Prevalence, antimicrobial susceptibility pattern, and associated factors of *Salmonella* and *Shigella* among food handlers in Adigrat University student’s cafeteria, Northern Ethiopia, 2018

**CURRENT STATUS:** UNDER REVIEW

*Tropical Diseases, Travel Medicine and Vaccines*  •  BMC

Haftom Legese  
Adigrat university

✉ legesehaftom2@gmail.com  
*Corresponding Author*  
ORCID: https://orcid.org/0000-0002-6280-1116

Tsega Kahi
Adigrat University College of Health Sciences

Aderajew Gebrewahd
Adigrat University College of Health Sciences

Brhane Berhe
Adigrat University College of Health Sciences

Berhane Fseha
Adigrat University College of Health Sciences

Senait Tadesse
Bahir Dar University College of Medical and Health Sciences

Guesh Gebremariam
Adigrat University College of Health Sciences

Hadush Negash
Adigrat University College of Health Sciences

Fitsum Mardu
Adigrat University college of Medicine and health Sciences

Kebede Tesfay
Adigrat University College of Health Sciences
DOI: 10.21203/rs.2.22938/v2

SUBJECT AREAS
Infectious Diseases

KEYWORDS
Antimicrobial susceptibility, Food handlers, Salmonella, shigella, Ethiopia
Abstract
Background: Food handlers play a significant role in the transmission of foodborne infection. Salmonella and Shigella are the most common foodborne pathogens and their infections are a major public health problem of the globe. Thus, this study was aimed to determine the prevalence, antimicrobial susceptibility patterns, and associated factors of Salmonella and Shigella among food handlers.

Methodology: A cross-sectional study was conducted from March to August 2018 at Adigrat University student cafeteria, Northern Ethiopia. Data on socio-demographic and associated factors were collected using a structured questionnaire. Fresh stool samples were collected from 301 food handlers and transported to Adigrat University Microbiology Laboratory. Bacterial isolation and antimicrobial susceptibility test were performed using standard bacteriological methods. Data analysis was performed using SPSS version 22 and P < 0.05 with a corresponding 95% confidence interval was considered statistically significant.

Results: A total of 301 food handlers were included in this study. The majority of study participants were females 265 (88.0 %). About 22 (7.3%) and 11 (3.7%) of food handlers were found to be positive for Salmonella and Shigella respectively. Hand washing after using a bathroom with water only, hand washing after using the bathroom, hand washing after touching dirty materials, hand washing before food handling and fingernails status were significant associated factors identified. None of the Salmonella and Shigella isolates was sensitive to ampicillin. On the other hand, low resistance was found for chloramphenicol, ceftriaxone, and ciprofloxacin.

Conclusion: The present study revealed that the prevalence of Salmonella and Shigella among food handlers found to be 22 (7.3%) and 11 (3.7%) respectively. Such colonized food handlers can contaminate food, drinks and could serve as a source of infection to consumers via the food chain. This indicates that the need for strengthened infection control measures to prevent Salmonella and Shigella transmission in the students’ cafeteria.

Background
Foodborne diseases are a major public health problem in the globe. The severity is higher among
developing countries due to low hygienic food handling practices, lack of environmental sanitation and poor access to safe drinking water [1]. In developing countries, approximately 70% of cases of diarrheal diseases are associated with the consumption of contaminated food [2].

*Salmonella* remains a major cause of foodborne infection in humans [3], which leads to approximately, 93 million infections every year [4, 5]. World Health Organization (WHO) estimates that there are around 16 million new cases and 600,000 deaths due to typhoid fever each year worldwide [6]. It causes bacterial bloodstream infections with a fatality rate of 20-25% [7]. The widespread nature of salmonellosis increases antibiotic resistance which in turn increases the treatment cost, hospitalization, morbidity, and mortality [8].

These bacteria are transmitted directly and indirectly through contaminated objects such as food, water, nails, and fingers, this indicates those microorganisms can be spread by faecal-oral human-to-human transmission [9, 10]. Bacteria are transmitted directly and indirectly through food, water, and fingernails compared to other parts of the hand, fingernails harbour the most microorganisms and difficult to clean easily.

*Shigella* continues to play a major role in etiology for inflammatory diarrhoea, and dysentery in food handlers [11]. The annual incidence of *Shigella* is estimated to be 164.7 million people, with 69% of all deaths attributable to shigellosis worldwide [12, 13]. The highest prevalence of shigellosis is observed in tropical and subtropical parts of the world [14].

*Salmonella* and *Shigella* are a significant cause of severe post diarrheal complications such as reactive arthritis, sepsis, reiter syndrome, myocarditis, inflammatory bowel diseases, irritable bowel syndrome, and peritonitis [15, 8, 16]. The emergence of antimicrobial-resistant *Salmonella* and *Shigella* becomes a significant threat to deliver reliable therapies [17, 18].

