

INTERNATIONAL JOURNAL OF ADVANCES IN MEDICAL BIOTECHNOLOGY

# Bacteriological assessment of stethoscope used by healt care personnel in attat hospital, SNNP, Gurage Zone, Ethiopia

# Tamirat Salile<sup>1</sup>

\*Corresponding author: e-mail address: tamewoldia@gmail.com

Abstract: The stethoscope has always been an important element of a physician's toolkit when it comes to examining patients. The widespread use of stethoscopes by health–care workers for patient examinations makes them a potential source of nosocomial infection transmission. The goal of this study was to see if stethoscopes used by different health–care professionals in Attat hospital may transmit bacteria. From April to June 2018, a cross–sectional study was done in the molecular laboratory of Wolkite University's department of biotechnology and biology. A total of 26 stethoscopes from health workers who had direct contact with patients were gathered during the study period. The sample was obtained using a sterile cotton–tipped applicator saturated in a sterile solution of physiologic saline (0.85 % sodium chloride) to swab the whole surface of the stethoscope's diaphragm and then inoculated into macconkey agar, tryptone soya agar, and blood agar medium. 18(69.2%) stethoscopes out of total collected stethoscopes had bacterial growth and 12 bacterial isolates were selected and characterized to genus level. Isolates include staphylococcus aureu s(37.5%), coagulase negative staphylococci (28.12%), Streptococcussp. (21.88%), and Bacillus sp.(12.5%). All isolates were susceptible to the co–trimoxazole and ciprofloxacine, while resistant to cifoxitine. They showed intermediate growth against vancomycine. All except streptococcus were found resistant against penicillin. Both S. aureus and CoNS were sensitive to the chloramphenicol; Streptococcus was intermediate while bacillus was resistant to the chloramphenicol. All stethoscopes (42.2 %) that had never been cleaned and were last cleaned a week ago were severely contaminated, while those washed multiple times a day and cleansed between each patient before the examination of the patients had lower levels of contamination (27 %).

Keywords: Bacterial Isolate. Nosocomial Infection. Stethoscope.

# Introduction Background of the study

Nosocomial infections have existed since the beginning of hospitals, and they continue to be a significant public health issue even in the modern era of antibiotics. When infections become clinically obvious during hospitalization (at least 72 hours after admission), they are classified as nosocomial<sup>[21]</sup>. Such infections are caused by a variety of factors, including the emergence and persistence of multidrug-resistant bacteria, patients' compromised immune systems, and mechanical transmission of microorganisms<sup>[12]</sup>, all of which result in high morbidity and mortality, prolonged hospitalization, increased antibiotic use, and increased costs<sup>[12]</sup>. According to studies, these infections occurred in 5% to 10% of all hospitalizations in Europe and North America, and in more than 40% of hospitalizations in Asia, Latin America, and Sub–Saharan Africa<sup>[34]</sup>.

According to<sup>[32]</sup>, more than 1.4 million people globally are infected with illnesses acquired in hospitals at any given time, and health–care personnel are possible sources of these infections. Because many infections can be spread through the hands, all health–care professionals must wash their hands before and after each patient encounter<sup>[35]</sup>. Diseases can be transmitted through contaminated medical devices, and outbreaks of hospital–acquired infections have been connected to electronic thermometers, blood pressure cuffs, stethoscopes, latex gloves, masks, neckties, pens, badges, and lanyards, white coats, computers, and keyboards<sup>[31]</sup>.

The sterilization and disinfection of intrusive equipment and devices prior to interventions are frequently overlooked. Stethoscopes are the most commonly utilized medical devices by health care personnel to examine the health of patients among those equipments. As a result, they frequently come into touch with a large number of patients and have been identified as potential nosocomial infection vectors in various regions of the world<sup>[26, 31, 27]</sup>.

According to a similar report from Jimma University Specialized Hospital, bacterial contamination of the stethoscope is significant and could be a vector for illness transfer between patients and health care staff [29]. Pathogens can adhere and establish themselves on the diaphragms of stethoscopes after contact with contaminated skin, and then be conveyed to other patients if the stethoscope is not cleansed <sup>[18]</sup>.

There are also more cases of antibiotic–resistant bacteria being transmitted from one patient to another via stethoscopes <sup>[31, 9, 20, 12]</sup>. In a hospital setting, these antibiotic–resistant organisms are capable of causing serious infections, necessitating contact isolation and rigorous treatment to limit the spread of the organis– ms<sup>[12]</sup>. Ceftazidime–resistant Klebsiella pneumonia, vancomycin–resistant enterococci, methicillin–resis–

<sup>&</sup>lt;sup>1</sup>Addis ababa university

Received 02 August 2021; Accepted 30 September 2021; Available online 1 October 2021. DOI: https://doi.org/10.52466/ijamb.v4i2.95

Salile

aeruginosa, gentamicin–resistant Pseudomonas aerugi– nosa, and penicillin–resistant pneumococci are examples of antibiotic–resistant organisms<sup>[16, 10, 22].</sup>

Infection transmission in hospitals (nosocomial infections) is a major issue caused by contaminated medical equipment and health-care workers (HCWs). Medical devices that have not been adequately sterilized/ disinfected may spread bacteria from one patient to the next. Due to rising morbidity and cost burden, health-care-acquired infections are becoming a major concern not only for doctors, but also for patients, and stethoscope disinfection is still not a widely accepted practice among most health-care workers.

Despite the fact that stethoscopes are a possible vector for the transfer of health-care-associated illnesses and resistant bacteria, health-care professionals fail to disinfect them<sup>[31]</sup>. Swiping stethoscopes with alcohol pads is the current gold standard for stethoscope decontamination<sup>[27]</sup>. To prevent nosocomial infections, medical devices such as stethoscopes should be tested for microbial colonization on a regular basis, and health care staff should be educated on proper cleaning procedures <sup>[7]</sup>.

