adult prenatal famine exposure and risk of MetS (44). This might be partly due to the different duration of famine Compared to the china famine, the great Ethiopian famine had relatively small duration (about 2 years) however china famine lasted 3 years. Racial difference and use of different diagnostic criteria for MetS diagnosis might be another potential reason to explain the different findings. Other possible reason could be the china study participants were retired workers and they are already survived the hard ship, whereas the present study participants were young age.

Our finding also disagree with the Dutch famine study, current study revealed significant association between adult prenatal famine exposure and risk of MetS (38). This might be partly due to the different duration of famine period, the great Ethiopian famine had a longer duration (about 2 years) however Dutch famine lasted six months which does not cover full gestational period, socio economic difference in which Dutch is developed country where early catch up would happened, use of different criteria to identify famine exposed group which Dutch study use daily ration and birth registration data while current study use birth date. Racial difference and use of different diagnostic criteria for MetS might be another potential reason to explain the different findings.

The current finding revealed low physical activity were associated with MetS. Similar finding support this result from EPHI, 2016 and National NCDs STEPS Survey, 2015. This could be explained in such a way that the life style practice of Ethiopian was changed because of the recent urbanization and modernization which favors for sedentary way of life and later risk for adult chronic disease. On other hand adult with prenatal exposure were further increase the risk of MetS in which explained by the synergetic effect of low physical exercise with previous famine exposure history

Current study revealed that there is significant difference in systolic (2.23 mmHg and diastolic (2.47 mmHg) blood pressure between fetal exposed and non-exposed group. The results were in line with the findings of the Dutch (systolic 2.77 mmHg and diastolic 1.27mmHg) (49) and china (systolic 2.2 mmHg and diastolic 0.9mmHg) blood pressure(48). Our finding is lower than study conducted in Nigeria in which in systolic (+7 mmHg) and diastolic (+5 mmHg) blood pressure in fetal exposed group as compared to counterpart (39). This discrepancy could be the Nigerian study SBP/DBP difference were from pooled fetal –infant exposure as compared with our finding in which the difference come from only prenatal famine exposed group. However, our finding is higher as compared to earlier china study (SBP/DBP was about 0.7/ 0.7 mmHg higher in subjects exposed to famine.(64) .However, siege of Leningrad study did not support our finding.(65). This could be Leningrad study had small sample size and had racial difference.

Diabetes mellitus related prevalence among adult who had prenatal famine exposed was 23.4% and non-exposed group was 11.6% in the current study. This finding is in line with study done in china in which 9.8% (non-exposed) and 22.6% were exposed group (53) and but lower to Holocaust famine study in which exposed (32.9%) and non-exposed group (17.4%) (50) however, it was higher than two china study (23,51). This difference could be explained by geographical and other differences between the population involved in our study and the previous study. Another possible explanation were use of different in measurement of bio chemical test.

# 7. Strength and limitation of the study

# 7.1. Strength of the study

Subjects born in the transitional period were excluded; there was no late gestational overlap with famine in the unexposed group.

Laboratory analyses were done in star regional laboratory branch outside the study area. Include all nutritional assessment method.

# 7.2 Limitation of the study

Lack of age matched true non-exposed group in the same area to see the real effect of famine on adulthood metabolic status.

Lack of birth weight data which is the most commonly used index for fetal undernutrition.

Age estimation were from self-reported of respondent's miss classification of exposed and non-exposed group would be occurred.

The data were self-report from the participants; thus subject to recall bias. However, due attention was given to the entire procedure of data collection; its effect might not be a threat to the findings of the study

# 8. Conclusions

Prenatal famine exposure is associated with increased risk of metabolic syndrome during adulthood. There is also a significance difference in waist circumference, systolic and diastolic blood pressure, fasting plasma glucose level among exposed adult compared to non-exposed adults.

Moreover, Low physical activity level was identified as one of the predictors for MetS among studied adults.

### 9. Recommendations

Our study indicates that early life environment is critical in development of the metabolic syndrome during adulthood. Recommendations we derived from the present study are:

For Ministry of health /regional health bureau, zonal health office

- > The ministry of health should give special due consideration for historical famine exposed area regarding the risk of chronic illness.
- Exceptional emphasis should be given for future primary prevention programs on malnutrition to includes interventions focused on pregnant women nutrition
- > Strengthening the existing programs on essential nutrition action
- > Strengthening current chronic illness programs and strategies with special focus on historically famine exposed regions

## For researchers

- Future studies consider molecular studies will provide to investigate the causative role of early life famine exposure in the programming of future MetS, hypertension and diabetes mellitus.
- ➤ Consider age matched comparison between famine exposed and non-exposed cohorts to see the real effect of famine on ongoing adulthood chronic disease.

#### Reference

- Gebreyes YF, Goshu DY, Geletew TK, Argefa TG, Zemedu TG, Lemu KA, et al.
   Prevalence of high bloodpressure, hyperglycemia, dyslipidemia, metabolic syndrome and their determinants in Ethiopia: Evidences from the National NCDs STEPS. 2018;1–18.
- 2. Alberti KGMM, Eckel RH, Grundy SM, Zimmet PZ, Cleeman JI, Donato KA, et al. Harmonizing the Metabolic Syndrome International Atherosclerosis Society; and International Association for the Study of Obesity. 2009;(120):1640–5.
- 3. The IDF consensus worldwide definition of the Metabolic Syndrome, 2006;
- 4. Cipullo P, Alberto L, Ciorlia S, Cesarino B, Moreira GC, Vilela-martin F. Prevalence of Metabolic Syndrome: Association with Risk Factors and Cardiovascular Complications in an Urban Population. PLOS/ONE. 2014;9(9):1–8.
- 5. Barker DJP, Osmond C, Kajantie E, Eriksson JG. Growth and chronic disease: findings in the Helsinki Birth Cohort. 2009;36(2):445–58.
- 6. Barker DJP. The developmental origins of chronic adult disease. 2004;(8):26–33.
- 7. Bleker LS, Rooij SR De, Painter RC, Velde N Van Der, Roseboom TJ. Prenatal Undernutrition and Physical Function and Frailty at the Age of 68 Years: The Dutch Famine Birth Cohort Study. 2016;71(10):1306–14.
- 8. Lindeboom M, Portrait F, Berg GJ Van Den. Long-run effects on longevity of a nutritional shock early in life: The Dutch Potato famine of 1846 1847. J Health Econ. 2010;29(5):617–29.
- 9. Birarra MK. Metabolic syndrome among type 2 diabetic patients in Ethiopia: a cross-sectional study. BMC. 2018;18(149):1–12.
- 10. Ford ND, Behrman JR, Hoddinott JF, Maluccio JA, Martorell R, Ramirez-zea M, et al. Exposure to improved nutrition from conception to age 2 years and adult cardiometabolic disease risk. Lancet Glob Heal. 2018;6(8):875–84.
- 11. Dercon S, Porter C. Live aid revisited: long-term impacts of the 1984 Ethiopian famine on children. 2010: 15-20
- 12. GILL P. Famine and foreigners Ethiopia since live AID. 2010
- 13. Devereux S. famine in the twenty century. 2000;1: 1-12.
- 14. Kidane A. Mortality estimates of the 1984-85 Ethiopian famine. Scand J SOC Mcd.

- 1990;18(6):281–6.
- 15. Chen Y, Zhou L. The long-term health and economic consequences of the 1959 1961 famine in China. 2007;26:659–81.
- 16. Wang N, Wang X, Li Q, Han B, Chen Y, Zhu C, et al. The famine exposure in early life and metabolic syndrome in adulthood. Clin Nutr 2017;36(1):253–9. Available from: http://dx.doi.org/10.1016/j.clnu.2015.11.010
- 17. The New England Journal of Medicine Downloaded from nejm.org at uc shared journal collection on December 10, 2014.
- 18. Lumey LH. Prenatal Famine and Adult Health. NIH Public Access. 2013;32:2–26.
- 19. Langley-Evans SC. Nutrition in early life and the programming of adult disease : a review. J Hum Nutr Diet. 2014;
- 20. Wang P, Wang J, Lei Y, Xiao L, Luo Z. Impact of Fetal and Infant Exposure to the Chinese Great Famine on the Risk of Hypertension in Adulthood. PLOS/ONE. 2012;7(11):1–8.
- 21. Anping YLI. Exposure to the Chinese Famine in Early Life and the Risk of Metabolic Syndrome in Adulthood. Diabetes Care 2011;34:1014–8.
- 22. Roseboom TJ, Painter RC, Abeelen AFM Van, Veenendaal MVE, Rooij SR De. Hungry in the womb: What are the consequences? Lessons from the Dutch famine. Maturitas 2011;70(2):141–5.
- 23. Li Y, He Y, Qi L, Jaddoe VW, Feskens EJM, Yang X, et al. Exposure to the Chinese Famine in Early Life and the Risk of Hyperglycemia and Type 2 Diabetes in Adulthood. 2010;59
- 24. Painter RC, Roseboom TJ, Bleker OP. Prenatal exposure to the Dutch famine and disease in later life: An overview. 2005;20:345–52.
- 25. Roseboom T, Rooij S De, Painter R. The Dutch famine and its long-term consequences for adult health. 2006
- 26. WHO Progress monitor report 2017. .
- 27. Ethiopia steps report on risk factors for non-communicable diseases and prevalence of selected ncds ethiopia steps report on risk factors for chronic non-communicable diseases and. 2016:183–203.
- 28. Tran A, Gelaye B, Girma B, Lemma S, Berhane Y, Bekele T, et al. Prevalence of

- Metabolic Syndrome among Working Adults in Ethiopia. Int J of Hypertension. 2011;2011:2–9.
- 29. Tachebele B, Abebe M, Addis Z, Mesfin N. Metabolic syndrome among hypertensive patients at University of Gondar Hospital, North West Ethiopia: a cross sectional study. BMC. 2014;14(177):1–9.
- 30. Khorrami Z, Etemad K, Yarahmadi S, Khodakarim S, Kameli M, Hezaveh M, et al. Urbanization and noncommunicable disease (NCD) risk factors: WHO STEPwise Iranian NCD risk factors surveillance in 2011. 2011;2011:469–79.
- 31. Biggs ML, Mukamal KJ, Luchsinger JA, Ix JH, Carnethon MR, Newman AB, et al. NIH Public Access. 2011;303(24):2504–12.
- 32. Suliga E. Dietary Patterns in Relation to Metabolic Syndrome among Adults in Poland : A Cross-Sectional Study. Nutrients. 2017;9(1366):1–15.
- 33. Cohen MP, Stern E, Rusecki Y, Zeidler A. High Prevalence of Diabetes in Young Adult Ethiopian Immigrants to Israel. Diabetes Care. 1988;37.
- 34. The Federal Democratic Republic of Ethiopia Ministry of health. Health Sector Transformation Plan. 2015;16–23.
- 35. Vickers MH. Early Life Nutrition, Epigenetics and Programming of Later Life Disease. Nutrients. 2014;6:2165–78.
- 36. Vaiserman AM. Early-Life Nutritional Programming of Type 2 Diabetes: Experimental and Quasi-Experimental Evidence. Nutrients. 2017;9(236):2–16.
- 37. Shia Z. Early life exposure to Chinese famine modifies the association between hypertension and cardiovascular disease. J Hypertens. 2017;35(1):1–7.
- 38. Rooij SR De, Painter RC, Holleman F, Bossuyt PMM, Roseboom TJ. The metabolic syndrome in adults prenatally exposed to the Dutch. 2018;(April):3–8.
- 39. Hult M, Tornhammar P, Ueda P, Chima C, Bonamy AE, Ozumba B, et al. Hypertension, Diabetes and Overweight: Looming Legacies of the Biafran Famine. 2010;5(10):1–8.
- 40. Barker D, Adair L. Early nutrition and adult outcomes. 2013;6736(13):10–1.
- 41. Baird J, Jacob C, Barker M, Fall CHD, Hanson M, Harvey NC, et al. Developmental Origins of Health and Disease: A Lifecourse Approach to the Prevention of Non-Communicable Diseases. 2017;5(14):2–12.
- 42. Anping YLI. Exposure to the Chinese Famine in Early Life and the Risk of Metabolic

