

# Bacteriological Profile of Wound Sepsis and Antimicrobial Pattern of Isolates at Federal Medical Centre, Bida, Niger State. Nigeria.

## Abstract

**Aim:** The study was aimed to identify etiology of bacteria associated with wound infections and antimicrobial susceptibility profile of the isolated organisms in the community.

**Study design and Methodologys:** It is a retrospective study; data were obtained from Medical Microbiology department register from May 2005 through October 2007 and was exempted from ethical approval. Swab samples were collected from 408 patients between age groups 0 through 75years from out patients and inpatients admitted in the wards for various injuries such as burns, post surgical wound, fracture and ulcer wound. Samples were culture within 1hour on macConkey agar, blood agar and chocolate agar, and incubated at 37° c for 18-24hours overnight. Data were coded and computed using SPSS 16.0 and p-value 0.05 was considered statistical significant. **Results:** Out of 408 swab samples, 338 (82.8%) yielded positive culture, overall highest isolates was found within age groups 31-40years with 69(94.5%) growth followed by 21-30years 61(85.9%) and the least growth was found in 51-60years 27(77.1%) and 0-10years 88(77.2%), and statistically not significant ( p-value 0.814, mean age =11.34, median =12.00, mode =12 and S.D±4.361). The highest single isolates was *Staphylococcus aureus* 122(42.5%) followed by *Escherichia coli* 108(37.6%), *Pseudomonas aeruginosa* 28(9.8%), *Proteus* species 15(5.2%) and lowest isolates were *Candida albicans* 3(1.0%), *Clostridium* species 2(0.7%), Coagulase negative *Staphylococcus* 2(0.7%) and *Streptococcus* species 2(0.7%).

*Escherichia coli* and *Staphylococcus aureus* had the most prevalent polymicrobial isolates with 28(54.9%) followed by *Escherichia coli* and *Proteus* species 8(15.7%).

*Staphylococcus aureus* the highest prevalent single isolates was susceptible to Ceftriazone 75(61.5%), Ciprofloxacin 71(58.2%), Ofloxacin 68(55.7%) and Clindamycin 83(68.0%).

**Conclusion:** The incidence rate of wound sepsis in the studied population is 338(82.88%) with incriminating single isolate of *Staphylococcus aureus* 122(42.5%). This is a serious burden to our patients which call for serious attention among stake holders.

**Recommendation:** Stake holders need to educate patients visiting hospital community on the danger of wound sepsis, and first aid treatment before visiting tertiary health care to reduce morbidity and mortality incidence rate.

Keywords: Sepsis, Infection, Susceptibility, Antimicrobial.

## Introduction

Chronic wound infection occurs in individual with an increased risk of bacteria invasion as a result of poor local factors such as arterial insufficiency, veinous hypertension, trauma and systemic disease like diabetic mellitus and rheumatoid arthritis(Falanga, 1993).<sup>1</sup>

Wound infection is important in the morbidity and mortality of patients irrespective of its cause; its delay healing and is associated with prolong hospital stay thereby increasing cost of

healthcare services (Lateef *et al*, 2003)<sup>2</sup>. It may occur as a result of exposure of subcutaneous tissue following a loss of skin integrity; wound provides a warm, moist, and nutritious environment that is favorable for microbial colonization and proliferation.

Wound colonization is most frequently poly-microbial, involving numerous microorganisms that are potentially pathogenic, wounds are at risk of becoming infected (Dai *et al*, 2010)<sup>3</sup>. In western world, studies on wound infections are focused on surgical sites infections because other types of wound infections are not problematic (Gaynes *et al*, 2001)<sup>4</sup> while in developing countries such as Africa continent, other types of wound infections are major causes of morbidity and mortality among the patients (Melta *et al*, 2007, and Anguzu, and Ohila, 2007)<sup>5,6</sup>. The incidence rate of different bacterial infected wounds varies, it exists inter-institutionally and intra institutionally (Fadeyi *et al*, 2008).<sup>7</sup> Bacterial infections in burn and wound patients are similar and are difficult to control (Armour *et al*, 2007)<sup>8</sup>. Wound infection constitutes major barrier to healing and have an adverse effect on the patient's quality of life as well as on the healing rate of the wound.

Infected wounds are likely to be more painful, hypersensitive and odorous, resulting in increased discomfort and inconvenience for the patient (Kotz *et al*, 2009)<sup>9</sup>. The prevalent organisms associated with wound infection include *Staphylococcus aureus* which account for 20-40% and *Pseudomonas aeruginosa* 5-15% of the nosocomial infection, with infection mainly following surgery and burns. Other pathogens such as Enterococci and members of the Enterobacteriaceae have been implicated, among immuno-compromised patients and following abdominal surgery (Taiwo *et al*, 2002)<sup>10</sup>. Also, Godebo *et al*, (2013)<sup>11</sup> and Mulu *et al*, (2006)<sup>12</sup> stated that *Staphylococcus aureus*, *Kelbsiella* species, *Escherichia coli*, *Proteus* species, *Streptococcus* species, Enterobacter species, *Pseudomonas* species and Coagulase negative *Staphylococci* were common pathogens in wound infection.