Infection transmission in hospitals (nosocomial infections) is a major issue caused by contaminated medical equipment and health-care workers (HCWs). Medical devices that have not been adequately sterilized/ disinfected may spread bacteria from one patient to the next. Because of rising morbidity and cost burden, health-care-acquired infections connected with stethoscopes are now a major concern for doctors as well as patients, and stethoscope disinfection is still not a widely accepted practice among most health-care workers. To our knowledge, there has only been one study on the function of stethoscopes in the transmission of nosocomial infections, and none has been conducted in Ethiopia's south. In Attat Hospital, a referral hospital serving people of ChehaWoreda and nearby communities in Southwest Ethiopia, we wanted to look into the role of stethoscopes as potential fomites for possibly dangerous bacteria. Therefore, this study was focus on the following objectives i) to establish the bacteriological agents responsible for stethoscope contamination in Attat Hospital, as well as to examine healthcare personnel's attitudes and knowledge about stethoscope hygiene behavior; ii) In order to identify and characterize bacterial isolates based on biochemical and morphological tests; iii) For determining a drug resistance profile of selective isolates; iv) To explore the behavior, attitudes and beliefs about stethoscope hygiene amongst healthcare personnel within the hospital's various clinical units.

# Literature Review The different types of nosocomial infections

The CDC and the National Healthcare Safety Network (NHSN) divide health–care–associated infection sites into 13 primary kinds based on clinical and biological criteria, with roughly 50 potentially specific infection sites for surveillance. Surgical wound and other soft tissue infections, urinary tract infections (UTI), respiratory infections, gastroenteritis, and meningitis are the most frequent nosocomial diseases that can arise in a hospital setting<sup>[25]</sup>. However, with the increased use of invasive procedures for therapeutic and diagnostic purposes, cancer chemotherapy, immunotherapy, and advancements in organ transplantation, changes in the distribution of nosocomial infection sites can be observed over time.

### **Epidemiology of nosocomial infections**

It is estimated that about 10% of hospital patients or more than 2-million hospitalized patients are annually suffering from hospital infection in the USA; and an estimated annual death rate is 20,000, which may reach even up to 88,000 deaths per year. Basic epidemiologic patterns can be used to guide prevention and control actions in hospital-acquired infections. The virus that causes hospital infection has reservoirs, can be transferred in predictable ways, and needs a vulnerable host<sup>[33]</sup>. The inanimate environment, such as surgical instruments and the operating room, and the animate environment, such as diseased or colonized health care staff, patients, and hospital visitors, could be reservoirs and sources of infection. Cross-infection from an endogenous flora present in the patient or autoinfection from an endogenous flora found in the patient are two possible modes of transmission for hospital acquired infection. For example surgical site infection can be caused by an endogenous flora that translocate to a normally sterile site or when the sterile peritoneal cavity is contaminated by spillage from the gastrointestinal tract; and by an exogenous source of microbial contamination that comes from the surgical team, surgical instrument and the theatre environment.

Furthermore, aseptic procedures were not followed strictly by the majority of the nurses and physicians in several practice areas and are found to be significant for the transmission of the infection<sup>[24]</sup>.

### Nosocomial infections: sources and transmission

Infections are caused by nosocomial microorganisms that can come from either endogenous or external sources. Hospital staff, other patients, visitors, food, water, fomites, urinary catheters, intravenous devices, respiratory apparatus, and other prosthesis are all examples of animate and inanimate sources of exogenous infections. Contact is the most common way for nosocomial illnesses to spread, generally directly but occasionally indirectly through bodily secretions. Air can also be a source of airborne nosocomial viruses that infect the respiratory tract (e.g., in droplet nuclei and aerosols). Food–borne and water–borne diseases can enter through the faeco–oral pathway. The oropharynx, gastrointestinal system, and urinary tract are the most prevalent reservoirs for nosocomial colonizers <sup>[26]</sup>.

# Nosocomial infection risk factors

For a variety of reasons, hospitalized patients are at an unusually high risk of infection. Intrinsic and extrinsic factors are roughly classified into two categories. Intrinsic risk factors are those that are present in the patient as a result of the underlying disease. Patient care may contain extrinsic risk factors. Concurrent infections, prosthetic devices, surgery, immunosuppressive medications, broad-spectrum antibiotic therapy, and the emergence of multidrug-resistant organisms are some of the general predisposing factors that make patients prone to nosocomial infections. Other risk factors include the patient's age, length of stay in the hospital, underlying conditions such as diabetes, malignancies, or ward congestion. The length of hospital stay is the most important risk factor for contracting a nosocomial infection among the multiple risk variables [17].

# **Nosocomial infection agents**

Infections in hospitals are caused by a wide range of bacteria, and any bacterium has the potential to cause an infection in hospitalized patients.

# Microorganisms

Nosocomial infections can be caused by a variety of microorganisms. The infecting organisms differ depending on the patient demographic, the health care setting, the facility, and the country.

# A. Bacteria

These are the most commonly found nosocomial pathogens in hospitals. There is a distinction to be made between commensal bacteria found in the typical flora of healthy humans and pathogenic bacteria. These provide an important protective role by preventing harmful germs from colonizing the area. If the native host is harmed, some commensal bacteria may cause illness. Intravascular line infection is caused by cutaneous coagulase negative staphylococci, and urinary infection is caused by intestinal Escherichia coli.

Pathogenic bacteria have a higher pathogenicity and, independent of host status, cause infections (sporadic or epidemic). Anaerobic Gram–positive rods (such as Clos– tridium) induce gangrene, for example.