- Syndrome in Adulthood. 2011;34:1014–8.
- 43. Wang N, Wang X, Li Q, Han B, Chen Y, Zhu C, et al. The famine exposure in early life and metabolic syndrome in adulthood. 2017;36 (4) 12-25.
- 44. Yu C, Wang J, Wang F, Han X, Hu H, Yuan J, et al. Exposure to the Chinese famine in early life and metabolic syndrome prevalence risk in adults Department of Occupational and Environmental Health and State Key Laboratory of Dongfeng Central Hospital, Nutrition. 2017;10 (2) 11-17. Available from: https://doi.org/10.1016/j.nut.2017.12.013
- 45. Zheng X, Wang Y, Ren W, Luo R, Zhang S, Zhang JH, et al. Risk of metabolic syndrome in adults exposed to the great Chinese famine during the fetal life and early childhood. Eur J Clin Nutr. 2011;66(2):231–6. Available from: http://dx.doi.org/10.1038/ejcn.2011.161
- 46. Li C, Lumey LH. Social and Economic Determinants Exposure to the Chinese famine of 1959 61 in early life and long-term health conditions: a systematic review and meta-analysis. Int J Epidemiol. 2017;46(4):1157–70.
- 47. Mph LK, Shasha-lavsky H, Eilat-zanani S, Edri-shur A, Shasha SM. Chronic Health Conditions in Jewish Holocaust Survivors Born during World War II. IMAJ. 2015;17.
- 48. Li Y, Jaddoe VW, Qi L, He Y, Lai J, Wang J, et al. Exposure to the Chinese famine in early life and the risk of hypertension in adulthood. J Hypertens 2011,. 2002;1085–92.
- 49. Stein AD, Zybert PA, Bruin KVDP, Lumey LH. Exposure to famine during gestation, size at birth, and blood pressure at age 59 y: evidence from the dutch famine. 2006;759–60.
- 50. Bercovich E, Keinan-boker L, Shasha SM. Long-Term Health Effects in Adults Born during the Holocaust. 2014;16:203–7.
- 51. Zhang Y, Liu X, Wang M, Song Y, Zhang L, You Y, et al. Risk of Hyperglycemia and Diabetes after Early-Life Famine Exposure: A Cross-Sectional Survey in Northeastern China. Int J Environ Res Public Heal. 2018;15(1125):2–10.
- 52. Wang N, Cheng J, Han B, Li Q, Chen Y, Xia F, et al. Exposure to severe famine in the prenatal or postnatal period and the development of diabetes in adulthood: an observational study. Diabetologia [Internet]. 2016; Available from: http://dx.doi.org/10.1007/s00125-016-4148-4
- 53. Wang N, Cheng J, Han B, Li Q, Chen Y, Xia F, et al. Exposure to severe famine in the prenatal or postnatal period and the development of diabetes in adulthood: an

- observational study. Diabetologia [Internet]. 2017;262–9. Available from: http://dx.doi.org/10.1007/s00125-016-4148-4
- 54. Cai H, Huang J, Xu G, Yang Z, Liu M, Mi Y, et al. Prevalence and Determinants of Metabolic Syndrome among Women in Chinese Rural Areas. plose ONE. 2012;7(5):1–11.
- 55. Stein AD, Rundle A, Wada N, Goldbohm RA, Lumey LH. Associations of Gestational Exposure to Famine with Energy Balance and Macronutrient Density of the Diet at Age 58 Years Differ According to the Reference Population Used. J Nutr. 2009;139(8):1555–61.
- 56. Ethiopia demographic and health survey 2016.
- 57. Nichols EK, Byass P, Chandramohan D, Clark SJ, Flaxman AD, Jakob R, et al. The WHO 2016 verbal autopsy instrument: An international standard suitable for automated analysis by InterVA, InSilicoVA, and Tariff 2016;1–9.
- 58. FAO. Dietary Assessment. 2018.
- 59. WHO\_1995\_The use and interpratation of Anthropometry.pdf.
- 60. Sinaga M, Worku M, Yemane T, Tegene E, Wakayo T, Girma T, et al. Optimal cut-off for obesity and markers of metabolic syndrome for Ethiopian adults. Nutr J. 2018;1–12.
- 61. Stanner SA, Bulmer K, Andrès C, Lantseva OE, Borodina V, Poteen V V., et al. Does malnutrition in utero determine diabetes and coronary heart disease in adulthood? Results from the Leningrad siege study, a cross sectional study. Bmj. 1997;315(7119):1342–8.
- 62. Survey NE. Alcohol Consumption and the Prevalence of the Metabolic Syndrome in the U . S . 2004;27(12).
- 63. Jaffe A, Giveon S, Wulffhart L, Oberman B, Freedman L. Diabetes among Ethiopian Immigrants to Israel: Exploring the Effects of Migration and Ethnicity on Diabetes Risk. 2016;81:1–11.
- 64. Wang P, Wang J, Lei Y, Xiao L, Luo Z. Impact of Fetal and Infant Exposure to the Chinese Great Famine on the Risk of Hypertension in Adulthood. 2012;7(11):1–8.
- 65. Stanner SA, Yudkin JS. Fetal Programming and the Leningrad Siege Study. 2001;4(5):287–92.

# Annex 1. Participant's consent informant

## Good morning/afternoon;

Hello. My name is ------I am interviewing participants in this zone to assess adulthood metabolic status among survived adult of the Ethiopian Great Famine in North wollo zones

#### Dear respondents;

Thank you for being cooperative to answer for the valid and effective completion. All what you have told will be kept in secret and you are not expected to tell your name and your name is not going to be registered. A code number will identify every participant. The interview is voluntary; you have the right to participate or not to participate at any time during the interview. Your refusal will not have any effect on the services that you or any of your family receives. Your participation is important to fulfil the study and in order to help to design appropriate intervention to reduce the effect of starvation during the prenatal in this zone and other similar settings. Therefore, you are kindly requested to give genuine answers according to the questions. You can ask questions to the interviewer for clarification.

Moreover, as part of this research project, we are asking people over the Kebeles to measure their height, weight, waist and hip circumference, blood pressure measurement and take blood samples for fasting blood glucose test and lipid profile tests. Besides, there will be interview on sociodemographic, behavioral risk factors, dietary habit assessment. This is to determine metabolic syndrome which comprises hypertension, dyslipidemia, diabetes mellitus and abdominal obesity. These problems are a serious health problem that usually results from starvation during intrauterine life, childhood period and adolescent period. This survey will assist the government to develop programs to prevent and treat chronic disease.

For these test, we will need a few drops of blood from a finger. The equipment used to take the blood is clean and completely safe. It has never been used before and will be thrown away after we take your blood. The blood will be tested for fasting blood glucose immediately, and the result will be told to you right away. The result will be kept strictly confidential and will not be shared with anyone other than members of our survey team.

Risk and /or Discomfort

By participating in this research project you may lose around \_\_\_ minutes. There is no risk in

participating in this research. Benefits

If you are participating in this research, there may not be direct benefit to you but your participation

is likely to help us in showing the effect of prenatal starvation on adulthood metabolic syndrome

that help in decision making for stakeholders. Incentives/Payments for Participating

You will not be provided any incentives or payment to take part in this project. Confidentiality

The information collected for this research project will be kept confidential and information about

you that will be collected by this study will be stored in a file, without your name, but a code

number assigned to it. This data will not be revealed to anyone except the principal investigator

and assistants will be kept locked with key.

Right to Refusal

You have full right to refuse from participating in this research and this will not affect you from

getting any kind of health service. You have also the full right to leave from this study at any time

you wish, without losing any of your right. Person to contact

This research project will be reviewed and approved by the ethical committee of the Jimma

University. If you want to know more information you can contact through the address below. If

you have any question you can contact any of the following individuals and you may ask at any

time you want.

1. Mr .Habtamu Hasen MSc student at Jimma University

Mobile: +25138641297 e.mail: habtamu130@gmail.com

Is the information/ objective clear?

1) Yes

2) No

Are you willing to participate in the interview? It is up to you to decide

1) Yes

2) No

Thank you!

# Annex 2: English version Questionnaire

**Instruction:** This is a structured questionnaire designed for the study of adulthood metabolic profile of the Great Ethiopian Famine survivors who in the age of 30- 36 years at the time of the famine in North Wollo provinces, Northeast Ethiopia, 2019.

Note: Get consent before interviewing performing any activity.