In addition, Arturson , (1985)<sup>13</sup> said infection causes 50% to 60% of deaths in burn patients in spite of intensive therapy with antibiotics both topically as well as intravenous, and wound can be infected by a variety of microorganisms ranging from bacteria to fungi and parasites (Bowler *et al*, 2001)<sup>14</sup>. Post-surgical wound infections are hospital acquired and vary from one geographical area to the other (Isibor *et al*, 2008)<sup>15</sup>. The emergence of high anti-microbial resistance among bacterial pathogens made the treatment of post-operative wound infections challenging (Andhoga *et al*, 2002)<sup>16</sup>. The situation is serious in developing countries due to irrational prescriptions of antimicrobial agents (Fadeyi *et al*, 2008)<sup>17</sup>. The emergence of drug resistant pathogens like Methicillin Resistant *Staphylococcus aureus* (MRSA) and Extended Spectrum Beta Lactamase (ESBL) leading to treatment failure (2011)<sup>18</sup>. The study was aimed to identify etiology of bacteria associated with wound sepsis and antimicrobial susceptibility profile of the isolated organisms in the community.

## Materials and Methods

### Study population

The research was a retrospective study; data were collated from May, 2005 through October 2007 from Medical Microbiology department register and exempted from ethical approval. Swab samples of four hundred and eight (408); female 191 and male 217 swab specimen were collected aseptically from different categories of patients both out-patient and in-patients from various wound site such as burns, ulcer, post operative wound and fracture wound, submitted to Medical Microbiology department for routine analysis. Subjects were between age groups 0 through 75years old.

### Analysis, Characterization and Identification of Bacteria from swab Samples

Swab samples were submitted for routine, gram stain, culture and sensitivity. Samples were culture within 1hour of submission on MacConkey agar, Blood agar and Chocolate agar

according to Chessbrough<sup>19</sup>. Samples were further gram stain directly to classify staining reaction (Cheesbrough, 2002)<sup>19</sup>. The bacterial isolates were characterized based on colonial morphology, growth on selective media and enriched media, and biochemical tests which include Gram's reaction, indole tests, methyl red, voges-proskauer, citrate utilization, motility, endospore, utilization of carbohydrates such as glucose, sucrose, mannitol, lactose and fructose, oxidase, catalase, coagulase and starch hydrolysis test (Oyeleke and Manga, 2008)<sup>20</sup>. Antimicrobial susceptibility test by disc diffusion methods according to clinical laboratory standard guidelines (Cheesbrough, 1991).<sup>21</sup> The antimicrobial disc used include Clindamycin (5mcg), Streptomycin(10mcg), Gentamycin (10mcg), Ceftriazone (30mcg), Erythromycin (5mcg), Ofloxacin (5mcg), Augmentin (30mcg), Ciprofloxacin (5mcg), Ampicillin (10mcg), Tetracycline (5mcg), Cotrimoxazole (10mcg), Azythromycin (30mcg) and Pefloxacin (5mcg). Susceptibility to antibiotics was measured by the method of Baker and Breach (Baker and Breach 1980)<sup>22</sup>. When the antibiotic agent was 16mm or higher, it was recorded susceptible, and resistance when less than 16mm. The susceptibility plates were incubated aerobically for 18-24hrs and zone of inhibition were recorded. Data was coded, computed and analyzed using SPSS version 16.0 and p values  $\leq 0.05$  was considered to be statistically significant.

**Results:** Table 1, showed four hundred and eight (408) patients enrolled; a total of three hundred and thirty eight 338(82.8%) yielded significant growth of isolates, and 70(17.2%) had sterile culture. Out of 338 (82.8%) positive culture, overall highest positive culture was found within age groups 31- 40years with 69(94.5%) growth followed by 21-30years 61(85.9%) and the least growth was found in 0-10years 88(77.2%) and 51- 60years 27(77.1%).

Table 2, showed the frequency of isolates in relation to age. Our research showed two categories of isolates, single pure isolates 287(84.9%) table 2A, and mixed growth isolates 51(15.1%) table 2B.

Table 2A: The highest single isolates was *Staphylococcus aureus* 122(42.5%) followed by *Escherichia coli* 108(37.6%), *Pseudomonas aeruginosa* 28(9.8%), *Proteus* species 15(5.2%) and lowest isolates were *Candida albicans* 3(1.0%), *Clostridium* species 2(0.7%), Coagulase negative *Staphylococcus* 2(0.7%) and *Streptococcus* species 2(0.7%). Also, a higher occurrence of single isolates was found within age groups 0-10years with 77(26.8%) followed by 31-40years 42(19.9%) and lowest isolates was in 61-70years 7(2.4%).