In Ethiopia, it is difficult to estimate the severity of salmonellosis and shigellosis as well as their antibiotic resistance due to limited scope of studies, lack of coordinated epidemiological surveillance system, poor reporting system and limited availability of culture facilities [19].

Determining the prevalence and antimicrobial susceptibility pattern of *Salmonella* and *Shigella* is very important for the proper selection of antimicrobial agents to control the spread of infection. However,
in the study area, there was a scarcity of data on the carriage of salmonellosis and shigellosis among food handlers. Therefore, the aim of this study is designed to assess the prevalence, antimicrobial susceptibility patterns, and associated factors of *Salmonella* and *Shigella* among food handlers in Adigrat University, Tigrai, Northern Ethiopia.

Materials And Methods

**Study Design, Area and Period**

A cross-sectional study was conducted among food handlers who were participated in food preparation, dispatch and store of Adigrat University student’s cafeteria from March to July 2018. The annual rainfall ranges from 400-600mm and the minimum and maximum temperature range from 6-21.8°C. Currently, the University enrolled more than 15,000 students who are getting dining services in the student cafeteria. There are six cafeterias and a total of 700 food handlers are working in the student’s cafeteria (Adigrat University human resource management and registrar office).

**Sample size determination and sampling technique**

**Sample size determination**

The sample size was determined by using a single population proportion formula.

\[
 n = \left( \frac{Z_{\alpha/2}}{d} \right)^2 P (1-P)
\]

The sample size was determined based on the prevalence of *Salmonella* among university food handlers done by Mama and Alemu at Arba Minch University, South Ethiopia (6.9%) [14] Then with a margin of error (5%), (\(d=0.03\)) and 95% level of confidence (\(z=1.96\)), the sample size was calculated as follows:

\[
 n = \left( \frac{1.96}{0.03} \right)^2 \times 0.069(0.931) = 274, \text{ with } 10 \% \text{ non response rate } = 301
\]

Therefore, a total of 301 food handlers were included in the study from all cafeteria of the university. A simple random sampling technique was employed. The lottery method was used to select the study subjects after a complete list of food handlers was obtained from a roster of cafeteria office Adigrat.
Eligibility criteria

Inclusion criteria

Food handlers working in Adigrat University student’s cafeterias were included in the study.

Exclusion criteria

Food handlers who have taken antibiotics within one week, antihelminthics, and those with clinical signs of typhoid fever were excluded from the study.

Data collection and Sample processing

Socio-demographic and Specimen Collection, Handling and Transportation

A structured questionnaire was used to collect the data regarding socio-demographic and associated factors. Questionnaires were checked for accuracy and completeness. After proper instruction, about 2 g of fresh stool specimens were collected from food handlers with a labelled wide-mouthed plastic container and a clean wooden applicator stick. Specimens were immediately transported to the laboratory using icebox.

Isolation and identification

The stool specimen was collected and transported to Adigrat University Medical Microbiology laboratory within one hour of collection. Stool specimens were immediately inoculated in Selenite F enrichment broth and incubated at 37°C for 24 hours, and then subculture onto selective media of xylose-lysine desoxycholate agar (XLD) and Hektoen enteric medium agar incubate at 37°C for 18-24 hours. The isolated colonies were differentiated and identified based on gram stain, colonial morphology and pigmentation, hemolysis on blood agar, catalase test, oxidase test, carbohydrate fermentation, H₂S production, motility, indole formation and urease production, citrate utilization and incubated for 24 to 48 hours at 37°C. Then colonies producing an alkaline slant with acid butt and hydrogen sulfide production on Triple Sugar Iron Agar, positive for lysine, negative for urea hydrolysis, negative for indole test, positive for citrate utilization and motility test were considered to be Salmonella. Colonies which were urease negative, indol positive/negative, in Triple Sugar Iron agar produce a pink-red slope and yellow butt with no blackening, Lysine decarboxylase negative and
citrate negative is identified as *Shigella* species. Finally, all of the confirmed *Salmonella* and *Shigella* isolates were examined for antimicrobial susceptibility.

**Antimicrobial susceptibility tests**

Antimicrobial susceptibility testing was performed using the modified Kirby- Bauer disc diffusion method according to Clinical and Laboratory Standards Institute (CLSI) guidelines, 2016 [20]. Using a sterile wire loop, 3-5 well-isolated colonies of the test organism was emulsified into a tube of 3-4 ml sterile physiological saline to get bacterial inoculums equivalent to 0.5 McFarland turbidity standards. Then the standardized suspension (test organisms) were uniformly swabbed within 15 minutes using a sterile cotton swab into Muller-Hinton agar and allowed to dry. After that, the antibiotic discs were placed manually on the medium and incubated at 37°C for about 18 hours and the zones of inhibition were measured using a calliper. The interpretation of the results was made based on the CLSI criteria as sensitive, intermediate and resistant [20]. The following antimicrobials are prioritized by considering local prescription; gentamicin (10 μg), ampicillin (30 μg), amoxicillin (30 μg), ciprofloxacin (5 μg), clarithromycin (30 μg), chloramphenicol (30 μg), cotrimoxazole (25 μg), amoxicillin-clavulanic acid (30 μg), and ceftriaxone (30 μg) [20].