Gram-positive bacteria: Staphylococcus aureus (a cutaneous bacterium that colonizes both hospital staff and patients' skin and nose) causes a wide range of lung, bone, heart, and bloodstream infections and is usually antibiotic-resistant; beta-haemolytic streptococci are also essential.

Gram-negative bacteria, such as E. coli, Proteus,

Klebsiella, Enterobacter, and Serratiamarcescens, can colonize places where the host's defenses are impaired (catheter insertion, bladder catheter insertion, cannula insertion) and cause significant infections (surgical site, lung, bacteraemia, peritoneum infection). They could also be extremely resistant. Gram–negative bacteria, such as Pseudomonas spp., are frequently found in wet and damp environments. They may colonize hospitalized patients' gastrointestinal tracts.

Other germs provide a distinct threat in hospitals. For example, Legionella species can cause pneumonia (sporadic or endemic) in people who inhale polluted water aerosols (air conditioning, showers, and therapeutic aerosols).

# **B. Viruses**

Many viruses, including hepatitis B and C (transfusions, dialysis, injections, and endoscopy), respiratory syncytial virus (RSV), rotavirus, and enteroviruses, can be transmitted nosocomially (transmitted by hand-to-mouth contact and via the faecal-oral route). Other viruses that can be transferred include CMV, HIV, Ebola, influenza viruses, herpes simplex virus, and varicella-zoster virus.

# C. Parasites and fungi

Some parasites, such as Giardia lamblia, are easily spread between adults and children. Many fungi and parasites are opportunistic organisms that cause infections when the immune system is suppressed by antibiotics (Candida albicans, Aspergillus spp., Cryptococcus neoformans, Cryptosporidium). In immune–compromised patients, they are a primary source of systemic infections. Contamination of the environment by airborne organisms that originate in dust and dirt, such as Aspergillus spp., is also a worry, particularly during hospital building. Sarcoptes scabies (scabies) is an ectoparasite that has caused outbreaks in health care on several occasions.

# Diagnosis

The diagnosis and identification of hospital–acquired infection involves interpretation of clinical and laboratory findings. Clinically, a patient is assessed based on clini– cal sign and symptoms developed due to the infection. Pain, soreness, redness, localized swelling, and purulent discharge from the wound are symptoms of a superficial incision site infection. The patient developed a fever (> 38°C), localized discomfort or tenderness, and puru– lent discharge from the incision if the infection was at a deep cut. Fever (>38°C), urgency, frequency, and dysu– ria were all reported in a patient with symptomatic UTI. However, patients under the age of one year may have hypothermia (37°C), apnea, bradycardia, lethargy, or vo– miting.

The pathogen was isolated by urine culture, which was used to make the laboratory diagnoses. The amount and

types of bacteria in the urine must be determined as part of the diagnostic process. If a mid-stream urine culture includes 105 organisms per ml and no more than two types of microbes, it is deemed positive.

### NOSOCOMIAL INFECTION PREVENTION

It is the responsibility of all individuals and services providing health care to prevent nosocomial infections. In addition, everyone must work together to limit the risk of infection for both patients and employees. Although not all hospital infections are preventable, the majority of them can be. Surveillance of NIs is an important aspect of infection control, and it is widely recognized as a first step toward prevention around the world. Reduced heal– th–care–associated infection rates, on the other hand, is dependent on a number of factors. Staff–related proce– dures, particularly hand hygiene, have recently received a lot of attention. Furthermore, there has been a growing understanding that environmental controls should be an important part of any overall plan for preventing health– care–associated illnesses [2].

Hand washing is still the most critical action in infection prevention. Gloves, gowns, and masks have a role in avoiding infections, but they are frequently misused, resulting in unnecessary service expenses. Many people are visibly disturbed when their inadequate hygiene practices are revealed, and many are outraged when it is claimed that they could be disease vectors, spreading dangerous bacteria among their patients, complicating infection control efforts [2].

### MATERIALS AND METHOD

### STUDY AREA AND PERIOD

The current research was carried out at the Attat Specialized Hospital, which is located 175 kilometers southwest of Addis Ababa, 17 kilometers from the town of Wolkite on the road to Hosanna in the Cheha Woreda of the Gurage Zone, SNNPRs, Ethiopia. Since its establishment in 1969, the hospital was managed and run by Medical Mission Sisters under the Eparchy of Emdibir. According to medical mission sisters, the hospital has ten wards (0–9) providing extensive integrated health services for more than 800,000 people with in and out of the operational area. Between April and June 2018, samples were evaluated at Wolkite University's Department of Biotechnology and Biology laboratory, which is part of the College of Natural and Computational Science.

### **STUDY DESIGN**

A cross–sectional descriptive study was conducted using a structured questionnaire and specimen asses– sment from the stethoscope of Attat Hospital healthcare workers.

### STUDY POPULATION

All healthcare personnel (doctors, nurses, health of-

ficers and students) having their stethoscope on-hand during data collection constitute the source population for the study. According to the report from the medical director of Attat hospital, they had around 50 stethoscopes for 10 wards, which were used as the sampling frame. A proportional sample size was determined for each department (including inpatients and OPD) and participants were selected using a simple random sampling.

### **S**AMPLE SIZE

There are 26 stethoscopes required for sample collection from different wards and from different professionals.

### **S**ELECTION CRITERIA

Healthcare personnel who were willing to give informed consent and the study included anyone who had their personal stethoscope on hand at the time of data collection. The study did not include those who:

- Do not have a stethoscope on hand during data collection.
- Already participated in the study while working in another ward.
- Refuse to give informed consent.