Table 2B was a polymicrobial isolates; overall highest isolates was in age groups 0-10years with 13(25.5%) followed by 31-40years 11(21.6%) and least isolates was 61-70years 2(3.9%). *Escherichia coli* and *Staphylococcus aureus* 28(54.9%) had highest mixed isolates followed by *Escherichia coli* and *Proteus* species 8(15.7%) and least isolates *Escherichia coli* and *Corynebacterium diptheriae* 1(2%), and *Proteus* species and *Klebsiella* species 1(2%).

Table 3A showed antimicrobial susceptibility pattern of the isolates; *Staphylococcus aureus* the highest prevalent isolate was susceptible to Ceftriazone 75(61.5%), Ciprofloxacin 71(58.2%), Ofloxacin 68(55.7%) and Clindamycin 83(68.0%), and least susceptible was Augmentin 5(4.1%) and Ampicillin 1(0.8%).

Second isolate *Escherichia coli* were susceptible to Ceftriazone 64(59.3%), Ciprofloxacin 59(54.6%) and Ofloxacin 55(50.9%) and least susceptible to Ampicillin 1(0.9%) and Augmentin 4(3.7%). *Pseudomonas aeruginosa* was susceptible to Ciprofloxacin 17(60.7%), Ofloxacin 15 (53.6%) and Ceftriazone 15(53.6%) and least susceptible to Cotrimoxazole 1(3.6%) and Azithromycin 3(10.7%)

Table 3B depict the antimicrobial activities of mixed isolates; the most prevalence was *Escherichia coli* and *Staphylococcus aureus* with susceptibility to Ciprofloxacin 16(57.1%), Ofloxacin 11(39.3%) and lowest susceptible to Augmentin 1(3.6%) and Tetracycline 1(3.6%).

**Table 1**

Frequency of Subjects in Relation to Age Showing Positive and Negative culture

| Age   | Number of Subjects | Positive Subjects | Negative Subjects |
|-------|--------------------|-------------------|-------------------|
| 0-10  | 114                | 88 (77.2%)        | 26(22.8%)         |
| 11-20 | 44                 | 35(79.4%)         | 9(20.5%)          |
| 21-30 | 71                 | 61(85.9%)         | 10(14.1%)         |
| 31-40 | 73                 | 69 (94.5%)        | 4(5.5%)           |
| 41-50 | 60                 | 50(83.3%)         | 10(16.7%)         |
| 51-60 | 35                 | 27(77.1%)         | 8(22.9%)          |
| 61-70 | 10                 | 8(80%)            | 2(20%)            |
| 71-75 | 1                  | 0                 | 1(100%)           |
| Total | 408(100%)          | 338(82.8%)        | 70(17.2%)         |

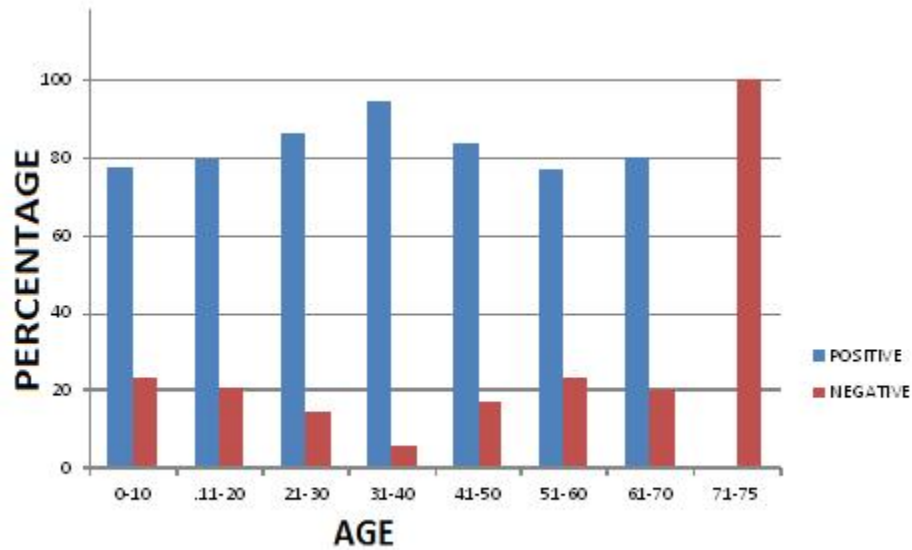
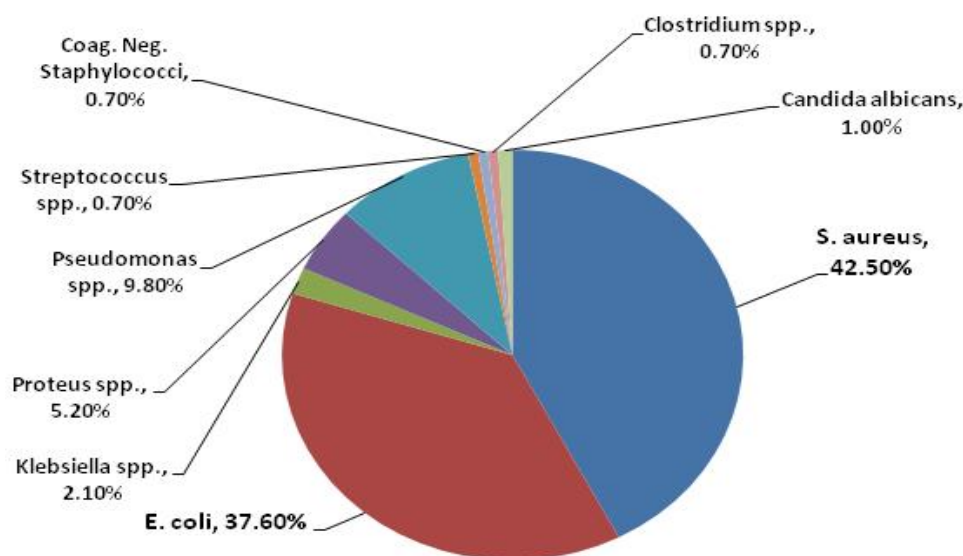