Questi	onnaire ID No/code		
Date o	of interview		
Zone_	KebeleGot	Villa	ige
GT GT			
	ION A: QUESTIONS RELATED TO 1975-77 E.C ETHIC	)PIAN GRI	EAT FAMINE and Verbal
	sy Questionnaire		
A1	Place of Birth		_
A2	Age	ye	ars
A3	Birth date	//_	
		I don't kno	OW
A3a	If <b>A3 unknown</b> : What was your age during 1975-77 E.C		month
	Great Ethiopian Famine (kifu ken)?		_ years
	Probe Use local calendar	0 pregnant	į.
	Age in completed years	I don't kno	OW
A3b	If <b>A3a</b> ( <b>pregnant</b> ): What was your gestational age?		weeks
A4	How do you feel about the burden of 1975-77 E.C Great	1. Ha	ardly
	Ethiopian Famine?	2. Li	ttle
	If he/she can remember?	3. I d	lon't know
A5	Did you have sufficient foods or resources to cope up the	1 Ye	es
	famine?	0 No	0
A6	Did anybody die due to famine in your home?	1 Ye	es
		0 No	0
A6a	If yes to A6, Number of males deceased?		number
		Id	lon't know
A6b	If yes to A6, Number of females deceased		number
		I don'	t know
A6c	If yes to A6, what was the place and date of death	Place of de	eath
		Date of de	ath
		I don't kno	ow
A6d	If yes to Question A6, how many people died in your home?	(1	number of people dies in the
		Household	1)
SECT	FION B. Household Wealth index		
Now I	I will ask you about some fixed assets that your house	ehold have	,
Does t	he household have any of the following properties? (Circle)	Yes	No
B1	Functioning radio/Tape recorder/CD player	1	0

B2	Functioning Television	1	0
В3	Solar light	1	0
B4	Kerosene stove	1	0
B5	Electric stove	1	0
B6	Bicycle	1	0
B7	Motor Cycle	1	0
B8	Cart/Gari	1	0
B9	Watch (Hand/Wall)	1	0
B10	Mobile phone	1	0
B11	Plough	1	0
B12	Sofa	1	0
B13	Spring mattress	1	0
B14	Sponge/Foam mattress	1	0
B15	Cotton mattress	1	0
B16	Grass Mattress	1	0
B17	Chair/Stool	1	0
B18	Generator	1	0
B19	Milling	1	0
B20	Water pump	1	0
	Does the household have any of the following animals?	1.Yes	How many?
		0. No	
B21	Oxen		
B22	Cows		
B23	Horse/mules/donkey/Camel		
B24	Goats/Sheep		

	SECTION C: Background information of the participants				
I first	I first ask you questions about yourself and your family:				
C1	What is the sex of the participant?	1 Male			
	(Record Male/Female as observed)	0 Female			
C2	Where is your place of residence?	1. Rural			
		2. Urban			
C3	What is your religion?	1. Orthodox			
		2. Muslim			
		3. Protestant			
		4. Catholic			
		5. Other specify)			
C4	What is your ethnicity?	1. Amhara			
		2. Oromo			
		3. Tigrie			
		4. Other(specify)			
C5	What is your educational status?	1. Can't read and write			
		2. Primary school (1-8)			
		3. Secondary school (9-12)			
		4. More than secondary (>12)			
C6	What is your current marital status?	1. Single			

		<ul><li>2. Married</li><li>3. Divorced/Separated</li></ul>
		4. Widowed
C7	What is your occupation?	Government employee
		2. Non-government employee
		3. Farmer
		4. Housewife
		5. Merchant
		6. Other (specify)
<b>C8</b>	Do you have a child less than 5 years?	1 Yes
		0 No
<b>C9</b>	How many family members including you in	number
	your house	

SECTIO	SECTION D: Substance use of the respondents				
1.	1. Tobacco Use				
Now I a	m going to ask you some questions about tobacco use.				
D1	Do you currently smoke any tobacco products, such as cigarettes, cigars or pipes, gaya?	1 yes 0 No, If no, go to D5			
D2	Do you currently smoke tobacco products daily?	1 Yes 0 No			
D3	How old were you when you first started smoking?	Age in years 77 Don't know			
D4	On average, how many of the following do you smoke each day/week? (IF LESS THAN DAILY, RECORD WEEKLY)  (RECORD FOR EACH TYPE)	<ol> <li>Manufactured         cigarettes</li> <li>Gaya</li> <li>Other (specify)         77 Don't know</li> </ol>			
D5	In the past 12 months, did you ever smoke any tobacco products?	1 Yes 0 No			
D6	In the past 12 months, did you ever smoke daily?	1 Yes 0 No			
D7	Do you currently use any smokeless tobacco such as [snuff, chewing tobacco]?	1 Yes 0 No If No, go to D10			
D8	Do you currently use smokeless tobacco products daily?	1 Yes 0 No			

D9	On average, how many times a day do you use (IF LESS THAN DAILY, RECORD WEEKLY). (RECORD FOR EACH TYPE, USE SHOWCARD)	1. Snuff, by mouth         2. Snuff, by nose         3. Chewing tobacco         4. Other (specify)         Don't Know
D10	In the past 12 months, did you ever use smokeless tobacco such as [snuff, chewing tobacco)?	1 Yes 0 No
D11	During the past 30 days, did someone smoke in your home?	1 yes 0 No
D12	During the past 30 days, did someone smoke in closed areas in your workplace (in the building, in a work area or a specific office)?	1 Yes 0 No
	Alcohol consumption	
The next	t questions ask about the consumption of alcohol.	
D13	Have you ever consumed any alcohol such beer, Tella, korefie, Tej, Arake, wine, [add other local examples]?	1. Yes 0 No if no go to D22
D14	Have you consumed an alcoholic drink within the past 12 months?	1 yes 0 No
D15	During the past 12 months, how frequently have you had at least one alcoholic drink?	<ol> <li>Daily</li> <li>5-6 days per week</li> <li>1-4 days per week</li> <li>1-3 days per month</li> <li>Less than once a month</li> </ol>
D16	Have you consumed an alcoholic drink within the past 30 days?	1. Yes 0. No If No, Go to D22
D17	During the past 30 days, on how many occasions did you have at least one alcoholic drink?	Number Don't know
D18	During the past 30 days, when you drank alcohol, on average, how many standard alcoholic drinks did you have during one drinking occasion?	Number 77 Don't know
D19	During the past 30 days, when you consumed an alcoholic drink, how often was it with meals? Please do not count snacks.	<ol> <li>Usually with meals</li> <li>Sometimes with meals</li> <li>Rarely with meals</li> <li>Never with meals</li> </ol>
D20	During the past 7 days, how many standard alcoholic drinks did you have?	Number Don't know
D21	During the past 7 days, did you consume any homebrewed alcohol, like Tella, Tej, Katikalla, korefie?	1 Yes 0 No
	at use	
Now I at D22	m going to ask you some questions about Khat chewing  Have you ever chewed Khat?	1 Yes
D22	Thave you ever chewed Khat?	0 No, if No go to E1

D23	Do you currently chew Khat?	1 Yes 0 No
D24	During the past 12 months, how frequently did you chew Khat?	<ol> <li>Daily</li> <li>5-6 days per week</li> <li>3-4 days per week</li> <li>1-2 days per week</li> <li>1-3 days per month</li> <li>Less than once a month</li> </ol>
D25	On average, how many bundles of Khat do you chew each/day week? (IF LESS THAN DAILY, RECORD WEEKLY)	Bundles of Khat Don't know
D26	Do you currently smoke tobacco products while chewing Khat?	1 Yes 0 No
D27	In the past 12 months, did you ever smoke tobacco products while chewing Khat?	1 Yes 0 No
D28	Do you currently drink alcohol after you chew Khat?	1 Yes 0 No

**SECTION E: PHYSICAL ACTIVITY:** I am going to ask you about the time you spent being physically active in the last 7 days. Please answer each question even if you do not consider yourself to be an active person. Think about the activities you do at work, as part of your house and yard work, to get from place to place, and in your spare time for recreation, exercise or sport.

**READ**: Now, think about all the *vigorous* activities which take *hard physical effort* that you did in the last 7 days. Vigorous activities make you breathe much harder than normal and may include heavy lifting, digging, aerobics, or fast bicycling. Think only about those physical activities that you did for at least 10 minutes at a time.

E1	During the <b>last 7 days</b> , on how	days	[Interviewer clarification:
	many days did you do	77 Don't	Think only about those physical
	vigorous physical activities?	Know/not	activities that you do for at least
		sure	10 minutes at a time.].
			[Interviewer note: If respondent
			answers zero, or does not know,
			skip to Question E3]
<b>E2</b>	How much time did you		If the respondent can't answer
	usually spend doing vigorous	Minutes per day	because the pattern of time spent
	physical activities on one of	77 Don't Know/not	varies widely from day to day,
	those days?	sure	ask Q E2-1
E2-1	"How much time in total	hours per week	
	would you spend over the last	Don't Know/not sure	
	7 days doing vigorous		
	physical activities?"		

**READ**: Now think about activities which take *moderate* physical effort that you did in the last 7 days. Moderate physical activities make you breathe somewhat harder than normal and may include carrying light loads, bicycling at a regular pace, or doubles tennis. Do not include walking. Again, think about only those physical activities that you did for at least 10 minutes at a time.

<b>E3</b>	During the <b>last 7 days</b> , on how	days	[Interviewer clarification:
	many days did you do	1. Don't Know/not sure	Think only about those physical
	<b>moderate</b> physical activities?		activities that you do for at least
			10 minutes at a time.]
			[Interviewer note: If respondent
			answers zero, refuses or does not
			know, skip to Question E5]
<b>E4</b>	How much time did you	Minutes per day	If the respondent can't answer
	usually spend doing moderate	77 Don't Know/not sure	because the pattern of time spent
	physical activities on one of		varies widely from day to day,
	those days?		ask Q E4-1
E4-1	What is the total amount of	hours per week	
	time you spent over the <b>last 7</b>	77 Don't Know/not sure	
	days doing moderate physical		
	activities?		

**READ**: Now think about the time you spent **walking** in the last 7 days. This includes at work and at home, walking to travel from place to place, and any other walking that you might do solely for recreation, sport, exercise, or leisure.