### **D**ATA AND SAMPLE COLLECTION

The investigators went to any inpatient or outpatient department for data and sample collection without any prior notice. Then, after obtaining consent, self-administered questionnaires were utilized to gather socio-demographic data characteristics (gender, profession and experience) of participants, use of stethoscopes, stethoscope cleaning habits, and perceived barriers to cleaning. Swam samples for bacteriological and antibio-tic resistance profiling were taken from the diaphragmatic section of stethoscopes using a sterile cotton-tipped applicator bathed in sterile normal saline (0.85 % w/v).

The obtained swabs were immediately placed in Amies transport media, and samples were sent to the laboratory in an ice box with proper and comprehensive labeling, along with the questionnaire.

# BACTERIAL PATHOGEN ISOLATION, ENUMERATION, AND IDENTI-FICATION

The material was inoculated in duplicate on Blood agar, Tryptone soya agar, and MacConkey agar and incubated aerobically at 370C for 48 hours after gentle mixing. The media were inspected for bacterial growth after incubation, and the total number of colonies for each sample was tallied. Significant growth was defined as a colony count of more than 20cfu/diaphragm [29], and the stethoscope was deemed contaminated. In tryptone soy broth and agar slant, representative colonies from contaminated stethoscopes were purified and stored. Following normal bacteriological techniques, the isolates were identified to the genus and species level.

### Salile

Based on colony characteristics (appearance, size, and color), cell morphology, Gram reactions and KOH test obtained further identification of bacteria was made by a series of biochemical tests. Mannitol salt agar and blood agar plates were used to cultivate Gram–positive cocci. Following that, catalase and tube coagulase tests were performed. Staphylococcus aureus was identified in iso– lates that passed all three tests. Isolates tested negative for tube coagulase was considered coagulase–negative staphylococci (CoNS).b Catalase negative gram positive cocci were cultured on blood agar and pattern of hemoly– sis (alpha, beta, and gamma) was observed.

### Antibiotic sensitivity test

The following antibiotics were used to determine the anti biogram of the isolates: Penicillin ( $10\mu g$ ), Chloram-phenicol ( $30\mu g$ ), Ciprofloxacin ( $5\mu g$ ), Cefoxitin ( $30\mu g$ ), Co-trimaxozole ( $25\mu g$ ), and Vancomycine ( $30\mu g$ ). The antibiotic discs were selected based on availability and current use in health facilities of Ethiopia. Direct colony suspension of the test organism in sterile saline solution were prepared, the turbidity of the inoculum was adjusted to a 0.5 McFarland standard (1.5x108CFU/ml). A new sterile cot-ton-tipped swab dipped in the suspension was used to wipe the surface of Mueller Hinton agar plates within 15 minutes of inoculum formation. Then, within 15 minutes of standard antibiotic discs was applied. MHA plates were

incubated at 370C for 18–24 hours, and the diameter of each antimicrobial disc's zone of inhibition was determined <sup>[8]</sup>.

### **Data analysis**

The SPSS v23 computer software was used to enter and evaluate the data. Categorical variables were displayed in tables and bar graphs, with frequencies and percentages summarized. A statistically significant difference was defined as a P-value of less than 0.05.

#### Results

#### Study participants' socio-demographic profiles

A total of 26 stethoscopes were tested for bacterial contamination by four separate professionals from different hospital wards. Medical ward (IPG & OPD) (8), Gynecology ward (6), Surgical ward (IPD&OPD) (5), Pediatrics ward (IPD&OPD) (3), Emergency ward (2), Neonatal (1), and Delivery ward (1) were among the 10 wards where these health professionals worked (1).

The study included equal number of males (13) and females (13), where the majorities (10, 39%) were doctors, followed by nurses (7, 27%), medical students (5, 19%) and health officers (4, 15%). Most of the participants years of experience were less than 2 years (11, 42%), followed by 2–5 years (9, 34), and 5–10 years and >10 years (3, 12% each) as shown on (Table1).

		UNCONTAMINATED	CONTAMINATED	
		Count (%)		— Total (%)
GENDER	Male	6 (75)	7 (39)	13 (50)
	Female	2 (25)	11 (61)	13 (50)
PROFFESSION	Doctor	4 (50)	6 (33)	10 (39)
	Nurse	1 (12.5)	6 (33)	7 (27)
	Student	2 (25)	3 (17)	5 (19)
	Но	1 (12.5)	3 (17)	4 (15)
EXPERIANCE	<2Yrs	4 (50)	7 (39)	11 (42)
	2–5Yrs	3 (38)	6 (33)	9 (34)
	5–10Yrs	0	3 (17)	3 (12)
	>10Yrs	1 (12)	2 (11)	3 (12)
WARD	Medical	2 (25)	6 (33)	8 (31)
	Gynecology	2 (25)	4 (22)	6 (23)
	Surgical	1 (12.5)	4 (22)	5 (19)
	<b>Pediatrics</b>	2 (25)	1 (6)	3 (11)
	Emergency	1 (12.5)	1 (6)	2 (8)
	Neonatal	0	1 (6)	1 (4)
	Delivery	0	1 (6)	1 (4)

Table 1 - Health-care personnel's socio-demographic features at Attat Hospital, Wolkite.

# Stethoscopes: knowledge, attitudes & practices (KAP) survey

Of 26 stethoscopes studied, almost one third (29%) of the owner reported that the last time they cleaned their stethoscope was last week. Other responded cannot recall (20%), never (17%), today (17%) and yesterday (17%). When asked how often they clean their stethoscope, the majority of respondents (23%) said once daily, followed by an equal number of respondents (19%) who said every patient, once weekly, or numerous times a day. Fewer (12% and 8%) responded rarely if ever and never, respectively. The agents they used for cleaning their stethoscope were alcohol wipes (92%) and antiseptic wipes (8%). No significant relation was identified among the use of disinfectant and bacterial contamination.