FIGURE 1: Chart Showing Frequency of Subjects In Relation To Age of Positive and Negative Culture

**Table 2A**

Incidence Rate of Single Isolates in Relation to Age Distribution of Subjects  
with Wound Infections.

| Isolates                                       | 0-10      | 11-20     | 21-30     | 31-40     | 41-50     | 51-60    | 61-70   | 71-80 | Total      |
|--|-----------|-----------|-----------|-----------|-----------|----------|---------|-------|------------|
| <i>S. aureus</i>                               | 36(29.5%) | 18(14.8%) | 20(16.4%) | 19(15.6%) | 15(12.3%) | 12(9.8%) | 2(1.6%) | 0(-)  | 122(42.5%) |
| <i>E. coli</i>                                 | 18(16.7%) | 11(10.2%) | 24(22.2%) | 25(23.1%) | 18(16.7%) | 7(6.5%)  | 5(4.6%) | 0(-)  | 108(37.6%) |
| <i>Klebsiella</i><br>species                   | 1(16.7%)  | 0(-)      | 1(16.7%)  | 3(50%)    | 1(16.7%)  | 0(-)     | 0(-)    | 0(-)  | 6(2.1%)    |
| <i>Proteus</i><br>species                      | 3(20%)    | 0(-)      | 4(26.7%)  | 5(33.3%)  | 3(20%)    | 0(-)     | 0(-)    | 0(-)  | 15(5.2%)   |
| <i>Pseudomonas</i><br><i>aeruginosa</i>        | 14(50%)   | 1(3.6%)   | 1(3.6%)   | 4(14.3%)  | 4(14.3%)  | 4(14.3%) | 0(-)    | 0(-)  | 28(9.8%)   |
| <i>Streptococcus</i><br>species                | 0(-)      | 0(-)      | 0(-)      | 1(50%)    | 1(50%)    | 0(-)     | 0(-)    | 0(-)  | 2(0.7%)    |
| <i>Coagulase neg.</i><br><i>Staphylococcus</i> | 1(50%)    | 0(-)      | 1(50%)    | 0(-)      | 0(-)      | 0(-)     | 0(-)    | 0(-)  | 2(0.7%)    |
| <i>Clostridium</i><br>Species                  | 1(100%)   | 0(-)      | 0(-)      | 0(-)      | 0(-)      | 0(-)     | 0(-)    | 0(-)  | 2(0.7%)    |
| <i>Candida</i><br><i>albicans</i>              | 3(100%)   | 0(-)      | 0(-)      | 0(-)      | 0(-)      | 0(-)     | 0(-)    | 0(-)  | 3(1.0%)    |
| Total  | 77(26.8%) | 30(10.5%) | 51(17.8%) | 57(19.9%) | 42(14.6%) | 23(8.0%) | 7(2.4%) | 0(-)  | 287(100%)  |