**Data Quality Assurance**

Data quality was ensured at various activities of the study by following a prepared standard operating procedure (SOP). Questionnaires were prepared in a clear and precise way and translated into the local language and back-translated to English to ensure the consistency of the questionnaires. The pretest was done on 5% of food handlers and modifications were made accordingly. To ensure general safety; universal bio-safety precautions were followed. American Type Culture Collection (ATCC) strains *P. aeruginosa* (ATCC 27853), *E. coli* (ATCC-25922) were used as control strains for the culture and antimicrobial susceptibility testing.

**Statistical Analysis**

After collection of socio-demographic characteristics, associated factors and laboratory data using a structured questionnaire and laboratory report format, data were edited, cleaned, entered and analyzed using statistical package for social science (SPSS) version 22. Descriptive statistics,
Bivariate, and multivariate logistic regression were performed. Bivariate logistic regression was employed to look association between the outcome variable and each independent variable. A binary logistic regression analysis was used to calculate the odds ratios (OR); crude odds ratio (COR) and adjusted odds ratio (AOR) to ascertain the degree of association between dependent and independent variables. In this study, multi-collinearity among independent variables was detected using the standard errors for regression coefficients. Finally, variables with p-value < 5% with a corresponding 95% confidence interval (CI) were considered as statistically significant.

Results

Socio-demographic characteristics

A total of 301 food handlers were included in the study. Out of the total respondents, 265 (88.0 %) were females. The age of study participants ranged from 19-38 years (23.51 ± 3.186 years). The majority of 241(80.1%) of the participants were between the ages of 21 and 30 years. One hundred fifty-six (51.8%) were enrolled in secondary school with an average of 3.74 years of work experience in the cafeteria. Out of the total study participants, 37(12.3%) were certified for training in food handling and 265(88.0%) had previously undergone a medical checkup stool microscopy examination (Table 1).

Prevalence and associated factors of Salmonella and shigella carriers

The prevalence of Salmonella and Shigella in this study was 22 (7.3%) and 11 (3.7%) respectively. In the current study, 13 independent variables were considered during the analysis of associated factors for Salmonella and Shigella carriers (Table 2). All variables with a significance value in the bivariate analysis were entered into multivariate logistic regression analysis. Accordingly in the multivariate analysis being hand washing after using the bathroom with water only (AOR=23.239, 95%CI:2.125-254.17, P<0.01), hand washing after using the bathroom (AOR=2.25, 95%CI: 5.11-77.34, P<0.001), hands washing after touching dirty materials (AOR=37.19, 95%CI: 5.66-244.45, P<0.001), hand washing before food handling (AOR=33.1, 95%CI: 4.958-220.52, P<0.001) and fingernail status (AOR=13.97, 95%CI: 3.404-57.362, P<0.001) were significant factors associated with Salmonella and Shigella carriers (Table 3).
Antimicrobial Susceptibility Patterns of *Salmonella* and *Shigella* isolates

Antimicrobial susceptibility patterns were performed for isolates against 9 antimicrobial agents. Of the isolates 22 (100%) *Salmonella* and 11(100%) *Shigella* were resistance to ampicillin 22(100%) and 11(100%) followed by gentamicin 22(100%) and 10(90.9%), amoxicillin 21(95.5%) and 11(100%) clarithromycin 9(40.9%) and 5(45.5%) respectively. However, all isolates were susceptible to ciprofloxacin. Resistance was observed for ceftriaxone 0(0.00%) and 1(9.1%), chloramphenicol 0(0.00%) and 2(18.2%) and cotrimoxazole 2(9.1%) and 1(9.1%) for *Salmonella* and *Shigella* respectively. None of the isolates showed intermediate resistance (Table 4). Multidrug resistance in this study is defined as resistance to at least three classes of antimicrobial agents and out of the thirty-three isolates 12 (54.54%) *Salmonella* and 10(90.9%) *Shigella* species were multi-drug resistant isolates (Table 5).