With regards to the ideal frequency of cleaning, 69% responded cleaning before and after every patient would be the best approach to keeping the stethoscope clean.81% of the participants believe that stethoscope could transmit infectious agents. For 19% of participants, cleaning their stethoscope at the start and end of the day was sufficient, while 12% had no notion what the appropriate frequency

of cleaning. Forgetfulness (46%), lack of time (18%) (18%) were identified as the major barrier of cleaning stethoscopes. Concern for damage and lack of supplies were the other barriers to cleaning.

# Bacterial contamination of stethoscope diaphragm

After two days of incubation, the diaphragms of all stethoscopes examined from ten wards revealed varying degrees of bacterial contamination. 18 (69.2%) of the 26 stethoscopes tested were heavily contaminated (>20 CFUs/diaphragm), while the other (30.8%) were not.

From 18 contaminated stethoscopes, 11 (61%) were from female health care personnel and 7 (39%) were from male. In terms of profession, the most frequently contaminated stethoscopes were those used by doctors (6, 33%) and nurses (6, 33%), followed by medical students and health officers. While analyzing the proportion, majority of stethoscopes used in Medical (6, 33%), Gynecology (4, 22%) Contamination was found in the Surgical ward (4, 22%). On stethoscope diaphragms from emergency, pediatrics, delivery, and neonatal wards, there was significantly less contamination (**Table 2**).

		UNCONTAMINATED	CONTAMINATED	Total
		Count (%)		
GENDER	Male	6 (75)	7 (39)	13 (50)
	Female	2 (25)	11 (61)	13 (50)
PROFFESSION	Doctor	4 (50)	6 (33)	10 (39)
	Nurse	1 (12.5)	6 (33)	7 (27)
	Student	2 (25)	3 (17)	5 (19)
	Но	1 (12.5)	3 (17)	4 (15)
EXPERIANCE	<2Yrs	4 (50)	7 (39)	11 (42)
	2-5Yrs	3 (38)	6 (33)	9 (34)
	5-10Yrs	0	3 (17)	3 (12)
	>10Yrs	1 (12)	2 (11)	3 (12)
WARD	Medical	2 (25)	6 (33)	8 (31)
	Gynecology	2 (25)	4 (22)	6 (23)
	Surgical	1 (12.5)	4 (22)	5 (19)
	Pediatrics	2 (25)	1 (6)	3 (11)
	Emergency	1 (12.5)	1 (6)	2 (8)
	Neonatal	0	1 (6)	1 (4)
	Delivery	0	1 (6)	1 (4)

Table 2 – Level of bacterial contamination in terms of gender, profession, experience and ward.

### Phenotypic characteristics of bacterial isolates

Based on colony appearance, size, and color, 13 (72%) of the contaminated stethoscope showed a single, uniform colony growth, while the rest (5, 28%) had polymicrobal growth. Of which all colonies identified were gram-positive organisms, while no gram-negative bacteria were observed. A total of twenty-five representative bacterial colonies were primarily selected, in which twelve distinct isolates were purified and preserved for further investigation.

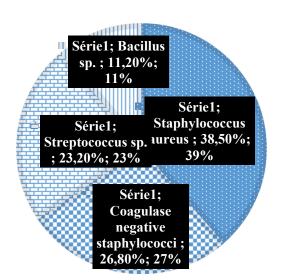
The selected isolates had cultural characteristics that were similar and different, and they were identified to ge– nus level based on their phenotypic and biochemical cha– racters, as well as hemolysis on blood agar and mannitol salt agar for fermentation analysis. The results are pre– sented in **Table 3 and Fig 1**. Among the isolates identi– fied, *Staphylococcus aureus* constitutes38.5%. Coagulase negative *Staphylococcus, Streptococcus* sp. and *Bacillus* sp. constitute 26.8%, 23.2% and 11.5% for the respective bacterial isolates (Fig 1).

For the isolation and identification of staphylococcus aureus from stethoscope samples that had previously grown in blood agar and tryptone soya agar, mannitol salt agar (MSA) was utilized as a selective and differential medium. This media was selective for staphylococcus aureus which ferment mannitol and produce yellow colonies with yellow zone around the colony; non mannitol fermented bacteria remain red to pink and colorless in the medium (Figure 3). The majority of the isolates (38.5%)

were show yellow zone in the medium and identified as S. aureus.

The action of bacterial hemolytic exotoxins on red blood cells was used to identify normal flora from pathogenic bacteria using a blood agar plate (BAP) as a bacterial growth medium. The isolates were described and identified as streptococcus species based on their hemolytic (Alpha hemolysis, Beta hemolysis, and Gamma hemolysis) patterns (Figure–4). Streptococcus pyrogenes was identified as the bacterium that caused beta hemolysis on blood agar. On BAP, alpha hemolysis indicated the growth of normal flora, while gamma hemolysis suggested that the growth on BAP had no effect on the agar's appearance. Streptococcus pyrogenes (13.2%) was identified from the isolates and streptococcus pneumonia (10%) as normal flora as it showed gamma hemolysis in BAP.

Table 4 shows bacterial colony counts by gender, profession, experience and ward. The mean colony count of different wards was 109, where the highest (220) and lowest (36) was recorded in Delivery and Emergency ward, respectively. The data also showed difference between female (151) and male (67) mean colony count, which showed a significant difference at P<0.05. Nurses had the greatest mean colony count (148), while doctors had the lowest (79). In contrast, a non–significant mean difference was found in respect to health care personnel's years of experience.



# **Figure 1** – Bacterial profile isolated from stethoscope diaphragm.

Staphylococcus aureus
 Coagulase negative staphylococci
 Streptococcus sp.
 Bacillus sp.