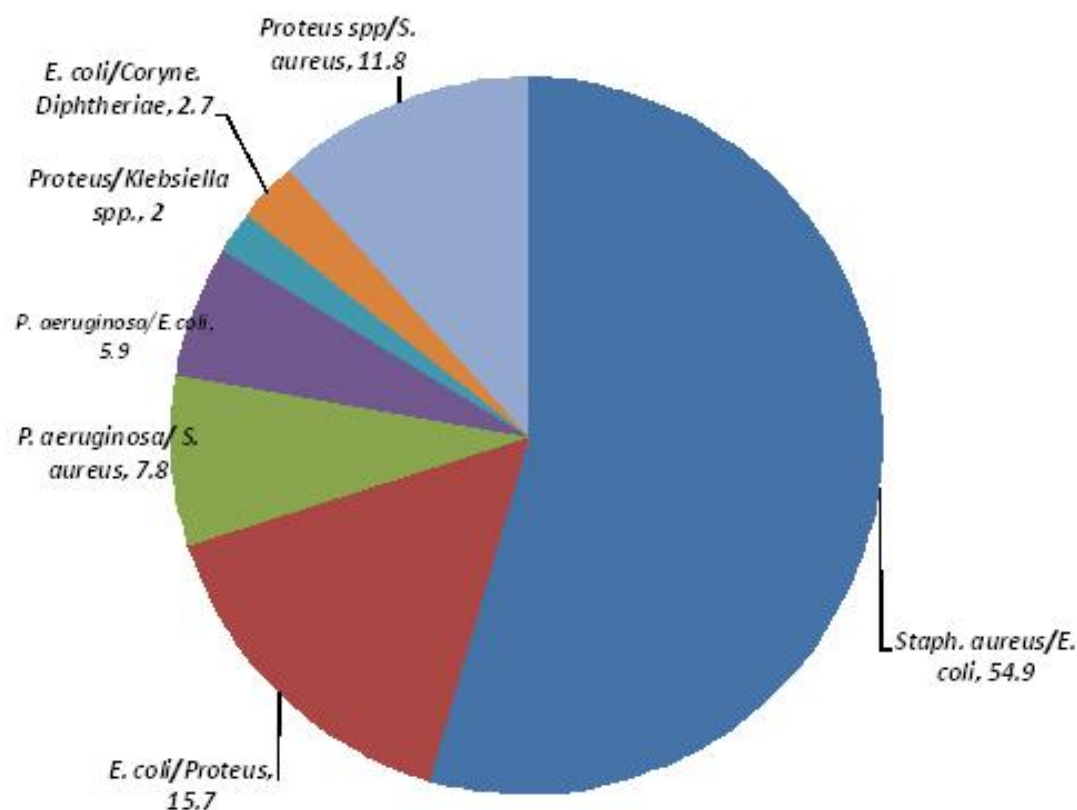
**Fig. 2A: Depicts Percentage Frequency of Isolated Organisms.**

**Table 2B**

Incidence Rate of Mixed Isolates in Relation to Age Distribution of Patients With Wound Infections.

| Isolates  | 0-10      | 11-20    | 21-30     | 31-40     | 41-50    | 51-60    | 61-70    | 71-75 | Total     |
|---|-----------|----------|-----------|-----------|----------|----------|----------|-------|-----------|
| <i>S. aureus</i> & <i>E. coli</i>               | 5(17.9%)  | 3(10.7%) | 9(32.1%)  | 5(17.9%)  | 3(10.7%) | 2(7.1%)  | 1(3.6%)  | 0(-)  | 28(54.9%) |
| <i>E. coli</i> & <i>Proteus spp.</i>            | 3(37.5%)  | 1(12.5%) | 0(-)      | 3(37.5%)  | 1(12.5%) | 0(-)     | 0(-)     | 0(-)  | 8(15.7%)  |
| <i>P. aeruginosa</i> & <i>S. aureus</i>         | 2(50%)    | 0(-)     | 0(-)      | 1(25%)    | 1(25%)   | 0(-)     | 0(-)     | 0(-)  | 4(7.8%)   |
| <i>P.aeruginosa</i> & <i>E. coli</i>            | 1(33.3%)  | 0(-)     | 0(-)      | 1(33.3%)  | 0(-)     | 0(-)     | 1(33.3%) | 0(-)  | 3(5.9%)   |
| <i>Proteus species</i> & <i>Klebsiella spp.</i> | 1(100%)   | 0(-)     | 0(-)      | 0(-)      | 0(-)     | 0(-)     | 0(-)     | 0(-)  | 1(2.0%)   |
| <i>E. coli</i> & <i>Coryne. Diphtheria</i>      | 0(-)      | 0(-)     | 1(100%)   | 0(-)      | 0(-)     | 0(-)     | 0(-)     | 0(-)  | 1(2.0%)   |
| <i>Proteus spp.</i> & <i>S.aureus</i>           | 1(16.7%)  | 1(16.7%) | 0(-)      | 1(16.7%)  | 2(33.3%) | 1(16.7%) | 0(-)     | 0(-)  | 6(11.8%)  |
| Total   | 13(25.5%) | 5(9.8%)  | 10(19.6%) | 11(21.6%) | 7(13.7%) | 3(5.9%)  | 2(3.9%)  | 0(-)  | 51(100%)  |