Discussion

In this study, the prevalence of *Salmonella* among food handlers was 22 (7.3%). This was similar to studies carried out in Southern Ethiopia, Arba Minch University (6.9%) [14] and Nigeria, Abeokuta (5.5%) [21]. However, it was higher than the studies reported from Ethiopia, Addis Ababa (3.5%) [22], Bahir Dar (1.6%) [23] and Gondar (3.1%) [24]. On the other hand, our result was lower than the studies reported from Ethiopia, Addis Ababa (10.5%) [25] and Nigeria (42.3%) [26]. The variation might be attributed to poor personal and environmental hygiene differences among the study areas. The rate of *Shigella* (3.7%) in our study is consistent with studies done in Southern Ethiopia, Arba Minch University (3%) [14], Ethiopia, Addis Ababa (4.5%) [25], and Gondar (3.1%) [26]. However, our finding was lower than a study conducted in Nigeria (15.5%) [27]. These might be due to the differences in hygiene practices of the food handlers.

In the present study, the practice of handwashing after using the bathroom among food handlers was significantly associated with *Salmonella* and *Shigella* carriers. Food handlers who hadn’t washed their hands after using the bathroom were more likely to be colonized with *Salmonella* and *Shigella* compared to those who were washed with water and soap after using the bathroom. This finding was similar to a study conducted in Ethiopia, Mekelle [28], Gondar [29] and Bahir Dar [30]. The acquisition
of *Salmonella* and *Shigella* is due to poor sanitary conditions, poor toilet facilities and availability of facilities used for handwashing practice.

Our finding also revealed that there is a statistically significant difference in handwashing after touching dirty materials among food handlers with *Salmonella* and *Shigella* carriers. Food handlers not washing their hands after touching dirty materials are twenty-eight fold more likely to be colonized with *Salmonella* and *Shigella* than those who washing with water and soap after touching dirty materials. This finding is consistent with a study conducted in Ethiopia, Bahir Dar [23]. This might be due to the absence of handwashing facilities within proximity of the food handler's workplace.

Our study showed that food handlers who were washed their hand with soap and water before touching food were less likely to be colonized with *Salmonella* and *Shigella* than food handlers who were not washing their hand with soap and water before food preparation. This is in line with the finding of a similar study reported from Ethiopia, Yebu Town [31]. In the majority of food handlers, hand washing before handling food was practiced. However, a very large proportion (42.8%) were washed their hands only by water. There are food handlers who apply some hygiene practice, though many of them do not use soap nor do they appreciate or understand the need handwashing [32].

Furthermore, in the current study, untrimmed fingernail was significantly associated with *Salmonella* and *Shigella* colonization among food handlers. This study is similar to studies conducted in Ethiopia, Yebu Town [31] and Arba Minch [14]. This result might due to the lifestyle of food handlers. Examination of fingernail contents of food handlers for *Salmonella* or *Shigella* is one way of indicating the possible contamination of food [31]. However, the current study did not assess the *Salmonella* and *Shigella* carriage of fingernail contents.

Antimicrobial susceptibility pattern data showed that ciprofloxacin, ceftriaxone, gentamicin, chloramphenicol and cotrimoxazole were effective against the *Shigella* isolates. Our finding was comparable with studies reported from Ethiopia, Haramaya University ceftriaxone(16.7%) [33], Jimma, gentamicin(1.3%) [34], and Harar (3.6%) [35]. Whereas our result showed lower resistance pattern compared to the studies conducted in Ethiopia, Addis Ababa gentamicin(75.6%) [36], and
Gondar, ciprofloxacin (8.9%) and cotrimoxazole(73.4%) [37,38]. This increase of resistance from those reports indicated that difference in the geographical area, study period and study design. Increased resistance was observed in our finding which is in line with a study reported from Harar, ampicillin (100%) [35], Arba Minch, amoxicillin (100%) [14].

In the current study, isolates of *Salmonella* species were sensitive to gentamicin, ciprofloxacin, chloramphenicol, ceftriaxone, cotrimoxazole and clarithromycin. This is consistent with reports from Gondar University, Ethiopia[24, 25, 38]. Increased resistance was observed in our findings for amoxicillin-clavulanic, amoxicillin and ampicillin which were supported by studies reported from Ethiopia, Arba Minch, Jimma and Bahir Dar [14,12,23,35]. This might be due to misuse or inappropriate use of these antibiotics for other infections in addition to the replacement of sensitive strains by resistant strains.

In the present study, the prevalence of multidrug resistance towards *Salmonella* and *Shigella* were observed. Of the total (54.54%) *Salmonella* and (90.9%) *Shigella* species of all the isolates were resistant at least to three antimicrobials. One isolate of *Shigella* was resistant to six classes of antimicrobial agents. This study is supported by a study conducted in Ethiopia Butajira [25], Addis Ababa [22], Haramaya University [33] and Gondar [24]. This increased multidrug resistance might be due to genetic variation by mutation, irrational use of antimicrobials and less hygienic practice of the food handlers.