**Figure 2** – Image of bacterial isolates that grown in mannitol salt agar.

Staphylococcus aureus

Staphylococcus epidermis

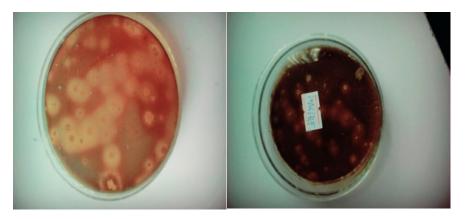


Figure 3 – Image of hemolytic streptococcus grown in blood agar.

Streptococcus pyrogenes

Streptococcus pneumonia

**Table 4** – Bacterial colony counts from culture of stethoscope diaphragm surface.

		<b>MEAN BACTERIAL COUNT</b>
GENDER	Male	67.7
GENDER	Female	150.8
	Doctor	79.2
PROFFESSION	Nurse	148.6
PROFFESSION	Student	112.6
	Но	111.3
	<2Yrs	111.1
EXPERIANCE	2–5Yrs	111.7
EXPERIANCE	5–10Yrs	74.3
	>10Yrs	130
	Medical	135.6
	Gynecology	106.7
	Surgical	140
WARD	Pediatrics	36.7
	Emergency	22.5000
	Neonatal	40
	Delivery	220

# Patterns of antimicrobial susceptibility in isolates

The isolates' antibiotic sensitivity pattern was examined for the following antibiotic discs: co-trimoxazole, chloramphenicol, penicillin, vancomycin, ciprofloxacin penicillin. Both S. aureus and CoNS were sensitive to the chloramphenicol; Streptococcus was intermediate while bacillus was resistant to the chloramphenicol (**Table 5 & Figure-4**).

 Table 5 – Bacterial isolates from stethoscopes were tested for antimicrobial sensitivity.

ANTIBIOTIC DISCS	TYPE OF ISOLATES				
	S. aureus	CoNS	Streptococcus sp.	Bacillus sp.	
Co-trimoxazole (25 µg)	S	S	S	S	
Vancomycin (30 µg)	I	I	I	I	
Cifoxitine (30 µg)	R	R	R	R	
Penicillin (10 µg)	R	R	I	R	
Ciprofloxacin (5 µg)	S	S	S	S	
Chloramphenicol (30µg)	S	S	I	R	

Key:R=Resistant; S=Susceptible; I=Intermediate

ANTIBIOTIC DISCS	TYPE OF ISOLATES WITH THEIR DIAMETER(CM)				
	S. aureus	CoNS	Streptococcus sp.	Bacillus sp.	
Co-trimoxazole (25 µg)	2.4	3.1	2.5	3.2	
Penicillin (10 µg)	1.3	1.5	1.7	1.3	
Cifoxitine (30 µg)	1.2	1.5	1.3	1.4	
Vancomycin (30 µg)	1.67	1.7	1.7	1.9	
Ciprofloxacin (5 µg)	3.5	2.9	2.8	3.2	
Chloramphenicol (30µg)	3.6	3.00	1.7	1.3	

# Figure 4 – Antimicrobial sensitivity test result (inhibition zone and diameter measurement).



# Discussion

The stethoscope is a piece of medical equipment that is utilized by all health–care workers. Our research indicated that 69.2 percent of the stethoscopes surveyed were infected, which is similar to previous findings that found 71 percent to 100 percent of stethoscopes were colonized by different bacteria<sup>[7, 29].</sup>

Doctors' and nurses' stethoscopes were found to be more polluted (33 percent apiece) than those used by other health workers. This research is comparable to that of [7] The fact that doctors and nurses use stethoscopes more frequently than other health care staff may explain the higher rate of bacterial colonization, even if the difference was not statistically significant <sup>[19, 7]</sup>. Nurses, on the other hand, had a greater mean microbial load (149) and medical students (113) and the lowest were recorded in doctors (79), which might be due to improved stethoscope cleaning habits in later case.

In this investigation, a swab of stethoscopes taken from clinicians in the medical ward (8) revealed the highest level of infection. Medical physicians may wear stethoscopes more frequently than others, which could explain why they have a greater prevalence of bacterial contamination.

A total of 25 colonies were isolated from 18 (69.2%) contaminated stethoscope diaphragms, although only 13 unique bacterial isolates were examined for further phenotypic characterization. Surprisingly, no gramnegative bacteria were found in any of the stethoscope diaphragms that were analyzed. Gram-positive bacteria were found in all of the isolates.

For bacterial growth and enumeration, blood agar, MacConkey agar, and tryptone soya agar media were utilized. Gram positive bacteria from four different species were recovered from both blood and tryptone soya agar. The largest number of bacterial isolates per diaphragm was three, while the lowest number was one. The majority of the isolates (40%) were identified to be potential pathogens. Staphylococcus aeurus species was the most prevalent isolate (38.5%), followed by coagulase negative staphylococci. although Staphylococcus epidermis and enterobacteracea were the most common organisms recovered from stethoscopes in investigations bv<sup>[7]</sup>. Co-trimoxazole and ciprofloxacine were determined to be effective against all gram-positive isolates based on their resistance profiles. Meanwhile cifoxitine and penicillin were not. That is all isolates were susceptible to the co-trimoxazole and ciprofloxacine, while resistant to cifoxitine. They showed intermediate growth against vancomycine. All except streptococcus were found resistant against penicillin. Both S. aureus and CoNS were sensitive to the chloramphenicol; Streptococcus was intermediate while bacillus was resistant to the chloramphenicol.

In this study, a questionnaire was used to analyze

knowledge, attitudes, participants' and practices regarding the role of stethoscopes as carriers of infectious organisms. We found that stethoscopes were contaminated with dangerous germs and that inadequate stethoscope cleaning/disinfection techniques were related with high contamination. In particular,34 stethoscopes cleaned on the same day as the data collection were uncontaminated, compared to 100 percent contamination in those who said they never/cannot recall. Because even brief contact with a patient's skin and the stethoscope can result in bacterial translocation<sup>[1]</sup>, measures to reduce bacterial contamination through better stethoscope cleaning habits are needed.