E5	During the last 7 days, on how	days per week	[Interviewer clarification:		
	many days did you walk for at	77 .Don't Know/not sure	Think only about those physical		
	least 10 minutes at a time?		activities that you do for at least		
			10 minutes at a time.]		
			If respondent answers zero,		
			refuses or does not know, skip to		
			Question E7]		
<b>E6</b>	How much time did you	Hours per days	If the respondent can't answer		
	usually spend walking on one	77 .Don't Know/not	because the pattern of time spent		
	of those days?	sure	varies widely from day to day,		
			ask Q E6-1		
E6-1	"What is the total amount				
	of time you spent walking over	77 .Don't Know/not sure			
	the last 7 days?"				
DEAD. N	<b>READ</b> : Now think about the time you spent sitting on week days during the last 7 days. Include time				
	· ·	•			
spent at v	work, at home, while doing cours	e work, and during leisure	time. This may include time spent		
spent at v	work, at home, while doing cours a desk, visiting friends, reading o	e work, and during leisure or sitting or lying down to w	time. This may include time spent yatch television.		
spent at v	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how	e work, and during leisure r sitting or lying down to w minutes per weekday	time. This may include time spent		
spent at v	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually	e work, and during leisure r sitting or lying down to w minutes per weekday	time. This may include time spent ratch television.  Interviewer clarification: Include time spent lying down		
spent at v	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend <i>sitting/reclining</i> on a	e work, and during leisure r sitting or lying down to w minutes per weekday	time. This may include time spent vatch television.  Interviewer clarification: Include time spent lying down (awake) as well as sitting]		
spent at v	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually	e work, and during leisure r sitting or lying down to w minutes per weekday	time. This may include time spent ratch television.  Interviewer clarification: Include time spent lying down (awake) as well as sitting] [Interviewer probe: If the		
spent at v	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend <i>sitting/reclining</i> on a	e work, and during leisure r sitting or lying down to w minutes per weekday	time. This may include time spent ratch television.  Interviewer clarification: Include time spent lying down (awake) as well as sitting]  [Interviewer probe: If the respondent can't answer because		
spent at v	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend <i>sitting/reclining</i> on a	e work, and during leisure r sitting or lying down to w minutes per weekday	time. This may include time spent ratch television.  Interviewer clarification: Include time spent lying down (awake) as well as sitting] [Interviewer probe: If the respondent can't answer because the pattern of time spent varies		
spent at v	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend <i>sitting/reclining</i> on a	e work, and during leisure r sitting or lying down to w minutes per weekday	Interviewer clarification: Include time spent lying down (awake) as well as sitting  [Interviewer probe: If the respondent can't answer because the pattern of time spent varies widely from day to day, go to		
spent at v sitting at E7	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend sitting/reclining on a weekday?	e work, and during leisure or sitting or lying down to w minutes per weekday 77 .Don't Know/not sure	time. This may include time spent ratch television.  Interviewer clarification: Include time spent lying down (awake) as well as sitting] [Interviewer probe: If the respondent can't answer because the pattern of time spent varies		
spent at v	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend sitting/reclining on a weekday?	e work, and during leisure or sitting or lying down to we minutes per weekday 77 .Don't Know/not sure	Interviewer clarification: Include time spent lying down (awake) as well as sitting  [Interviewer probe: If the respondent can't answer because the pattern of time spent varies widely from day to day, go to		
spent at v sitting at E7	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend <i>sitting/reclining</i> on a weekday?  "What is the total amount of time you spent <i>sitting</i> last	e work, and during leisure or sitting or lying down to warminutes per weekday 77 .Don't Know/not sure Hours on WednesdayMinutes on	Interviewer clarification: Include time spent lying down (awake) as well as sitting  [Interviewer probe: If the respondent can't answer because the pattern of time spent varies widely from day to day, go to		
spent at v sitting at E7	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend sitting/reclining on a weekday?	e work, and during leisure or sitting or lying down to we minutes per weekday 77 .Don't Know/not sure Hours on WednesdayMinutes on Wednesday	Interviewer clarification: Include time spent lying down (awake) as well as sitting  [Interviewer probe: If the respondent can't answer because the pattern of time spent varies widely from day to day, go to		
spent at v sitting at E7	work, at home, while doing cours a desk, visiting friends, reading of During the last 7 days, how much time did you usually spend <i>sitting/reclining</i> on a weekday?  "What is the total amount of time you spent <i>sitting</i> last	e work, and during leisure or sitting or lying down to warminutes per weekday 77 .Don't Know/not sure Hours on WednesdayMinutes on	Interviewer clarification: Include time spent lying down (awake) as well as sitting  [Interviewer probe: If the respondent can't answer because the pattern of time spent varies widely from day to day, go to		

# **SECTION F:** Food frequency questionnaire (FFQ)

Thinking back on the last 12 months, please tell me the how often you consumed each of the following items. **0 IF NEVER CONSUMED THE ITEM.** 

Food list	Frequency	Food list	Frequency
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
F1. Teff	3-4 times per week		3-4 times per week
ri. ren	5-6 times per week	F2. Maize	5-6 times per week
	1 time per day	1.72. Waize	1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
F3. Barely	3-4 times per week		3-4 times per week
13. Barciy	5-6 times per week	F4. Wheat	5-6 times per week
	1 time per day	including bread	1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
F5.	2-3 times per month		2-3 times per month
Sorghum/mil	1-2 times per week	F6. Rice	1-2 times per week
let	3-4 times per week		3-4 times per week
	5-6 times per week		5-6 times per week
	1 time per day		1 time per day

	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
G7. Beef	3-4 times per week	G8. Chicken	3-4 times per week
G7. Beel	5-6 times per week	Go. Chicken	5-6 times per week
	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less	G10. Liver	1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
G9. Fish	3-4 times per week		3-4 times per week
G). I isii	5-6 times per week	G10. Livei	5-6 times per week
	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
G11.Goat/la	2-3 times per month	G12.preserved	2-3 times per month
mb	1-2 times per week	meat	1-2 times per week
ino	3-4 times per week	meat	3-4 times per week
	5-6 times per week		5-6 times per week

	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
F13. Milk	3-4 times per week	F14. Cheese	3-4 times per week
1 13. WIIIK	5-6 times per week	1 14. Cheese	5-6 times per week
	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
F15. Butter	3-4 times per week	F16. Egg	3-4 times per week
113. Butter	5-6 times per week	110. 255	5-6 times per week
	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
F17. Nuts	1-2 times per week	F18. Oil	1-2 times per week
	3-4 times per week		3-4 times per week
	5-6 times per week		5-6 times per week
	1 time per day		1 time per day

	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
E10 Poons	1-2 times per week		1-2 times per week
F19.Beans,	3-4 times per week	F20. Sweet potato	3-4 times per week
peas, lentils, chickpeas	5-6 times per week	120. Sweet potato	5-6 times per week
cinckpeas	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
F21.Potatoes	3-4 times per week	F22. Carrot	3-4 times per week
121.1 otatoes	5-6 times per week	122. Carrot	5-6 times per week
	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
F23.Tomato	1-2 times per week	F24. Cauliflower	1-2 times per week
	3-4 times per week		3-4 times per week
	5-6 times per week		5-6 times per week
	1 time per day		1 time per day

	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
F25.Leafy	1-2 times per week		1-2 times per week
green	3-4 times per week	F26. Avocado	3-4 times per week
vegetables	5-6 times per week	120. Avocado	5-6 times per week
vegetables	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
F27. Papaya	3-4 times per week	F28. Bananas	3-4 times per week
127.1 apaya	5-6 times per week	1 20. Dananas	5-6 times per week
	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
F29.	2-3 times per month		2-3 times per month
Pineapple	1-2 times per week	F30. Oranges	1-2 times per week
Тіпопрію	3-4 times per week		3-4 times per week
	5-6 times per week		5-6 times per week
	1 time per day		1 time per day

	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
F31. Onion	Never consumed	F32. Garlic	Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
	3-4 times per week		3-4 times per week
	5-6 times per week		5-6 times per week
	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
	1-2 times per week		1-2 times per week
F33. Ginger	3-4 times per week	F34. Chills	3-4 times per week
133. Ginger	5-6 times per week	1 54. Cillis	5-6 times per week
	1 time per day		1 time per day
	2-3 times per day		2-3 times per day
	4-5 times per day		4-5 times per day
	6 or more times per day		6 or more times per day
	Never consumed		Never consumed
	1 time per month or less		1 time per month or less
	2-3 times per month		2-3 times per month
F35. Coffee	1-2 times per week	F36.Local drink	1-2 times per week
	3-4 times per week		3-4 times per week
	5-6 times per week		5-6 times per week
	1 time per day		1 time per day

	2-3 times per day		2-3 times per day	
	4-5 times per day		4-5 times per day	
	6 or more times per day		6 or more times per day	
	Never consumed		Never consumed	
	1 time per month or less		1 time per month or less	
	2-3 times per month		2-3 times per month	
	1-2 times per week		1-2 times per week	
F37. Soft	3-4 times per week	F38. Salt	3-4 times per week	
drink	5-6 times per week	1 736. Sait	5-6 times per week	
	1 time per day		1 time per day	
	2-3 times per day		2-3 times per day	
	4-5 times per day		4-5 times per day	
	6 or more times per day		6 or more times per day	
	Never consumed	F40. Over the past	1 yes	
	1 time per month or less	12 months, did you	0 No	
	2-3 times per month skip any of your 1-2 times per week meals?			
F39. Eating	3-4 times per week	F40a. If yes, which	1. Breakfast	
outside the	5-6 times per week	of the following	<ul><li>2. Lunch</li><li>3. Dinner</li></ul>	
home	1 time per day	meals did you		
		skip?		
	2-3 times per day			
	4-5 times per day			
	6 or more times per day			
F41. What type of in your household	oil or fat is most often <i>used for meal</i> d?	2. Pal 3. Bu	getable oil m oil tter or ghee her ( <i>specify</i> )	
F42. do you eat the breakfast, lunch ar	nat were not prepared at a home? By and dinner.		ner (specify)	

	for high blood pressure prescribed by a doctor or other health worker?	0 No	
J (2)	Has any of your family members (biological	1yes	
	parents, siblings or children) ever had raised	0 No	
	blood pressure or hypertension?		
	History of Diabetes		
J (3)	Are you currently receiving any treatments/advice	1 Yes	
	for diabetes prescribed by a doctor or other health	0 No	
	worker?		
J (4)	Has any of your family members (biological	1 Yes	
	parents, siblings or children) ever had raised	0 No	
	blood sugar or Diabetes?		
SECTIO	ON K: ANTHROPOMETRIC MEASURENT		
K1	Height	(cm)[for measurement 1	
	Double measurement	(cm)[for measurement 2]	
K2	Weight	kg for measurement 1	
	Double measurement	kg for measurement 2	
<b>K3</b>	Waist circumference	Centimeters (cm)	
	No need to measure for pregnant mothers		
K4	Hip circumference	Centimeters (cm)	
	TION L: BLOOD PRESSURE (BP) MEASUREME	<b>-</b>	
L1	Reading 1	Systolic (mmHg)	
		Diastolic(mmHg)	
L1-1	Reading 2	Systolic (mmHg)	
		Diastolic (mmHg)	
SECTIO	ON M: BIOCHEMICAL MEASUREMENTS		
First, I	will measure your blood glucose level.		
This test	is taken to measure raised blood sugar levels which a	are a risk factor for diabetes. For this purpose, blood	
is going	to be collected from a small prick on the finger Durin	ng the past 8 hours have you had anything to eat or	
drink, oti	her than water?		
If the an	swer is No, I will not measure your blood glucose	level	
Time of a	day blood specimen taken (24-hour clock) Hours		
M3	Fasting blood glucose Choose accordingly: mmol/l		
		mg/dl	
M3	Fasting blood glucose Choose accordingly: mmol/l	mg/dl Refused	
	Fasting blood glucose Choose accordingly: mmol/l  Today, have you taken insulin or other drugs (medical)	mg/dl Refused ation) that have 1Yes	
M3	Fasting blood glucose Choose accordingly: mmol/l	mg/dl Refused ation) that have 1Yes	

1 Yes

SECTION J: HISTORY OF HYPERTENSION, DIABETES

J (1)

I First ask you questions about History of Raised Blood Pressure

Are you currently receiving any treatments/advice

Lipids	<b>Lipids profiles:</b> This is to know your blood cholesterol level. Triglyceride test is taken to measure the fasting		
levels o	levels of natural fats in the bloodstream. For this purpose, the blood will be withdrawn from your fingertip.		
M5	M5 Total cholesterol Choose accordingly: mmol/l or mg/dlmg/dl Refused		
M6	During the past two weeks, have you been treated for raised cholesterol with drugs (medication) prescribed by a doctor or other health worker?	1. Ye 0 No	
M7	Triglycerides Choose accordingly: mmol/l or mg/dl	mg/dl Refuse	
M8	HDL Cholesterol Choose accordingly: mmol/l or mg/dl	mg/dl Refuse	

Thank you so much!