**Conclusion**

The overall prevalence of *Salmonella* and *Shigella* in the study area found to be 7.3% and 3.7% respectively. The *Salmonella* and *Shigella* carriage was significantly associated with washing hand after touching dirty, washing hands after using the bathroom, fingernails status. chloramphenicol, ceftriaxone and ciprofloxacin were sensitive antimicrobials. The majority of *Salmonella* and *Shigella* were multidrug-resistant. Regular medical checkup, improve personal hygiene and environmental sanitation and consistent training about food preparation and handling for the food handlers of Adigrat University is very important to prevent the risk of infection for the University community having close contact with those carriers.
Limitation

Fingernail content examination could not be identified, this may support to know the contamination due to poor fingernail hygiene and food handling practices.

Despite this limitation, the methods used to isolate and characterize the antimicrobial susceptibility pattern of *Salmonella* and *Shigella* spp. are comprehensive.

Abbreviation

**ATCC**: American Type Culture Collection; **CI**: confidence interval; **CLSI**: Clinical and Laboratory Standards Institute; **MDR**: Multi-drug resistant Multi-Drug Resistant; **SOPs**: Standard Operating Procedures **USA**: United States of America; **WHO**: World health organization;

Declarations

**Ethics approval and consent to participate**

The study was approved by the College of medicine and health sciences Research ethical review committee of Adigrat University, Ethiopia (Consent Ref Number AGU/CMHS/044/2018 approval dated 07/04/2018 Official letter was obtained from Adigrat University (Consent Ref Number AGU/CMHS/RCSH/19/2018 approval dated 25/04/2018. Written informed consent was sought from each study participant before sample collection and maintained throughout the study. All participants were given code numbers to keep their identity confidential.

**Consent for publication**

Not applicable

**Availability of data and materials**

All data collected and analyzed during this study were included in the manuscript. But if the full paper is needed, it will be shared upon request by the editor from the corresponding author.

**Competing interests**

The authors’ declared that there were no competing interests

**Funding**

Not applicable

**Authors’ contributions**
HL designed the study, collection, analysis, and interpretation of data, and drafted the manuscript.

TK, AG, BB, and BF designed the study, supervised data collection both on-field and in the laboratory, and prepared the manuscript. ST, GG, HN, and GA read and approved the final manuscript.

Acknowledgements

The authors gratefully acknowledge the food handlers of Adigrat University for their participation in the study. We also wish to extend our deep appreciation to Adigrat University, College of Medicine and Health Sciences for providing us with the opportunity to do this research and allowing the finance.

References

1. Centres for Disease Control and Prevention (CDC). Surveillance for foodborne disease outbreaks-United States, 2008. MMWR 2010; 59(31):1277-1280.

2. World Health Organization (WHO). (2007). Fact Sheet Number 237: Food safety and foodborne illness. Geneva, Switzerland: World Health Organization. Available at http://www.who.int/media_centre/factsheets/fs237/en/. accessed June 26, 2017

3. Coburn, G. A. Grassl, and B. B. Finlay, “Salmonella, the host and disease: a brief review,” Immunology and Cell Biology 2007; 85:2:112-118

4. Majowicz SE, Musto J, Scallan E, Angulo FJ, O'Brien SJ, Jones TF, The global burden of nontyphoidal Salmonella gastroenteritis. Clin Infect Dis. 2010; 50:882-9

5. Centres for Disease Control and Prevention. Surveillance for foodborne disease outbreaks the United States, 2008. MMWR Morb Mortal Wkly Rep 2011; 60(35):1197-1202

6. World Health Organization (WHO) laboratory identification and antimicrobial susceptibility testing of bacterial pathogens of public health concern in the developing world Geneva, Switzerland: World Health Organization; 2003

7. Smith SI, Fowora M.A, Goodluck H.A, Nwaokorie F.O, Aboaba O.O,Opere B. Molecular typing of Salmonella spp isolated from food handlers and animals in Nigeria Int J Mol
8. Feasey NA, Dougan G, Kingsley RA, Heyderman RS, and Gordona MA. Invasive non-typhoidal salmonella disease: an emerging and neglected tropical disease in Africa. *Lancet* 2012; **379**(9835): 2489-2499.

9. Pala K, Ozakin C, Akis N, Sinirtas M, Gediko S, Aytekin H: Asymptomatic carriage of bacteria in food workers in Nilüfer district, Bursa, Turkey. *Turk J Med Sci* 2010; **40**(1):133-139.

10. Khurana S, Taneja N, Thapar R, Sharma M, Malla N: Intestinal bacterial and parasitic infections among food handlers in a tertiary care hospital of North India. *Trop Gastroenterol* 2008; **29**:207-209.