**Fig 2b: chart showing polymicrobial isolate**

**Table 3A**

Percentage Antimicrobial Profile of Isolated Organisms from Wound Infections

| Antibiotics   | <i>P. aeruginosa</i><br>N=28 | <i>S. aureus</i><br>N=122 | <i>E. coli</i><br>N=108 | <i>Kleb.</i><br>species<br>N=6 | <i>Proteus</i><br>species<br>N=15 | <i>Strept.</i><br>species<br>N=2 | Coag. Neg. <i>Staph.</i><br>N=2 | Clostridium<br>species<br>N=1 |
|---------------|------------------------------|---------------------------|-------------------------|--------------------------------|-----------------------------------|----------------------------------|---------------------------------|-------------------------------|
| Ampicillin    | NA                           | 1(0.8%)                   | 1(0.9%)                 | 0(-)                           | 0(-)                              | 0(-)                             | 0(-)                            | 0(-)                          |
| Erythromycin  | NA                           | 48(39.3%)                 | NA                      | NA                             | NA                                | 1(50%)                           | 1(50%)                          | 0(-)                          |
| Tetracycline  | NA                           | 45(36.9%)                 | 18(16.7%)               | 2(33.3%)                       | 0(-)                              | 1(50%)                           | 0(-)                            | 0(-)                          |
| Augmentin     | 0(-)                         | 5(4.1%)                   | 4(3.7%)                 | 2(33.3%)                       | 0(-)                              | 0(-)                             | 0(-)                            | 0(-)                          |
| Azythromycin  | 3(10.7%)                     | 64(52.5%)                 | 45(41.7%)               | 4(66.7%)                       | 3(20%)                            | 2(100%)                          | 2(100%)                         | 0(-)                          |
| Streptomycin  | 3(10.7%)                     | 33(27.0%)                 | 30(27.8%)               | 0(-)                           | 6(40%)                            | 2(100)                           | 0(-)                            | 0(-)                          |
| Gentamycin    | 12(42.9%)                    | 72(59.0%)                 | 40(37.0%)               | 2(33.3%)                       | 8(53.3%)                          | 2(100%)                          | 1(50%)                          | 0(-)                          |
| Ciprofloxacin | 17(60.7%)                    | 71(58.2%)                 | 59(54.6%)               | 3(50%)                         | 9(60%)                            | 2(100%)                          | 1(50%)                          | 0(-)                          |
| Ofloxacin     | 15(53.6%)                    | 68(55.7%)                 | 55(50.9%)               | 4(66.7%)                       | 8(53.3%)                          | 1(50%)                           | 1(50%)                          | 0(-)                          |
| Ceftriazone   | 15(53.6%)                    | 75(61.5%)                 | 64(59.3%)               | 4(66.7%)                       | 10(66.7%)                         | 2(100%)                          | 1(50%)                          | 0(-)                          |

|     |               |           |           |           |        |          |        |         |         |
|-----|---------------|-----------|-----------|-----------|--------|----------|--------|---------|---------|
| 256 | Cotrimoxazole | 1(3.6%)   | 37(30.3%) | 13(12.0%) | 0(-)   | 3(20%)   | 1(50%) | 0(-)    | 0(-)    |
| 257 | Clindamycin   | NA        | 83(68.0%) | NA        | NA     | NA       | 1(50%) | 1(50%)  | 1(100%) |
| 258 | Pefloxacin    | 15(53.6%) | 62(50.8%) | 42(38.9%) | 3(50%) | 7(46.7%) | 1(50%) | 2(100%) | 0(-)    |
| 259 |               |           |           |           |        |          |        |         |         |

**Table 3B**  
Percentage Antimicrobial Susceptibility Profile of Mix Isolates from Wound Infections.

|     | Isolates              | Amp. | Tet.    | Aug.    | Azm.    | Strep.  | Gen.     | Cip.     | Oflo.    | Cro.     | Cot.    | Pef     |        |
|-----|-----------------------|------|---------|---------|---------|---------|----------|----------|----------|----------|---------|---------|--------|
| 265 | <i>E. coli</i> &      | -    | 1(3.6)  | 1(3.6)  | 9(32.1) | 4(14.3) | 10(35.7) | 16(57.1) | 11(39.3) | 10(35.7) | 4(14.3) | 7(25)   |        |
| 266 | <i>S. aureus</i>      |      |         |         |         |         |          |          |          |          |         |         |        |
| 267 |                       |      |         |         |         |         |          |          |          |          |         |         |        |
| 268 | <i>E. coli</i> &      | -    | -       | 1(12.5) | 1(12.5) | 2(25)   | 4(50)    | 4(50)    | 6(75)    | 4(50)    | 1(12.5) | 3(37.5) |        |
| 269 | <i>Proteus</i> spp.   |      |         |         |         |         |          |          |          |          |         |         |        |
| 270 |                       |      |         |         |         |         |          |          |          |          |         |         |        |
| 271 | <i>P. aeruginosa</i>  | -    | -       | -       | 1(25)   | 2(50)   | 1(25)    | 3(75)    | 3(75)    | 2(50)    | 2(50)   | -       | 4(100) |
| 272 | & <i>S. aureus</i>    |      |         |         |         |         |          |          |          |          |         |         |        |
| 273 |                       |      |         |         |         |         |          |          |          |          |         |         |        |
| 274 | <i>S. aureus</i>      |      |         |         |         |         |          |          |          |          |         |         |        |
| 275 | & <i>Proteus</i> spp. | -    | 1(16.7) | -       | 2(33.3) | 1(16.7) | 2(33.3)  | 5(83.3)  | 5(83.3)  | 5(83.3)  | 1(16.7) | 3(50)   |        |
| 276 |                       |      |         |         |         |         |          |          |          |          |         |         |        |
| 277 | <i>P. aeruginosa</i>  | -    | -       | -       | -       | -       | 3(100)   | 2(66.7)  | 1(33.3)  | 2(66.7)  | 1(33.3) | 1(33.3) |        |
| 278 | & <i>E. coli</i>      |      |         |         |         |         |          |          |          |          |         |         |        |
| 279 |                       |      |         |         |         |         |          |          |          |          |         |         |        |
| 280 | <i>Kleb. spp.</i>     | -    | -       | -       | -       | -       | 1(100)   | 1(100)   | 1(100)   | 1(100)   | 1(100)  | 1(100)  |        |
| 281 | & <i>Proteus</i> spp. |      |         |         |         |         |          |          |          |          |         |         |        |
| 282 |                       |      |         |         |         |         |          |          |          |          |         |         |        |
| 283 | <i>E. coli</i> &      | -    | 1(100)  | -       | -       | -       | -        | 1(100)   | 1(100)   | -        | -       | 1(100)  |        |
| 284 | <i>Coryn. Diphth.</i> |      |         |         |         |         |          |          |          |          |         |         |        |