#### AMHARIC VERSION QUESTIONERS

በጥናቱ ለመሳተፍ ስለጥናቱ ለተሳታፊዎች የሚሰጥ መረጃ

የአሰራር ሂደት፡ የተጠቀምንበት የጥናት ናሙና አወሳሰድ ቴክኒክ ሕርስዎን እንድናካትት አድርጎናል፡፡ ከሕርሶ መረጃውን በቃለ መጠይቅ፤ ከደም ናሙና መረጃ ሕሰበስባለሁ፡፡ የመረጃ ሰብሳቢዎች የሜታቦሲክ (ተሳላፊ ያልሆኑ ) በሽታና ተያያዥነት ያላቸዉን ምክንያቶች በአዋቂ የስኳር በሽተኞችን ፡ዴምግፊትሕናየሰዉነትስብመጠን ለማወቅ ጥያቄዎች ይጠየቃሉ ፡፡ ጥያቄዎችም ማህበራዊና ስነ- ህዝባዊ ፣ ማህበራዊ- ኢኮኖሚያዊ፣ ጤናነክ

እና ባህሪያዊ *ጉዳ*ዮች ዙሪያ በተ*መ*ለከተ ይጠይቃሉ። ከእርሶ 5 ሚሊ ሊትር ደም የስካ*ር* እና የሰዉነትስብ መጠኖን ለመለካት እንጠቃመለን። በዚህ ጥናት አንደሚካፈሉ ተስፋ ሕናደር*ጋ*ስን ፡፡መጠይቁ ከ 30 -60 ደቂቃዎች ይወስዳል ፡፡ከዚህም በኃላ ይጠናቀቃል ፡፡ አደ*ጋዎች* ወይም አስመመቸት ና የሚጠበቁ ጥቅሞች በዚህ ጥናት በመሳተልዎ የሚያጋጥምዎት አደጋ የለም ፡፡ ከእርሶ 5 ሚሊ ሊትር ደም የስካር መጠኖን ለመለካት እና ለሴሎች የሜታቦሲክ ችግር አመሳካች መረጃዎችችን ለመስብስብ ፡፡ በጥናቱ ላይ እርሶ የስካርና ደም ግፊት ችግር ከተገኛበዎት ወደ ሆስፒታል ሰበለጠ ህክምና ይላካሉ። በጥናቱ ወቅትም ምን ማድረግ እንዳለበዎት ይነገሮታል።እርስዎ በጥናቱ በመካፈልዎና በሚሰጡት መረጃ የሚያገኙት ልዩ ጥቅም የለም። የስካርና ዴም ግፊት በሽታና ተያያዥነት ያሳቸዉን ምክንያቶች በስኳርና ደም ግፊት በሽተኞቸች ምክንያቶች በተመለከተ መረጃ ለማቅረብና ችግሮቻቸዉን ለመቅረፍ ያስችል ዘንድ ስልቶችን ለመንደፍ ይጠቅማል ፡፡ በነዚህ በስኳርና ደም ግፊት በሽተኞች ህክምና፤ ድጋፍና እንክብካቤ ዙሪያ ለሚሰሩ መንግስታዊና መንግስታዊ ያልሆኑ ድርጅች ፕሮግራሞቻቸውን ለማሻሻልም ይረዳል፡፤ ሚስጢር መጠበቅ የሚሰጡት መልስም ሆነ የጥናቱ ውጤት በሚስጢራዊነት ይጠበቃል። ለዚህ ጥናት የሚሰበሰበው አርስዎን የሚመለከት መረጃ በማህደር የሚቀመጥ ሲሆን ማህደሩን በርስዎ ስም ሳይሆን በተለየ ኮድ ስለሚቀመጥ ከዋናው ተመራማሪ በስተቀር ለማንም አይገለጽም። *ልቃ*ደኝነት

ሕርስዎ በጥናቱ ውስጥ መካፈል ካልፈስጉ መሳተፍ የሰበዎትም ፡፡ ከጥናቱ ውስጥ በጣንኛውም ጊዜ አቋርጠው መውጣት መብትዎ ነው፡፡ በጥናቱ መካፈል በማቋረጥዎ ምንም ነገር አይባሱም በጤና ባለሙያዎች የሚሰጥዎት ድጋፍና እንክብካቤም አይጓደልብዎትም ፡፡ ጥያቄ ካለዎት መረጃ ሰብሳቢውን ይጠይቁ፡፡ ወደፊትም ቢሆን ያልተረዱት ነገር ካለ ሲጠይቁ ይችላሉ፡፡ ተመራጣሪዎቹ በጥናቱ ጊዜ አዲስ መረጃዎች ካጋጠሟችሁ ያሳውቅዎታል፡፡ የተመራጣሪው አድራሻ ከላይ ከተሰጠዎት መረጃዎች ውስጥ ግልጽ ያለሆነለዎት ከሆነ ወይም ተጨጣሪ መረጃ የሚፈልጉ ከሆነ ከዚህ በታች ባለው አድራሻ ተመራጣሪውን ሲያገኙ ይችላሉ፡፡
1.ዶ/ር ቃልኪዳን ሀሰን: ስልክ ቁጥር 0911370862 ኢ-ሜይል newewi333@gmail.com

- 2. ጌታቸዉ አራጌ: ስልክ ቁጥር 0910435581 ኢ-ሜይል getachewarage2004@gmail.com
- 3. ሀብታው ሀሰን: ስልክ ቁጥር 09038641297 ኢ-ሜይል <u>habtamu130@gmail.com</u>
- 4. ከማል ሀጂማህሙድ: ስልክቁጥር 0912255990 ኢ-ሜይል Kemal.mehmoud@gmail.com በዚህ ጥናት ላይ ችግር ከገጠሞት የጂማ ዩንቨርስቲ የጤናና ህክምና ሣይንስ ኮሌጅ የጥናትና ምርምር ስነ ምግባር ኮሚቴ በቢሮ ስልክ ቁጥር0471111450 ወይም በ ፖ.ሣ.ቁ 278 ጂማ መረጃ ሲያገኙ ይችላሉ።

በጥናትና ምርምሩ ለመሳተፍ ፍቃደኞተ*ዎን* ስለመስጠት ከላይ ስለ ጥናቱ *ያ*ለው*ን መ*ረጃውን ተነቦልኛል ወይም በሚገባኝ ቋንቋ ስለጥናቱ አላማ፤የአሰራር ሂደት፤ *ጉዳ*ቶች ወይም አለመመቸት

ና የሚጠበቁ ጥቅሞች፤ሚስጢር መጠበቅ ፤ፌቃደኝነት፡ ለበለጠ መረጃ ዩኒቨርሲቲ የጤናና ህክምና ሣይነስ ኮሌጅ የጥናትና ምርምር ስነ ምግ ተነግሮኛል ፡፡ ስለዚህ ጥናቱ ላይ ለመሳተፍ ተስማምችአለሁ፡፤ የጥናቱ ተሳታፊ ፊርማ	· · · · · · · · · · · · · · · · · · ·
የአማርኛ መጠይቅ	
የዚህ ምርመራ ፕሮጀክት አላማ ተላላፊ ያልሆኑ በሽታዎችና ተያ, ምክንያቶች ለማጥናት ነዉ።ለዚህም ጥናት የሚሳተፉት እድሜያ በላይየሆኑ አዋቂዎች ናቸዉ።በነዚህየህብረተሰብክፍሎችየስኴርናደያ ምን ያህል እንደሆነ እና ተያያዥ ምክንያቶችን በመለየት ለሚመለያ ቻስፒታልወይምየክልሉጤናቢሮበማሳየትትኩረትእንዲሰጡትማድረግ	ቸዉክዓ <i>መቅ</i> ናክዚያ ምግፊት በሽታ ከታቸው ክፍሎች
የጥያቌዉመስያኮድ	
መረጃ	ቀን:
ሰብሳቢው፡ስም ፊርማ የተቆጣጣሪ፡	ቀን፡ ሰዓት

<sub>-</sub>ፌርማ

ጥያቌዉየተጀመረበት**ቀ**ን\_\_

ዞን	ቀበሌ	ታጥ	መንደር	
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<b>ክፍል A</b> 1975-	: ከ 77 ዓ.ም የኢትዮጵያ ታላቁ ሬ <b>ሃብ <i>ጋ</i>ር የተ</b> ያያዙ <sup>(</sup>	<i>ጥያቄዎች</i>	
A1	የተወሰዱበት ቦታ		
A 2	እድ <i>ሜ ዎት</i> ስንት ነወ		_ዓመት
A 3	የተወሰዱበት ጊዜ		ቀ <i>ን</i> /ወር/ዓ.ም
А За	የተወሰዱበት ጊዜ ከሳወቁት በ <sup>1976-77</sup> ዓ.ም የኢትዮጵያ ታላቁ ረዛብ እድሜህ ሽ ስንት ነበር?	7	ዓ <i>ሙት</i> ወር ተረ <b>ግ</b> ኘና ነበር <sup>77</sup> አሳውቀዉም
A3b	<i>መ</i> ልስዎ ተረግኘና ነበር ከሆነ	 አሳው <i>4</i>	
A 4	የሚያስታዉሱ ከሆነ የ <sup>1976-77</sup> ታላቁ የኢትዮጵያውያን ረሃብ <i>እን</i> ደት ይገልጹታል?	<i>መ</i> ካከ <i>ስ</i>	ባም ከባድ 2. ነኛ <sup>ሙ</sup> ንም አይደ <b>ለ</b> ም
A 5	ረዛቡን <b>ለ</b> መቋቋም በቂ ምግብ ወይም ሀብት ነበረወት?	<sup>1</sup> አው 0	
A 6	በቤትዎ ዉስጥ በረዛብ ምክንያት የሞተው ሰው ነበር?	1 አወ 0 የ <b>ሰ</b> ዎ	
A 6a	መልሰዎ ለ A6 አወን ከሆነ የጧች እድሜ ስንት ነበር?		ዓ <i>መት</i> '
A 6b	ለ ጥያቄ A6 መልሰዎ አወን ከሆነ የጧች ጾታ	1 ወን 0 ሰ	
A 6c	መልስዎ ለ A6 አወን ከሆነ የሞተበት ቦታና ጊዜ የት ነው ?	የሞተበት	ቦታ ጊዜ አ <b>ሳ</b> ዉቀዉም
A 6d	መልሰዎ ለ A6 አወን ከሆነ በቤትዎ ውስጥ ስንት ሰዎች ሞተዋል?	ቁጥር	
	የቤተሰብ <i>ሀብት መ</i> ለኪ <i>ያ አሁን</i> ስለቤተሰብ <i>ዎአንዳን</i> ድቋሚንብ		I .
	ሀብት መስኪያ	አ <i>ዎን</i>	የስም
	የሚሰራ ሬዲዮ ቴፕሪኮርደር ሲዲጣጫወቻ አለዎት?	1	0
B 2	የሚሰራ ቴሌቪዥን አለዎት?	1	0
B 3	ሶላር መብራት አለዎት?	1	0
B 4	የነዳጅ ምድጃ አለዎት?	1	0