11. Ao TT, Feasey NA, Gordon MA, Keddy KH, Angulo FJ, Crump JA. Global Burden of Invasive Nontyphoidal *Salmonella* Disease *Infect Dis* 2015; **21**(6):441-449 DOI: http://dx.doi.org/10.3201/eid2106.140999

12. Beyene G, Tasew H. Prevalence of intestinal parasite, Shigella and Salmonella species among diarrheal children in Jimma health centre, Jimma southwest Ethiopia: a cross-sectional study. *Ann Clin Microbiol Antimicrob*. 2014; **13**: Published 2014 Feb 5. doi:10.1186/1476-0711-13-10

13. Mokhtari W, Nsaibia S, Majouri D, Ben Hassen A, Gharbi A, and Aouni M, Detection and characterization of Shigella species isolated from food and human stool samples in Nabeul, Tunisia, by molecular methods and culture techniques *J Applied Microbiol* 2012; **113**: 209-222 doi: 10.1111/j.1365-2672.2012.05324

14. Mama M, Getaneh A. Prevalence, antimicrobial susceptibility patterns and associated risk factors of *Shigella* and *Salmonella* among food handlers in Arba Minch University, South Ethiopia. *BMC Infect Dis*. 2016; **16**:686

15. Scallan E, Hoekstra RM, Angulo FJ, Tauxe RV, Widdowson MA, Roy SL, Jones JL, Griffin
PM. Foodborne illness acquired in the United States-major pathogens. *Emerg Infect Dis.* 2011;**17**(1):7-15

16. Bonkoungou IJO, Haukka K, Österblad M, Hakanen AJ, Traoré AS, Barro N, et al. Bacterial and viral etiology of childhood diarrhoea in Ouagadougou, Burkina Faso. *BMC Pediat.* 2013; **13**(36):1-6

17. Afeworki G, Lirneneh Y. Multiple drug resistance within Shigella serogroups. *Ethiop Med J.* 1980; **18**:7-11.

18. Roma B, Worku S, T/Mariam S, Langeland N. Antimicrobial susceptibility pattern of Shigella isolates in Awassa. *Ethiop J of Health Dev.* 2000; **14**:149-54.

19. Beyene G., Asrat D., Mengistu Y., Aseffa A. and Wain J. Typhoid fever in Ethiopia. *J Infect Developing Countries* 2008; **2**:448-453

20. Clinical and Laboratory Standards Institute (CLSI): Performance standards for antimicrobial susceptibility testing; Twenty-sixth Informational Supplement. *Wayne PA* 19087 USA 2016; **36**(1).

21. Mobolaji OA, Olubunmi OF. Assessment of the hygienic practices and the incidence of enteric bacteria in food handlers in small businesses in an urban area in Abeokuta. *Int J Microbiol Res.* 2014;**5**(3):41-9

22. Aklilu A, Kahase D, Dessalegn M, Tarekegn N, Gebremichael S, Zenebe S. et al. Prevalence of intestinal parasites, salmonella and shigella among apparently health food handlers of Addis Ababa University student’s cafeteria, Addis Ababa, Ethiopia. *BMC Res Notes* 2015; **8**:17:1-6

23. Abera B, Biadegelgen F, Bezaeh B. Prevalence of Salmonella typhi and intestinal parasites among food handlers in Bahir Dar Town, Northwest Ethiopia. *Ethiop J Health Dev.* 2010;**24**(1):46-50

24. Garedew-Kifelew L, Wondafrash N, Feleke A. Identification of drug-resistant
Salmonella from food handlers at the University of Gondar, Ethiopia. *BMC Res Notes*. 2014;7(1):545

25. Mengistu G, Mulugeta G, Lema T, Aseffa A. Prevalence and Antimicrobial Susceptibility Patterns of *Salmonella* serovars and *Shigella* *J Microb Biochem Technol*. 2014; S2: 006. doi:10.4172/1948-5948.S2-006

26. Ifeadike C, Ironkwe O, Adogu P, Nnebue C, Emelumadu O, Nwabueze S, Ubajaka C. Prevalence and pattern of bacteria and intestinal parasites among food handlers in the Federal Capital Territory of Nigeria. *Niger Med J*. 2012;53(3):166

27. Andargie G, Kassu A, Moges F, Tiruneh M, Huruy K: Prevalence of bacteria and intestinal parasites among food handlers in Gondar Town, Northwest Ethiopia. *J Health Popul Nutri* 2008;26(4):451-455.