## Discussion

Wound sepsis provides a moist, warm, nutritive environment conducive for microbial colonization, proliferation, and infection (Fauci *et al*, 2008)<sup>23</sup>. Sepsis is a major cause of morbidity and mortality among burn patients and sometimes result to opportunistic infection (Cochran *et al*, 2002)<sup>24</sup>. Out of 408 studied population our research showed prevalence of (82.8%) wound infection among the patients, and (17.2%) had sterile culture, and statistically not significant (p-value =0.814, mean age =11.34, median =12.00, mode =12 and S.D±4.361). Our report is higher than Sewunet *et al*, (2013)<sup>25</sup> who reported (42%) sepsis among burn infected wound patients in Ethiopia. Also, Kyati, *et al*, (2014)<sup>26</sup> reported (67.14%) and (32.85%) isolates in gram positive and gram negative isolates among post-surgical wound infection in Index Medical College hospital, India. But our report is similar to Aynalem *et al*, (2017)<sup>27</sup> who reported incidence of (83.9%) isolates among in-patients and out-patients attending university of Gondar referral hospital, NorthWest Ethiopia. However, our report is lower than Lakshmi *et al*, (2015)<sup>28</sup> who reported (93%) burn infected wound in King Gorge hospital, India. According to 2002 survey report by the Nosocomial Infection National Surveillance Service (NINSS), which covers the period of October 1997 through



September 2001, indicates that the incidence of hospital acquired infection (HAI) related to surgical wounds is 10%. These infections complicate illness, and causes anxiety, increases patient discomfort and sometimes lead to death of our patients<sup>29</sup>.

Highest overall isolates was found within age groups 31-40years with (94.5%) isolates followed by 21-30years (85.9%). Contrarily, Mama *et al*, (2014)<sup>30</sup> reported highest isolates of (89.5%) among age groups 45-59years in Jimma university specialized hospital, South-West, Ethiopia.

Our research showed two categories of isolates in relation to age groups. Single isolates showed (84.9%) table 2A and mixed isolates (15.1%) table 2B. The highest single isolates was *Staphylococcus aureus* (42.5%) followed by *Escherichia coli* (37.6%). Our report is similar to Kyati *et al*, (2014)<sup>26</sup> who reported (58.6%), Damien *et al*, (2015)<sup>31</sup> reported (45.2%) in North Central, Nigeria and Aynalem *et al*, (2017)<sup>27</sup> reported (34%) of *Staphylococcus aureus* has the most prevalent organism. However, Sewunet *et al*,<sup>25</sup> reported Coagulase negative *Staphylococci* (42.8%) while Lakshmi *et al*, (2015),<sup>28</sup> Alharbi and Zayed (2014)<sup>32</sup> both reported *Pseudomonas* species (33.6%) and (36.14%) as the highest single isolates. Also, *Escherichia coli* and *Staphylococcus aureus* had highest mixed isolates of (54.9%) followed by *Escherichia coli* and *Proteus* species (15.7%). Mengesha *et al*, (2014)<sup>33</sup> reported multiple bacterial infections in post surgical wound infection (23.95%) with *Staphylococcus aureus* and *Proteus* species as most occurrence isolate. The high prevalence rate of enterobacterial isolates in our study could reveal faecal contamination due to poor personal hygiene<sup>(34)</sup> or due to post procedural contamination<sup>(35)</sup>.

We observed that the organisms isolated from all the wound infected patients both in-patients and out-patients were normal flora of the gastrointestinal tracts. According to Davis *et al*, (1969)<sup>36</sup> and Wormald (1970)<sup>37</sup> research, both observed that most important reservoirs for microorganisms that colonized the burn wounds of newly admitted patients are from the gastrointestinal (GI) tracts of the patients. In addition, microorganisms can be transmitted through the hands of health care workers, by fomites and hydrotherapy water (Rutala, *et al*, 1983, Sherertz and Sullivan 1985)<sup>38,39</sup> and, through the air (Rutala, *et al*, 1983)<sup>38</sup>.