Disposable stethoscopes, especially in clinical high-risk contexts, and the placement of a single-use silicone membrane over the stethoscope head to provide a prophylactic barrier have been proposed as ways to reduce infection transmission from stethoscopes<sup>[23]</sup>. Although these measures could reduce the risk of infection transmission via stethoscope, they are out of reach for the majority of health workers and health facilities in developing nations, including Attat Hospital. Instead, hospitals should implement more stringent stethoscope disinfection programs and practices as a standard of care [28]. Health personnel who strictly follow stethoscope disinfection procedures will reduce cross-contamination and increase patient safety in hospitals.

### Conclusion

The current study found a higher percentage of bacterial contamination on the stethoscope diaphragm, indicating that there is a risk of nosocomial pathogen transmission. Many of the bacteria found in our study's stethoscopes (e.g., Staphylococcus aureus, CoNs; S.epidermis, *Streptococcus sp.; S.pyrogene*, and *Bacillus subtilis*) were known to cause serious infections in hospitalized patients. Staphylococcus and Bacillus species showed increased resistance to the drugs tested, however Streptococcus species did not. Infected stethoscopes were discovered in all parts of the hospital and among all types of medical professionals. The study also suggests that hospital employees should be alerted and educated about the potential health concerns linked with medical equipment.

To reduce infection transmission through stethoscopes, various techniques have been proposed, including the use of disposable stethoscopes, particularly in clinical high-risk areas, and the placement of a single-use silicone membrane over the stethoscope head to establish a prophylactic barrier. Although these measures could reduce the risk of infection transmission via stethoscope, they are out of reach for the majority of health workers and health facilities in developing nations, including Attat Hospital. Instead, hospitals should implement more stringent stethoscope disinfection programs and practices as a standard of care. Health personnel who strictly follow stethoscope disinfection procedures will reduce cross-contamination and increase patient safety in hospitals. Training and motivating health care providers to put their knowledge into practice could be the next step in drastically lowering the bacterial load from the stethoscope, which would immediately reduce cross-contamination and improve patient safety in the hospital setting.

# **The Study's Limitations**

The sample size was tiny (26 people), and it came from only one hospital. The frequency with which the stethoscopes were used differed from one participant to the next. In this investigation, the colonization of stethoscopes was not linked to hospital–acquired illnesses. Other contaminants such as anaerobic bacteria, fungi, and viruses were not investigated. The length of time the stethoscope was in touch with the patient's skin/ clothing was unknown. Bacterial identification was done using phenotypic characterization, which is not as reliable as molecular approaches. The identification of such contaminating organisms and their role as nosocomial infections should be the focus of future research.

# Recommendation

- The bacterial four isolates types of microbes need to be fully characterized using molecular techniques.
- Further study is important to identify other microbes from large enough sample size of different wards with their drug sensitivity tests is needed.
- Design instrument processing of stethoscope like other health service instruments
- There is a need of training for health personnel to increase the culture of decontaminations of their stethoscope.

# References

- [1] Adetunji, C., Makanjuola, O., Lateef, A., Oloke, J., Arowora, K., Adetunji, J., Ajani, A., Africa–Purino, F., Dy, E. and Coronel, R. (2001). Stethoscopes: a po– tential source of nosocomial infections. *Phil. J.Mi– crobiol. Infect. Dis.* **29**: 9–13.
- [2] Al-hamad A, Maxwell S. 2010 How clean is clean? Proposed methods for hospital cleaning assessment. J Hosp Infect; 70: 328–33
- [3] Alothman A, Bukhari A, Aljohani S, Muhanaa A. 2009 Should we recommend stethoscope disinfection before daily usage as an infection control rule? The Open Infectious Diseases Journal; 3: 80–2.
- [4] Aneja, K. (2003). Experiments in Microbiology, Plant Physiology and Biotechnology, 4<sup>th</sup>edn. New Age International, New Delhi.

- [5] Aslanzadeh, J. (2006). Biochemical profile-based microbial identification systems. In: Advanced Techniques in Diagnostic Microbiology, pp. 84–116, (Tang, Y. and Stratton, C., eds). Springer Science+Business Media, New York.
- [6] Atlas, R. (2010). Handbook of Microbiological Media, 4<sup>th</sup>edn. CRC Press Taylor & Francis Group, Boca Raton.
- [7] Chigozie, J., Annayo, O., Patrick, G. and Christian, M. (2010). Bacteria contamination of stethoscopes used by health workers: public health implications. *J. Infect. Dev.Ctries.*;**4**:436–41.
- [8] Cheesbrough M. (2006). District Laboratory Practice in Tropical Countries, Part 2, Cambridge University Press, United Kingdom, PP 60–64.
- [9] Fenelon, L., Holcroft, L. and Waters, N. (2009). Contamination of stethoscopes with MRSA and current disinfection practices. J. Hosp. Infect. 71: 376–378.
- [10] Gastmeier, P., Groneberg, K., Weist, K. and Rüden, H. (2003). A cluster of nosocomial *Klebsiella pneumonia* bloodstream infections in a neonatal intensive care department: Identification of transmission and intervention. *Am. J. Infect. Contr.***3**: 424–430.
- [11] Gregorson, G. (1978). Rapid method for distinction of gram negative from gram positive bacteria. *Euro– pean J. Appl. Microbiol.***5**: 123–127.
- [12] Gupta, A., Della–Latta, P., Todd, B., San Gabriel, P., Haas, J., Wu, F., Rubenstein, D. andSaiman, L. (2004). Outbreak ofextendedspectrum beta–lacta– mase–producing *Klebsiella pneumonia* in a neona– tal intensive care unit linked to artificial nails. *Infect. Contr. Hosp. Epidemiol.***25**: 210–215.
- [13] Harisha, S. (2007). *Biotechnology Procedures and Experiments Handbook*. Infinity Science Press, Hin-gham.
- [14] Harley, J. and Prescott, L.(2002). *Laboratory Exercise in Microbiology*,5<sup>th</sup>edn. The McGraw–Hill Companies, 466p.
- [15] Holt, J.G., Krieg, N.R., Sneath, P.H.A. and Staley, J.T. (1994). Bergey's Manual of Determinative Bacterio– logy, 9thedn. Williams and Wilkins company, Balti– more, MD, USA, pp: 255–273.
- [16] Kerr, J.R., Martin, H., Chadwick, M.V., Edwards, A., Hodson, M.E. and Geddes, D.M. (2002). Evidence against transmission of *Pseudomonas aeruginosaby*