B 5	የኤሌክትሪክ ምድጃ		1	0
B 6	ብስክሌት አለዎት?		1	0
B 7	ሞተር ሳይክል አለዎት?		1	0
B 8	<i>ጋ</i> ሪ አለዎት?		1	0
B 9	<i>ሠ</i> ኣት የእጅ <i>የግድግዳ</i> 〉 አለዎት?		1	0
B 10	ሞባይል አ <b>ሰ</b> ዎት?		1	0
B 11	<i>ሕረሻ መሬት አ</i> ለ <i>ዎት</i> ?		1	0
B 13	ሰፋ አስ <i>ዎት</i> ?		1	0
B 14	የስፕሪንግ ፍራሽ አለዎት?		1	0
B 15	ስፖንጅ ፍራሽ አለዎት?		1	0
B 16	የጥጥ ፍራሽ አለዎት?		1	0
B 17	<i>ግር</i> ፍራሽአለዎት?		1	0
B 18	ወንበር በርጩ <b>ማ</b> አለ <i>ዎት</i> ?		1	0
B 19	ጀ <b>ነ</b> ሬተርአለዎት?		1	0
B 20	ወፍጮ አለዎት?		1	0
B 21	የውዣፓምፕአሰዎት?		1	0
	ቤት ዉስጥ የቤት <i>እን</i> ስሳት አሉ?		አወን የለም	ስንት ናቸዉ በቁጥር?
B 22	በሬዎች			
B 23	ስም			
B 24	<b>ፌ</b> ረስ <sup>/</sup> በቅሎዎች <sup>/</sup> አህያ/ግመል			
B 25	ፍየሎች በጎች			
	ክፍል C: ስለተሳታ <i>ኤዎች ማ</i> ህበራዊ መ	ረጃ		
ስለራስዖ	<sup>ያ</sup> እና ስለ ቤተሰብዎ ጥ <i>ያቄዎችን እን</i> ጠይቅዎታለን	•		
C 1	ፆታ	1 (	ወንዶ	
		0 .	ሴት	
C 2	መኖርያ በታ	1. <i>ገ</i> ጠር ከተ <b>ማ</b>	2.	
C 3	<b>ሀይማ</b> ኖት·?	<sup>1</sup> አርቶድክስ <sup>2.</sup> ሙስሲም 3. ፕሮቴስታነት 4. ካቶሊክ 5. ሌላ ካለ <u></u> ይጠቀስ <sup>)</sup>		)
C4	ብሄርስብ	1. አማራ አሮሞ 3. 4. ሌላ ካ		

C5	የትምህርት ደረጃ?	1. ማንበብ ናመጸፍ አይችሎም
		2. የመጀመሪያ ደረጃ (1-8) 3.
		ሁለተኛ ደ <b>ረ</b> ጃ (9-12)

		4. h12รั	<sup>፡</sup> ክፍል በሳይ
C6	የኃብቻ ሁኔታ	1. ያላንባ ያንባ 3. የተፋቱ 4 የሞተበት	
C7	የስራ ሁኔታ?	1. የመንግስት ሰራተኛ 2. የግል መስሪያ ቤት ሰራተኛ 3. አርሶ አደር 4. የቤት አመቤት 5. ነ <i>ጋ</i> ደ 6. ሌላ ካለ ይጠቀስ )	
C8	እድሜያቸዉ ከ <i>ዓመት</i> በታች ህጻን አለወት ?	<b>አ</b> ወ	0 <i>ን</i> የለም
C9	አንተን /አንቺን ጨምሮ በቤት ዉስጥ ስንት ሆናችሁነ ዉ የምትኖሩት?	በቁጥር	
ክፍል	D: ሱስና ሱስነክ ስነም <b>ግ</b> ባር ባህር <i>ያትን</i> በተ <i>መ</i> ከተ		
1	. ትምባሆን በተመለከተ		
D1	በአሁት ጊዜ የትምባሆ ምርቶችንሲ <i>ጋራ፣ ጋያ</i> ይጠ	ቀ <b>ማ</b> ስ-?	1 አ <i>ዎን</i> <sup>0</sup> የለም. የለም ከሆነ ወደ <i>D5</i> ይሂዱ
D2	በአሁት ጊዜ የትምባሆ ምርቶችን, በየቀት ይጠቀማ	<del>ያ</del> ሉ-?	1 አ <i>ዎ</i> ን 0 የለም
D3	ሲያጨሱ ሰመጀመሪያ ጊዜእድ <i>ሜዎ</i> ስንት ነበር?		አመት 77 <b>አሳውቀዉም</b>
D4	በአ <i>ማ</i> ካኝ በሳም <i>ንት</i> በወር ምን ያህል ያጨሳሉ?		1. የፋብሪካሲ <i>ጋራ</i> 2. <i>ኃያ</i> 3.ሴላ ካለ ይጠቀስ <b>አሳውቀዉም 77</b>
D5	ባለፉት <sup>12</sup> ወራት የሲ <i>ጋ</i> ራ ምርቶችን ተጠቅመዋል?		<sup>1</sup> አ <i>ዎ</i> ን <sup>0</sup> የስም
D6	ባለፉት ወራት በየቀኑ ያጨሱ ነበር?		አ <i>ዎ</i> ን <sup>0</sup> የስም

D7	ጭስ አልባ <i>ትን</i> ባሆ በአፍንጫ ማማማ - ትንባሆማኘክ ይጠቀማሉ?	1 አ <i>ዎን</i> 0 የለም መልሰወ የለምክሆነ ወደ <b>ቁጥር</b> <i>D10</i> ይሂዱ ጥያቄ
D8	በየቀኑ <del>ሜ</del> ስ አልባ ትንባሆ በአፍንጫ ማማግ ትንባሆ ማኘክ ይጠቀማሉ?	አ <i>ዎ</i> ን <sup>0</sup> የለም
D9	በአማካኝ ስንት ጊዜ ጭስ አልባ ትንባሆ ይተቀማሉ ?	1. በአፍመሳብ 2. በአፍንጭመሳብ 3. ማኘክ 4. ሴሳካለ ይጠቀስ 77 አሳውቀዉም
D10	ባለፉት ወራት <del>ጭ</del> ስ አልባ ትንባሆ በአፍንጫ ማማግ ትንባሆ ማኘክ ተጠቅመዋል	1 አ <i>ዎን</i> <sup>0</sup> የ <b>ለ</b> ም
D11	ባለፉት <sup>30</sup> ቀናት ቤት ዉስጥ ያጨሱ ነበር?	1 አ <i>ዎን</i> 0 የለም
D12	ባስፉት ቀናት ሰወች ዝግ በሆነ ቦታ ስራቦታ ፣ ቢሮ ያጨሱ ነበር?	1.አ <i>ዎን</i> 0 የ <b>ለ</b> ም
2	አልኮል አወሳሰድን በተመለከተ	
D13	አልኮል ቢራ ጠሳ ከረፌ ጠ፯ አረቌ ወይን ጠጥታችሁ ታዉቃሳችሁ?	1 አ <i>ዎን</i> 0 የለምየ <b>ለምከሆነወደ <i>D23</i> ይሂ</b> ዱ
D14	ባለፉት ቼራት አልኮል ቢራ ጠሳ ክረፌ ወይን › ጠጂ› አረቃ ስፕራይት ጠጥታችሁ ታዉቃሳችሁ?	0 አ <i>ዎን</i> 0 የሰም
D15	ባለፉት ወራት፣አልኮልን ስትጥጠጡ በየስንት ጊዜዉ ይጠጡ ነበር ?	1. ቀንበቀን 2. 5-6 ቀንበሳምንት 3. 1-4 ቀንበሳምንት 4. 1-3 ቀንበወር 5. ከአንድጊዜበታችበወር
D16	ባለፉት <sup>ቆ</sup> ናት አልኮል ቢራ ጠሳ ከረፌ ጠጂ አሪቌ ወይን ) ጠጥታችሁ ታዉቃላችሁ??	1 አዎን <sup>0</sup> የ <b>ለ</b> ምከሆነ <b>ወደ</b> <i>D22</i> ይሂዱ
D17	ባለፉትቀናት አልኮል ሲጠቀሙ በአንድ ጊዜ ስንት	በቁጥር

ባለፉት ቆዩት አልኮል ሲጠቀሙ በዓማካኝ በአንድ ጊዜ ስንት

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በቁጥር \_\_\_\_

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ይጠጣሉ

ይጠጣሉ?

D18

D19	ባለፉትቀናት አልኮል ሲሲጠቀሙ ምን ያህል ክምግብ <i>ጋር</i> ተጠቀሙ?	1. አብዛኝውንጊዜከምግብ <i>ጋር 2.</i> አልፎአልፎከምግብ <i>ጋር 3.</i> በጥቂቱከምግብ <i>ጋር</i> 4. ከምግብ <i>ጋር</i> አልጠቀምም
D20	ባለፉትቀ <b>ኛ</b> ት አልኮል ሲጠቀሙ በአንድ ጊዜ ስንት ይጠጣሉ?	በቁጥር 77 አሳውቀውም
D21	ባለፉት ቀናት አልኮል ሲጠቀሙ በቤት ዉስጥ የተዘ <i>ጋ</i> ጁ ለምሳሌ ጠላ ጠጂ አረቄ ቦርደ ተጠቅመዋል?	1 ኣወን 0 የስም

3	ጫትን በተመለከተ	
D22	ጫት ቅመህ ሽ ታውቂያለህ ሽ	1 ኣወን
		0 የስም የስም <b>ከሆነወደ <i>E1</i> ይሂ</b> ዱ
D23	በአሁት ጊዜ ጫት ትቅመህ ሽ	1 ኣወን
		0 የለም የለም <b>ከሆነወደ <i>E1</i> ይሂ</b> ዱ
D24	ባለፉት ወራት፣ጫትን ስትጥጠቀሙ በየስንት ጊዜዉ ነበር?	1. ቀንበቀን
		2. <i>5-6</i> ቀ <b>ንበሳምንት</b>
		3. 3-4 <i>ቀን</i> በሳምንት
		4. 1-2 <b>ቀንበሳምን</b> ት
		5. 1-3 <b>ゆ</b> 3
		Λ <i></i> Ο <b>C</b>
		6. ከአንድጊዜበታችበወር
D25	በአ <i>ማ</i> ካኝ በ ወር <sup>/</sup> በሳም <i>ንት</i> ስ <i>ንት የታ</i> ሰረ <i>ጫት ይቅጣ</i> ሉ	የታሰረጫት ብዛት
		77 አሳውቀውም
D26	በአሁት ጊዜጫት እየቃምክ ሽ ታጨሳሰህ ሽ	1 አወን
		0 የስም
D27	በባለፉት ወራት ጊዜ ጫት እየቃምክ ሽ ታጨሳለህ?ሽ	1 አወን <sup>0.</sup>
	12	የስም
D28	በአሁት ጊዜ ጫት ከቃምክ ሽ በኋላ አልኮል ትጠጣላችሁ?	1 አወን
		0 የ <b>ሰም</b>