28. Nigusse D and Kumie A. Food hygiene practices and prevalence of intestinal parasites among food handlers working in Mekelle university student’s cafeteria, Mekelle. *Global Adv Res J of Soc Sci (GARJSS)*, 2012;1(4):065-071

29. Dagnew M, Tiruneh M, Moges F, Gizachew M. Bacterial Profile and Antimicrobial Susceptibility Pattern among Food Handlers at Gondar University Cafeteria, Northwest Ethiopia. *J Infect Dis Ther*. 2013;1: 105. doi:10.4172/2332- 0877.1000105

30. Abera B, Yitayew G, Amare H. *Salmonella* serotype Typhi, *Shigella*, and intestinal parasites among food handlers at Bahir Dar University, Ethiopia *J Infect Dev Ctries* 2016; 10(2):121-126

31. Tefera T, Mebrie G. Prevalence and Predictors of Intestinal Parasites among Food Handlers in Yebu Town, Southwest Ethiopia. *PLoS ONE*. 2014; 9 (10): e110621. doi:10.1371/journal.pone.0110621

32. Zain MM, Naing NN. Sociodemographic characteristics of food handlers and their knowledge, attitude and practice towards food sanitation: A preliminary report.
Southeast Asian J Trop Med Public Health 2002; 33(2): 410-7.

33. Marami D, Hailu K, and Tolera M. Prevalence and antimicrobial susceptibility pattern of Salmonella and Shigella species among asymptomatic food handlers working in Haramaya University cafeterias, Eastern Ethiopia. BMC Res Notes. 2018; 11:74

34. Mache A. Salmonella serogroups and their antibiotic resistance patterns isolated from diarrhoeal stools of pediatric out-patients in Jimma Hospital and Jimma Health Center, South West Ethiopia. Ethiop J Health Sci. 2002; 37: 37-45

35. Reda AA, Seyoum B, Yimam J, Andualem G, Fiseha S, Vandeweerd J-M. Antibiotic susceptibility patterns of Salmonella and Shigella isolates in Harar, Eastern, Ethiopia. J Infect Dis Immun. 2011; 3: 134-139

36. Asrat D. Shigella and Salmonella serogroups and their antibiotic susceptibility patterns in Ethiopia. East Mediterr Heal J. 2008;14:760-7

37. Yismaw O, Negeri C, Kassu A. A five-year antimicrobial resistance pattern observed in Shigella species isolated from stool samples in Gondar University Hospital, northwest Ethiopia. Ethiop J Heal Dev. 2006;20:194-8

38. Tiruneh M. Serodiversity and antimicrobial resistance pattern of Shigella isolates at Gondar University teaching hospital, Northwest Ethiopia. Jpn J Infect Dis. 2009; 62(2):93-7

Tables

Table 1 Socio-demographic characteristics of food handlers in Adigrat University student cafeteria, Tigrai, North Ethiopia March to August 2018 (N=301)
## Demographic characteristics

| Characteristics                  | Male | Female |
|----------------------------------|------|--------|
| **Sex**                          |      |        |
| Male                             | 1(2.7) | 36(97.3) |
| Female                           | 32(12.1) | 23(92.7) |
| **Age**                          |      |        |
| ≤ 20                             | 2(3.7) | 52(96.3) |
| 21-40                            | 31(12.5) | 216(87.4) |
| **Marital status**               |      |        |
| Single                           | 16(12.6) | 111(87.4) |
| Married                          | 17(9.7) | 157(90.2) |
| **Educational level**            |      |        |
| Lower than Secondary school      | 16(8.5) | 171(91.5) |
| Higher than secondary school     | 17(14.9) | 97(85.1) |
| **Certified in food preparation and handling** | No | Yes |
| **Medical check-up**             |      |        |
| No                               | 15(23.8) | 48(76.2) |
| Yes                              | 2(1.8) | 11(98.2) |

Table 2 Bivariate logistic regression analysis of factors associated with *Salmonella* and *Shigella* infections among food handler’s working at Adigrat University Students’ Cafeteria, Tigrai, Northern Ethiopia, March to August 2018 (N=301)
|                                | Yes only with water | No |  
|--------------------------------|---------------------|----|  
| Finger nail status            | Trimmed             | 4(2.2) | 17  
|                                | untrimmed           | 29(23.6) | 94  
| After blowing nose            | Yes with water and soap | 7(8.0) | 81  
|                                | Yes only with water | 11(11.7) | 83  
|                                | No                  | 15(12.6) | 10  
| Touch food with bare hands    | No                  | 16(14.0) | 98  
|                                | Yes                 | 17(9.1) | 17  
| Years of service              | ≤1                  | 9(25.0) | 27  
|                                | 1-2                 | 9(21.4) | 33  
|                                | >2                  | 15(6.7) | 20  
| Certified in food preparation and handling | No | 29(11.5) | 224  
|                                | Yes                 | 4(8.3) | 44  
| Medical check-up              | No                  | 10(18.5) | 44  
|                                | Yes                 | 23(9.3) | 22  

Key: \(^{a}\)(COR=Crude odds ratio); \(^{b}\)(CI=Confidence interval); 1(referent).