Also, age groups 0-10years had the most prevalent single isolates (26.8%) while age groups 61-70years had (2.4%) least isolates. Furthermore, the highest polymicrobial isolates was within 0-10years (25.5%) followed by 31-40 years (21.6%). Gould (2009)<sup>40</sup> stated that within a community, health care acquired infections (HCAIs), can arise across a wide range of clinical conditions and affect patients of all ages. However, certain groups of patients are at an increased risk of infections including: elderly, very young, people with cancer, and other malignant diseases, people with impaired immunity, invasive devices, very ill and surgical patients.

The predominant single isolate *Staphylococcus aureus* was susceptible to Ceftriazone (61.5%), Ciprofloxacin (58.2%), Ofloxacin (55.7%), Clindamycin (68%) and least susceptible to Ampicilin (0.8%). Our report contradict Aynalem *et al*, (2017)<sup>27</sup> who reported susceptibility pattern of *staphylococcus aureus* to Ceftriazone (79.5%), Ciprofloxacin (79.4%) and Penicilin (15.4%), Lakshmi *et al*, (2015)<sup>28</sup> reported Ofloxacin (73.9%), Mama *et al*, (2014)<sup>30</sup> reported susceptibility to Ceftriazone (85.17%) and Ciprofloxacin (96%). However, our report is higher than Mengesha *et al*, (2014)<sup>33</sup> who reported susceptibility of *Staphylococcus aureus* to Ceftriazone (10%) and Nazneen *et al*, (2017)<sup>41</sup> reported Fluoroquinolones (38.47%) in post operative wound infection.

The highest polymicrobial isolates; *Staphylococcus aureus* and *Escherichia coli* were both susceptible to Ciprofloxacin (57.1%), Ofloxacin (39.3%), Ceftriazone (37.5%) and Gentamycin (35.7%), and least susceptible to Cotrimoxazole (14.3%) and Augmentin (3.6%). Our research showed polymicrobial multi-drug resistance isolates. According to W.H.O (2009)<sup>42</sup>, which stated that emergence of resistance in microorganisms is due to

indiscriminate use of antibiotics in general, and use of broad spectrum antibiotics. In addition, the spread of multidrug resistance organisms (MDROs) in health-care settings occurs mostly via health-care workers' (HCWs) contaminated hands, contaminated items, equipments and environment, often leading to outbreaks and serious infections especially in critically ill patients. Hand hygiene performance is the most important measure among standard precautions.

Enteric organisms are the predominant isolates in our research, and are ubiquitous organisms found in soil, water and vegetation, and are part of the normal intestinal flora of animals, and including humans. We suggest that hand hygiene advocate should not be limited to health care providers; it should be extended to our patients and their relations. This will help in the control of both community and hospital acquired infections. Lee *et al*, (2012)<sup>43</sup> stated in his research that good quality surveillance data on antimicrobial resistance (AMR), and the feasibility and impact of interventions based on hand hygiene promotion compliance are needed in low and middle income countries such as African continent. In addition, AMR is a cross cutting problem affecting global health care settings and our communities. The role of patients and the civil society in combating AMR is crucial at different levels and hand hygiene is one of the measures that can be practice and advocated to control the menace. Chen *et al*, (2011)<sup>44</sup> advocate increase in hand hygiene in a hospital setting in Taiwan from 43.3% to 95.6% there was 8.9% decrease in hospital acquired infections (HAIs) and a decline in blood stream infection caused by Methicillin Resistance *Staphylococcus aureus* (MRSA) and extensive drug resistance *Acinetobacter baumannii*. Al-Tawfiq *et al*, (2013)<sup>45</sup> in Saudi Arabia hospital increases hand hygiene compliance from 38% in 2006 to 83% in 2011, there was significant reduction of MRSA infection from 0.42% to 0.08% and catheter associated urinary tract infection from 7.1% to 3.5%. Also, Carboneau *et al*, (2010)<sup>46</sup> in U.S.A, advocated increase in hand hygiene from 65% to 82%, there was 51% decrease in hospital acquire MRSA cases during the 12 months period. According to Chen *et al*,<sup>44</sup> who stated that every US \$1 spent on hand hygiene promotion could result in a US \$23.7 benefit. In addition, there should be in-service training for health care providers such as post graduate training, workshop and conferences this will expose stake holders to modern facilities and equipments, research methodology and improve method of practice to foster good health care service delivery this will invariably reduce medical tourism in African continent.

In conclusion, overall prevalence rate of (82.8%) wound infection, and monomicrobial isolates of *Staphylococcus aureus* (42.5%), and polymicrobial of (15.1%) in the studied population is alarming. Policy makers need to advocate importance of hand hygiene in our communities and good sanitary disposal. This can be achieved through media in various indigenous languages, hand bills and periodic education of our patients on admission. Also, there is need to strengthen infection control units in our hospitals and government need to encourage research in health industry at all level.

Limitation: The outcome of our research is limited to sample size, there is need to carry out surveillance data of antimicrobial drug resistance, root cause and infection control in our community. This will enable policy makers to budget appropriately in terms of staff training, employments and research.

Disclaimer: - This manuscript title was presented in the conference.

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