ክፍልE፡			
ከዚህ	ቀጥዬ የተለያዩ የአካል እንቀሰቃሴ በጣ	ነካሄዴ የሚያሳልፉ <b>አቸ</b> ዉን '	ኒዜያት በተ <b></b> ጣለከተ እጠይቅዎታለሁ።
ከስራ ;	<b></b> ፖር የተያያዙ አካላዊ <b>እንቅስቃሴ</b> ወች		
<b>E</b> 1	ባለፉት 7 ቀናት ብርቱ ጉልበት የሚይ		ቢያንስ ለ10 ደቂቃ ስለሚያደርጋ
	ቁ አካላዊ እንቅስቃሴዎችን ለስንት ቀ	77 አሳውቅም	ቸው አካሳዊ እንቅስቃሴዎች እንዲ
	ናት አደረጉ?		ያስቡ አማዝ
			ምሳሹ ዜሮጠጠ ወይም አሳውቅም
			ከሆነ ወደ <u>"ጥያቄ</u> E3 <u>" እለፍ/ፊ</u>
<b>E2</b>	በነዚያ ቀናት ብርቱ ጉልበት የሚ∩ይ	በቀን ሰዓት	አሳውቅም ከሆነ ወደ <u>"ዋያቄ</u> E3 <u>"</u>
	ቁ አካሳዊ እንቅስቃሴዎችን በማድረግ		<u> </u>
	ምንያህል ጊዜ ያሳልፋሉ?	በቀንደቂቃ	
		አሳውቅም77	
E2-1	ባለፉት 7 ቀናት ብርቱ ጉልበት	ሰዓት በሳምንት	
	የሚጠይቁ አካሳዊ እንቅስቃሴ ዎችን	ደቂቃ በሳምንት	
	በማድረግ በድምሩ ምን ያህል ጊ	77 አሳውቅም	
አቤ3	ዜ አሳለፉ? ባለፉት 7 ቀናት ውስጥ በነበሩት የሥራ	ቀናት <i>መ</i> <b>ለ</b> ነኛ ጌልበት	የማ ሉ የ ል ኔክለው እንቅስ ቀለ ወችን
	· ስሳሳለፉት ጊዜ አስቡ. ቀላል ሽክም ,ሞ		
<b>E3</b>	ባለፉት 7 ቀናት መጠነኛ ጉልበት የ		
	ሚጠይቁ አካሳዊ እንቅስቃሴዎችን ለ		ወደ ተያቄ E 5"  እለፍ/ፌ]
	ስንት ቀናት አደረጉ?	77 አሳውቅም	-
<b>E4</b>	በንዚያ ቀናት <i>መ</i> ጠነኛ ጉልበት የሚ	በቀን ሰዓት	ምሳሹ ዜሮወይም አሳውቅም ከሆ
	ይጠቁ አካሳዊ እንቅስቃሴዎችን በማ		ን ወደ <u>"ጥያቄ</u> E5 <u>" እለፍ/ፊ</u>
	ድረግ ምንያህል ጊዜ ያሳልፋሉ?	በቀንደቂቃ	
		77 አሳውቅም	
E4-1	ባለፉት 7 ቀናት <i>መ</i> ጠነኛ ጉልበት	ሰ <b>ዓት</b> በሳምንት	
	የሚጠይቁ አካሳዊ እንቅስቃሰ <sub>5</sub> ዎችን	ደቂቃ በሳምንት	
	በማድረግ በድምሩ ምን ያህል	77 አሳውቅም	
L.10 (	ጊዜ አሳለፉ?	u 130	L. Cour
ለሁን ፣	ነለፉት 7 ቀናት በእግር ጉዞ ስሳሳለፉት 1 በመዝናኛ፣ በስፖርት ወይም በዕረባ	•	
	וויט וויט ניין איניין שניין וויטן שניין	ቶ <i>ገ</i>	4.1.1.2 W. 197.197
E5	ባለፉት 7 ቀናት በአንድ ጊዜ ቢያንስ	በሳምንት ለ ቀናት	ምላሹ ዜሮወይም አላውቅም ከ
	የ10 ደቂቃ የአግር ጉዞ ለስንት ቀና		ሆን ወደ <u>"ዯያቄ</u> E7 <u>" እለፍ/ፊ</u>
	ት አደረጉ?	77 አሳውቅም	

E6	ከንዚያ ቀናት በአንዱ የእግር ጉዞ በጣ ድረግ በአማካኝ ስላሳለፉት ጊዜ ለጣ ወቅ ይፈለጋል። ጊዜ ከቀን ቀን በጣም የሚለያይ በመ ሆኑ ሳቢያ ለመመለስ ካልቻሉ 'ባለፉ ት 7 ቀናት የእግር ጉዞበማድረግ በድ ምሩ ምን ያህል ጊዜ አሳለፉ?"	ሰዓት በሳምንት ደቂቃ በሳምንት 77 አሳውቅም	ምላሹ ዜሮወይም አላውቅም ከ ሆነ ወደ <u></u>
E6-1	ባለፉት 7 ቀናት የእግር ጉዞ በጣድ ረግ የሚይቁ አካሳዊ እንቅስቃሴዎች ን በጣድረግ በድምሩ ምን ያህል ጊዜ አሳለፉ?	ሰዓት በሳምንት ደቂቃ በሳምንት 77 <b>አሳ</b> ውቅም	
በቤት፣	በለፉት 7 ቀናት ውስ <b>ተ በነበ</b> ሩት የሥራ ቁ በትምህርት፣በዕረፍትያሳለፉትንጊዜያጠቃ ወቀ፣ በመተ <b>ኛ</b> ት ወይም ቴሌቪዥን በወ	ያልላል።ይ <mark>ህበፅሁፍ ሥራ፣</mark>	ደኞችን በ <i>ሙ</i> ጎብኘት ፣ በማንበብ ፣
E7	ባለፉት 7 ቀናት ከነበሩት የሥራ ቀና ት በአንዱ ምን ያህል ጊዜ ቁጭ በማ ለት አሳለፉ?	ሰዓት በሳምንት ደቂቃ በሳምንት 77 <b>አሳውቅም</b>	ምላሹ ዜሮወይም አላውቅም ከ ሆነ ወደ "ጥያቀ G እስፍ/ፌ
E8	ባለልው ረቡዕ ቁ <del>ጭ</del> ብለው በድምሩም ን ያህል ጊዜ አሳለፉ?	ረቡዕ ዕለት ለ _ ሰዓት ረቡዕ ዕለት ለ ደቂ <i>ቃ</i> 77 አሳውቅም	

ክፍል F: አመ*ጋገብን* በተመለከተ፡ባለፉት <sup>12</sup> ወራት ወደ ኋላ መለስ ብለው ሲያስቡ በየቀኮ <sup>1</sup> በየሳምንቱ ወይም በየወሩ ምንያህል በተደ*ጋጋሚ እያንዳንዳ*ቸውን ምግቦች እንደተመገቡ ንገሩን

ምግብ ዝርዝር	ድግግሞሽ	ምግብዝርዝር	ድግግሞሽ
	ምንም አልተመገቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
	h 2-3 ጊዜ በወር		h 2-3 ጊዜ በወር
F 1.	ከ1-2 ጊዜ በሳምንት	F 2. በቆሎ	ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳምንት		ከ3-4 ጊዜ በሳምንት
	5-6 <i>ጊ</i> ዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		ከ2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		h 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በላይ በቀን		6 ጊዜ እና ከዚያ በላይ በቀን

	ምንም አልተመገቡም አንድ ጊዜና ከዚያ በታች በወር		ምንም አልተመገቡም አንድ ጊዜና ከዚያ በታች በወር
F 3. 7-11ስ	h 2-3 ጊዜ በወር	F 4. ስንደ ና ዱቄት	ከ 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳምንት		ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን

		1	
	ከ2-3 ጊዜ በቀን		h2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በላይ በቀን		6 ጊዜ እና ከዚያ በላይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
F5. <b>ማ</b> ሽላ/ዘ <i>ን,ጋዳ</i>	h 2-3 ጊዜ በወር	F 6. ሩዝ	h 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	h3-4 ጊዜ በሳምንት	-	ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		ከ2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በላይ በቀን		6 ጊዜ እና ከዚያ በላይ በቀን
F 7. የበሬ ስ <i>ጋ</i>	ምንም አልተመገቡም	F 8. የዶሮ ስ <i>ጋ</i>	ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
	h 2-3 ጊዜ በወር	-	ከ 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳምንት	-	ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		ከ2-3 ጊዜ በቀን

	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በሳይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
F 9. አሳ	አንድ ጊዜና ከዚያ በታች በወር	F 10. <i>ጉ</i> በት	አንድ ጊዜና ከዚያ በታች በወር
	ከ 2-3 ጊዜ በወር		h 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳምንት		ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		ከ2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ ሕና ከዚያ በላይ በቀን		6 ጊዜ እና ከዚያ በሳይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
F11.ፍየል/ <b>ግ</b> ልንል	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
	h 2-3 ጊዜ በወ <b>ር</b>	F12 የታሸን ስ <i>ጋ</i>	h 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	h3-4 ጊዜ በሳምንት		ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	h2-3 ጊዜ በቀን		h2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በሳይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
F 13. ወተት	አንድ ጊዜና ከዚያ በታች በወር	F 14. አይብ	አንድ ጊዜና ከዚያ በታች በወር
	h 2-3 ጊዜ በወር		h 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳም <i>ንት</i>

	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		h2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ ሕና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በላይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
F 15. ቅቤ	አንድ ጊዜና ከዚያ በታች በወር	F 16.	አንድ ጊዜና ከዚያ በታች በወር
	h 2-3 ጊዜ በወር		ከ 2-3 ጊዜ በወር
	ከ1-2 <i>ጊ</i> ዜ በሳም <i>ንት</i>		ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳምንት		ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		ከ2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን

	6 ጊዜ ሕና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በላይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር	F 18. ዘይት	አንድ ጊዜና ከዚያ በታች በወር
F 17. ስዉዝ	h 2-3 ጊዜ በወር	1 10. 11,57	h 2-3 ጊዜ በወ <b>ር</b>
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳም <i>ንት</i>		ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		ከ2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ ሕና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በሳይ በቀን