hands and stethoscopes in a cystic fibrosis unit. *J. Hosp. Infect.***50**: 324–326.

- [17] Lahsaeizadeh S, Jafari H, Askarian M. Health care associated infection in Shiraz, Iran 2004–2005. J Hosp Infect. 2009; 69:283–7.
- [18] Madar, R., Novakova, E. and Baska, T. (2005). The role of noncritical health-care tools in the transmission of nosocomial infections. *Bratisl.Lek. Listy.* 106: 348– 350.
- [19] Marinella MA, Pierson C, Chenoweth C1997. The Stethoscope– a potential source of nosocomial in– fection? Arch Intern Med; **157:786**–70.
- [20] Merlin, M.A., Wong, M.L., Pryor, P.W., Rynn, K., MarquesBaptista, A., Perritt, R., Stanescu, C.G. and Fallon, T. (2009). Prevalence of methicillin–re– sistant *Staphylococcus aureus* on the stethoscopes of emergency medical services providers. *Prehosp. Emerg. Care.***13**: 71–74.
- [21] Orrett, F.A., Brooks, P.J. and Richardson, E.G. (1998). Nosocomial infections in a rural regional hospital in a developing country: infection rates by site, service, cost, and infection control practices. *Infect. Contr. Hosp. Epidemiol.* **19**: 136–140.
- [22] Parmar, R.C., Valvi, C.C., Sira, P. andKamat, J.R. (2004). A prospective, randomised, double-blind study of comparative efficacy of immediate versus daily cleaning of stethoscope using 66% ethyl alcohol. *Indian J. Med. Sci.* 58: 423–430.
- [23] PatentStorm (2004) Disposable cover for stethoscope head. Available: http://www.freepatentsonline. com/5747751.html. Accessed 15 October 2009.
- [24] Rahman L, and Anson KR. (2004). Bacterial contamination of hospital physicians' stethoscopes. Infect. Control Hospital Epidemiol. **20**(9): 626–628.
- [25] Raka L, Zoutman D, Mulliqi G, Krasniqi S, Dedushaj I, Raka N, Ahmeti S, Shala M, VishajA, Elezi Y. Prevalence of nosocomial infections in high-risk units in the University Clinical Center of Kosovo. Infect ContrHospEpidemiol 2006; 27: 421–423.
- [26] Saloojee, H.andSteenhoff, A. (2001). The health professional's role in preventing nosocomial infections. *Postgrad. Med. J.* 77: 16–19.
- [27] Schroeder, A., Schroeder, M.A. and D'Amico, F. (2009).What's growing on your stethoscope? (and what you can do about it). *J. Fam. Pract.***58**: 404–

409.

- [28] Sengupta S, Sirkar A, Shivananda PG (2000) Stethoscopes and nosocomial infection. Indian J Pediatr 67: 197–199.
- [29] Shiferaw, T., Beyene, G., Kassa, T. and Sewunet, T. (2013). Bacterial contamination, bacterial profile and antimicrobial susceptibility pattern of isolates from stethoscopes at Jimma University Specialized Hospital. Annals of Clinical Microbiology and Antimicrobials 12:39.
- [30] Singh G ,. Urhekar A.D, Hodiwala A. V, Singh N , Das B. 2013 Bacterial contamination of stethoscopes used by health care workers in a tertiary care hospital in Navi Mumbai. IJPBS ;3(1) 186–193
- [31] Uneke, C.J., Ogbonna, A., Oyibo, P.G. and Ekuma, U. (2008). Bacteriological assessment of stethoscopes used by medical students in Nigeria: Implications for nosocomial infection control. *World Health Popul.* **10**: 53–61.
- [32] Vincent, J.L. (2003). Nosocomial infections in adult intensive care units.*Lancet***361**: 2068–2077.
- [33] Weinstein RL, Restrepo RD, Bourne KC, Daher N. 2005. Contamination level of stethoscopes used by physicians and physicians Assistants. The J of Physician Assistant Edu. 18:41–3.
- [34] WHO (2002). Prevention of Hospital–Acquired In– fections: A Practical Guide. Malta: Department of Communicable Disease, Surveillance and Response.
- [35] World Health Organization (2009) WHO Guidelines for Hand Hygiene in Health Care. First Global Patient Safety Challenge Clean Care is Safer Care. Geneva: WHO, 270p
- [36] WHO (2009). Save Lives Clean Your Hands–Guide to Implementation. A Guide to the Implementation of the WHO Multimodal Hand Hygiene Improvement Strategy WHO/IER/PSP/2009.02. Geneva: WHO 48p.
- [37] Yemane, T. (1967). Statstics: An Introductory Analysis, 2<sup>nd</sup>edn. New York: Harper and Row, S.A. (2002). Stethoscope: a friend or an enemy? Sao Paulo Med. J.120: 13 15.

### Salile