Table 3: Multivariate logistic regression analysis of factors associated with Salmonella and Shigella isolates among food handler’s working at Adigrat University Students’ Cafeteria, Tigrai, Northern Ethiopia, March to August 2018 (N=301)
| Variables | Growth of *Salmonella* and *Shigella* | COR\(^a\) (95% CI) |
|-----------|--------------------------------------|---------------------|
| **Negative** | **Positive** | **COR\(^a\)** |
| N (\%) | N (\%) | | |
| Hand washing using after using the bathroom | | |
| Yes with water and soap | 112 (99.1) | 1 (0.9) | 1 |
| Yes only with water | 108 (86.4) | 17 (13.6) | 17.630 (2.3) |
| No | 48 (76.2) | 15 (23.8) | 35.000 (4.4) |
| Hand washing after touching dirty materials | | |
| Yes with water and soap | 141 (96.6) | 5 (3.4) | 1 |
| Yes only with water | 113 (89.0) | 14 (11.0) | 3.494 (1.2) |
| No | 14 (50.0) | 14 (50.0) | 28.200 (8.8) |
| Hand washing before food handling | | |
| Yes with water and soap | 112 (98.2) | 2 (1.8) | 1 |
| Yes only with water | 129 (90.8) | 13 (9.2) | 5.643 (1.2) |
| No | 27 (60.0) | 18 (40.0) | 37.333 (8.1) |
| Finger nail status | | |
| Trimmed | 174 (97.8) | 4 (2.2) | 1 |
| untrimmed | 94 (76.4) | 29 (23.6) | 13.420 (4.5) |

Key: \(^a\)(COR=Crude odds ratio); \(^b\)(CI=Confidence interval); \(^c\)(AOR=Adjusted odds ratio); 1(referent).

Table 4 Antimicrobial susceptibility patterns of *Salmonella* and *Shigella* isolated from food handlers Of Adigrat University Students’ Cafeteria, Tigrai, Northern Ethiopia, March to August 2018 (N=33)

| Isolates(n) | Sensitivity pattern n(\%) | Antibiotic agents N (\%) | | | | |
|-------------|--------------------------|--------------------------|---|---|---|---|---|
|              | GM | AML | AMP | CIP | CLR | TS |
| *Salmonella* n(22) | | | | | | | |
| S | 22 (100) | 1 (4.5) | 0 (0.00) | 22 (100) | 18 (81.8) | 20 (90.9) |
| R | 0 (0.00) | 21 (95.5) | 22 (100) | (0.00) | 4 (18.2) | 2 (9.1) |
| *Shigella* n (11) | | | | | | | |
| S | 10 (90.9) | 0 (0.00) | 0 (0.00) | 11 (100) | 2 (18.2) | 10 (90.9) |
| R | 1 (9.1) | 11 (100) | 11 (100) | 0 (0.00) | 9 (81.2) | 1 (9.1) |

Key: S = Sensitive R = Resistant, GM Gentamicin, AML Amoxicillin, AMP Ampicillin, CIP=Ciprofloxacin, CLR Clarithromycin, TS Cotrimoxazole, AMC Amoxicillin-clavulanic acid CHL Chloramphenicol, CRO
### Ceftriaxone

Table 5 Multidrug-resistant of *Salmonella* and *Shigella* isolated from food handler's working at Adigrat University Students’ Cafeteria, Tigrai, Northern Ethiopia, March to August 2018 N=33

| Antimicrobials | *Salmonella* | *Shigella* |
|----------------|--------------|------------|
| For three      |             |            |
| AML,AMP,CLR    | 2 (16.67)   | 4 (40)     |
| AML,AMP,CHL    | -           | 1 (10)     |
| AML,CLR,AMC    | 7 (58.34)   | -          |
| AMP,CLR,AMC    | 1 (8.33)    | -          |
| AML,AMP,TS     | 1 (8.33)    | -          |
| FOR FOUR       |             |            |
| AML,AMP,CLR,AMC| 1 (8.33)    | 3 (30)     |
| FOR FIVE       |             |            |
| AML,AMP,CLR,AMC,CHL | -          | 1 (10)     |
| FOR SIX        |             |            |
| GM,AML,AMP,CLR,TS,AMC | -          | 1 (10)     |
|                | 12 (100)    | 10 (100)   |

Key: GM Gentamicin, AML Amoxicillin, AMP Ampicillin, CIP=Ciprofloxacin, CLR Clarithromycin, TS Cotrimoxazole, AMC Amoxicillin-clavulanic acid CHL Chloramphenicol, CRO, Ceftriaxone

MDR multidrug resistant; MDR definition for *Salmonella* and *Shigella* percent is computed from total number of *Salmonella* and *Shigella*