F 19.ባቄላ, አተር, ምስር, አኩሪ አተር	ች አንድ ጊዜና ከዚያ በታች በወር
h 2-3 2H በወር	
	ከ 2-3 ጊዜ በወር
ከ1-2 ጊዜ በሳምንት	ከ1-2 ጊዜ በሳምንት
ከ3-4 ጊዜ በሳምን <i>ት</i>	ከ3-4 ጊዜ በሳምንት
5-6 ጊዜ በሳምንት	5-6 ጊዜ በሳምንት
አንድ ጊዜ በቀን	አንድ ጊዜ በቀን
ከ2-3 ጊዜ በቀን	ከ2-3 ጊዜ በቀን
ከ 4-5 ጊዜ በቀን	ከ 4-5 ጊዜ በቀን
6 ጊዜ እና ከዚያ በሳይ በቀን	6 ጊዜ እና ከዚያ በላይ በቀን
F 21.ድንች ምንም አልተመገቡም F 22. ክሮት	ምንም አልተመገቡም
አንድ ጊዜና ከዚያ በታች በወር	አንድ ጊዜና ከዚያ በታች በወር
ከ 2-3 ጊዜ በወር	ከ 2-3 ጊዜ በወር
ከ1-2 ጊዜ በሳምንት	ከ1-2 ጊዜ በሳምንት
ከ3-4 ጊዜ በሳምንት	ከ3-4 ጊዜ በሳምንት
5-6 ጊዜ በሳምንት	5-6 ጊዜ በሳምንት
አንድ ጊዜ በቀን	አንድ ጊዜ በቀን
ከ2-3 ጊዜ በቀን	h2-3 ጊዜ በቀን
ከ 4-5 ጊዜ በቀን	ከ 4-5 ጊዜ በቀን
6 ጊዜ እና ከዚያ በሳይ በቀን	6 ጊዜ እና ከዚያ በላይ በቀን

	<i>ምንም አልተመገ</i> ቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
F 23. ቲ <b>ጣ</b> ቲም	h 2-3 ጊዜ በወር	F 24. አበባ <i>ጎመን</i>	ከ 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳምንት		ከ3-4 ጊዜ በሳምንት
	5-6 <b>ጊዜ በሳም</b> ንት		5-6 <b>ጊዜ በሳም</b> ንት

	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	h2-3 ጊዜ በቀን		h2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን	_	ከ 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በሳይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
F25.ቅጠል፣ አረንጓደ አትክልት	h 2-3 ጊዜ በወር	F 26. አቮካዶ	h 2-3 ጊዜ በወር
na / i a ni ni ni	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳም <i>ንት</i>	_	ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን	-	አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን	-	ከ2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን	-	ከ 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በሳይ በቀን	-	6 ጊዜ ሕና ከዚያ በሳይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
F 27. ፓፓያ	ከ 2-3 ጊዜ በወር	F 28. <i>ሙዝ</i>	h 2-3 ጊዜ በወር
	ከ1-2 <i>ጊ</i> ዜ በሳም <i>ንት</i>		ከ1-2 ጊዜ በሳምንት
	ከ3-4 ጊዜ በሳም <i>ንት</i>		ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		ከ2-3 ጊዜ በቀን
	h 4-5 ጊዜ በቀን	1	h 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በሳይ በቀን	1	6 ጊዜ እና ከዚያ በሳይ በቀን
	ምንም አልተመንቡም		ምንም አልተመገቡም

	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
F 29. አናናስ	h 2-3 ጊዜ በወር	F 30. ብርቱካን	h 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት

	ከ3-4 ጊዜ በሳምንት		ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	h2-3 ጊዜ በቀን		h2-3 ጊዜ በቀን
	h 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ ሕና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በሳይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
F31.ቀይ ሽንኩረት	h 2-3 ጊዜ በወር	F32.ነ <del>ሜ</del> ሽንኩረት	h 2-3 ጊዜ በወር
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	h3-4 ጊዜ በሳምንት		h3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	h2-3 ጊዜ በቀን		ከ2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በሳይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር
F 33. ዝንጂብል	h 2-3 ጊዜ በወር	F 34. <i>ቃሪያ</i>	h 2-3 ጊዜ በወ <b>ር</b>
	ከ1-2 ጊዜ በሳምንት		ከ1-2 ጊዜ በሳምንት
	h3-4 ጊዜ በሳምንት		ከ3-4 ጊዜ በሳምንት
	5-6 ጊዜ በሳምንት		5-6 ጊዜ በሳምንት
	አንድ ጊዜ በቀን		አንድ ጊዜ በቀን
	ከ2-3 ጊዜ በቀን		h2-3 ጊዜ በቀን
	ከ 4-5 ጊዜ በቀን		ከ 4-5 ጊዜ በቀን
	6 ጊዜ እና ከዚያ በሳይ በቀን		6 ጊዜ እና ከዚያ በሳይ በቀን
	ምንም አልተመገቡም		ምንም አልተመገቡም
	አንድ ጊዜና ከዚያ በታች በወር		አንድ ጊዜና ከዚያ በታች በወር

F 35. ቡና	F36.የአካባቢ <i>መ</i> ጠዣ	ከ 2-3 ጊዜ በወር ከ1-2 ጊዜ በሳም ከ3-4 ጊዜ በሳም 5-6 ጊዜ በሳምን አንድ ጊዜ በቀን ከ2-3 ጊዜ በቀን ከ 4-5 ጊዜ በቀን	31· 31· 1·
	F 38. ጨወ.	ምንም አልተመን አንድ ጊዜና ከዚ, ከ 2-3 ጊዜ በወር ከ1-2 ጊዜ በሳም ከ3-4 ጊዜ በሳም 5-6 ጊዜ በሳምን አንድ ጊዜ በቀን ከ2-3 ጊዜ በቀን ከ 4-5 ጊዜ በቀን 6 ጊዜ እና ከዚ	ያ በታች በወር ፡ ፡ ፡ ፡ ፡ ት ፡
F 39. ከቤት ዉጭ መመንብ	F 40. ባለፉት 12 መረ ሳይመንቡ ይዘሉ/ያሳልፉ F40a. መልሰወ አወን የቱንያዘልሳሉ? F41. በቤትዎ ውስጥ በመደበኛነት ብዙጊዜ ም ወይም ቅባት ነው ጥቅ የሚያዉሉት	ነበር? ከሆነ ስምግብነት ን አይነት ዘይት	1 አወን 0 የለም 1. ቁርስ 2. ምሳ 3 አራት የአትክልት ዘይት 2.የረጋ ዘይት ቅቤ 4.ሽኖ ለጋ

	5. ሴላ ካለ ይ <b>ግ</b> ለጹ <i>)</i>
1. ቀርስቀናት 2. ምሳቀናት 3 አራትቀናት	

በመጀመሪ	ያ ስለ ታካሚው የደም ግፊትን ታሪክ ጥያቄዎች እጠይቅዎታለሁ	
J1	በአሁን ጊዜ በሀኪምዎ ወይም በሴሳ የጤና ባለሙያ የታዘዘውን የደም ግፊት ማንኛውንም ሕክምና ምክር ይቀበሳሉ?	1 አ <i>ዎን</i> <i>0</i> የሰም
J2	ሕርስዎ ከደም ግፊት <i>ጋር</i> የተ <i>ያያ</i> ዘ የቤተሰብ ታሪክ አለዎት? ወሳጅ፣ወንድም ደም ግፊት ያለበት አለ?	አ <i>ዎን</i> <i>0</i> የለም
የስኳር	በሽታን በተመስከተ	
J3	በአሁት ወቅት በሀኪምዎ ወይም በሴሳ የጤና ባለሙያ የታዘዘውን ማንኛውንም የስኳር ህክምና/ ምክር ይቀበሳሱ?	1 አ <i>ዎን</i> <i>0</i> የሰም
J4	ሕርስዎ ክስኳር ህመም <i>ጋ</i> ር የተያያዘ የቤተሰብ <b>ቃ</b> ሳጅ <i>▶ሪክ አለዎት?</i>	1 <i>አዎን</i> <i>0</i> የ <b>ሰ</b> ም

ክፍል K	: የአካል ልኬትን በተመለከተ፣ ቁመት: ክተ	በደ <i>ት</i> :የወንብ <b>ና</b> የዳሌ ልኬት
K1	ቁ <b>መ</b> ት	(ሳሜ) ለ ልኬት 1
		(ሳሜ) ለ ልኬት 2
K2	ክብደት	ኪማ ለ ልኬት 1
		ኪማ ለልኬት 2
K3	የወንብ ልኬት	<u>በሣንቲ ሜትር (ሣሜ)</u> ልኬት <sup>1</sup>
		<u>በሣን</u> ቲ <i>ሜትር (ሣሜ</i> ) ልኬት <sup>2</sup>
K4	የዳሌ ልኬት	<u>በሣንቲ ሜትር (ሣሜ)</u> ልኬት <sup>1</sup>
		<u>በሣን</u> ቲ <i>ሜትር (ሣሜ</i> ) ልኬት <sup>2</sup>
ክፍል L	የደም ግፊትን በተመለከተ	
L1	የደም ግፊት ልኬት 1	ሲስቶ <b>ሊክ <i>ሚሜሜር</i>ኩሪ</b> <u>)</u>
		ዲያስቶሊክ <i>(ሚሜሜር</i> ኩሪ <u>)</u>
L1-1	የደም ግፊት ልኬት <i>2</i>	ሲስቶ <b>ሊ</b> ክ <i>ሚሜሜር</i> ኩሪ <u>)</u>
		ዲያስቶሊክ <i>ሚሜሜር</i> ኩሪ

# ክፍል M: ባዩ ሜዲካል ልኬትን በተመለከተ፡

መጀመ	ሪያ የደም ስኳር መጠን መ <b>ሰ</b> ካት	
M1	ባለፉት <i>ჩዓታት</i> ዉስጥምግብፈሳሽነ <i>ገር</i> ወስደዋል ?	1 አዏን <i>0</i> የሰም
M2	የደም ናሙና የተወሰደበት ስአት	ሰአት
M3	ምግብ ከመበሳተዎ በፊት የደም ስኳር መጠን	ሚ ሞል/ ሲትር ሚ ግ/ዲሲ ሲትር ፍቃደኝ አይደለሁም
M4	በአሁት ወቅት የደም ስኳርዎ በመጨመሩ ምክንያት በሀኪምዎ ወይም በሌሎች ጤና ባለሞያወች ኢንሱሊን መርፌ ወይም ክኒን ታዝወሎት ይወስዳሉ?	አዏን <i>0</i> የሰም
የደም ስብ	በ <i>መ</i> ጠን ልኬ <i>ት</i> :	
M5	ቶታል ኮሴስትሮል	ሚሞል/ ሲትር ሚግ/ዲሲ ሲትር ፍቃደኝ አይደስሁም
M6	በአለፉት ሁለት ሳምንታት ውስጥ ስብ መጠንዎ በመጨመሩ ምክንያት በሀኪም ወይም በሌሎወች ጤና ባለሞያወች ክኒን ወይም መርፌ ታዝወሎት ታክመዉ ያዉቃሉ?	1 አዏን 0 የሰም
M7	ትራይ ማሲሰራይድ	
M8	ኤች ዲ ኤል <i>ኮ</i> ሌስትሮል	

ሕና*መ*ሰማናስን!