Bacterial Contamination of Stored Blood Ready for Transfusion at a Referral Hospital in Ethiopia

Ahmed Esmael1*, Zewdu Dagnew2 and Genet Degu3

1Department of Microbiology, Immunology & Parasitology, Debremarkose University, Ethiopia
2Public health department, College of medicine and health science, Debre Markos University, Ethiopia
3Nursing department, College of medicine and health science, Debre Markos University, Ethiopia

Abstract

Background: After the discovery of Human Immunodeficiency virus, screening of blood donors practically reduced viral pathogens. However, transfusion associated bacterial sepsis, which causes high mortality and morbidity remain an important public health concern, has been received very little attention in the African set up including Ethiopia.

Objective: The aim of this study was to determine the prevalence of bacterial contamination of blood and their antibiotic susceptibility pattern at Debre Markos referral hospital, North West Ethiopia.

Methods: A facility based cross-sectional study was conducted using randomly sampled 120 whole blood units. The blood samples were obtained from screened, stored whole blood. All laboratory activities were carried out as Clinical and Laboratory Standards Institute (CLSI) protocol. Data were entered and analyzed using SPSS version 16 software. P<0.005 is statistically significant.

Results: The prevalence of bacterial contamination among stored blood was 12.5%. Gram positive bacteria (S. pneumoniae, S. aureus, coagulase-negative Staphylococcus, viridian streptococcus) and gram negative (S. typhi, E. coli and K. pneumonia) were the common isolates identified.

The isolated bacterial organisms showed varying susceptibility to the antibiotics tested. All isolated gram positive organisms were resistance to Tetracycline and susceptible to Ceftriaxone. Similarly, all the gram negative organisms isolated were resistance to Cotrimoxazole and susceptible to Ciprofloxacin and Cefoxitin.

Conclusion: In our study, we conclude the existence of this serious clinical issue (contamination of blood, development of drug resistance) and need for further surveillance and/or study.

Keywords: Bacterial contamination; Sepsis; Debre Markos referral hospital

Introduction

The field of transfusion medicine has made very rapid progress after the discovery of circulation of blood in 1628 by William Harvey. Soon afterwards the first dog to dog and subsequently, lamb to human blood transfusions were attempted. During World War II and the immediate post war period the demand for blood and blood components in the USA increased substantially. This resulted in the establishment and growth of blood banks transfusion services and other blood laboratory support services [1,2].

Blood bank and transfusion services collect, process, store and provide human blood intended for transfusion [2,3]. Although, ideally blood transfusion is a safe process (i.e. that saves lives and improves the quality of life in a large range of clinical conditions), there are a number of risks associated with transfusion such viral, bacterial and parasitic infection on recipient [3,4].

Since the 1980’s when the Human Immunodeficiency Virus (HIV) was recognized, rigorous screening of blood before it is supplied to recipients was instituted [5]. Those considerable efforts (national policies, improved donor selection and newer screening techniques) directed towards reducing transmissible pathogens have yielded a major reduction of viral agents especially in developed countries [6-9].

However, transfusion transmitted bacterial infection was identified as the commonest cause of complications associated with transfusion [10]. During blood transfusion bacterial infections might be originate from the environment, from the skin of the transfused subject, or from a donor bacteriaemia. Most commonly, contamination occurs during blood collection (insufficient disinfection of venipuncture site), or during handling of blood products (leaky seals) [11].

The most predominant bacteria isolated are usually commensals of the skin or gastrointestinal tract flora and the majority of isolates were Gram-positive aerobic pathogens (nearly 75%) [12].

For instance, in the United States, bacterial contamination of blood accounted for 15.9% of all transfusion related fatalities [13]. However, only few countries such as Ghana [14], Uganda [15] and, Nigeria [16] in Africa have documented records of bacterial contamination of blood/blood products. For instance, in Ghana, 9-17.5% of donor bloods were contaminated by bacteria. The major bacterial isolate identified were K. pneumoniae, E. coli, Y. enterocolitica, P. fluorescens, P. aeruginosa and Gram positive bacteria including Bacillus species and S. aureus [16,17].

In Ethiopia, blood transfusion service started in 1962 and each year more than 50,000 unit of blood has been transfused [18]. Moreover, researchers in Ethiopia depicted transfusion transmissible infectious...
agents (HIV, HBV, HCV, and T. pallidum) have been highly prevalent among blood donors across the country and still pose a threat for public health [19-21]. For instance, Baye et al. showed that the overall prevalence of HBV, HCV and malaria parasites among blood donors in Amhara and Tigray region were 6.2%, 1.7% and 1% respectively [20]. Moreover, Tsega et al. documented that the prevalence of antibody to HCV (anti-HCV) in healthy adult Ethiopian blood donors was 1.4% [21].

To the best of our knowledge, in Ethiopia, no research has been conducted to determine blood transfusion-associated infections. Therefore, the aim of this study was to determine the prevalence of bacterial contamination of blood, to identify the types of contaminating bacteria and determine their antibiotic susceptibility pattern at Debre Markos referral hospital, North West Ethiopia.

Materials and Methods

Study design and period

A facility based cross sectional study was conducted in Debre Markos referral hospital from February 2013 to June 2013.

Sampling

Primarily we mix stored blood and randomly sampled 120 units of blood from storage which was collected in Debre Markos referral hospital blood bank. The blood samples were obtained from screened, stored whole blood. All expired blood was excluded. Each unit of blood was mixed before sampling and the tubing was cleaned with 70% alcohol and cut with sterile scissors to remove any clotted blood. 5ml of blood was mixed before sampling and the tubing was cleaned with 70% alcohol and cut with sterile scissors to remove any clotted blood. 5ml of blood from storage which was collected in Debre Markos referral hospital blood bank. The blood samples were obtained from screened, stored whole blood. All expired blood was excluded. Each unit of blood was mixed before sampling and the tubing was cleaned with 70% alcohol and cut with sterile scissors to remove any clotted blood. 5ml of blood was mixed before sampling and the tubing was cleaned with 70% alcohol and cut with sterile scissors to remove any clotted blood. 5ml of blood was mixed before sampling and the tubing was cleaned with 70% alcohol and cut with sterile scissors to remove any clotted blood.

Bacterial isolation and identification

The broths were incubated at 37°C up to 7 days before they were discarded. After overnight incubation, sterile loopfuls of broth were sub-cultured on to blood agar and MacConkey agar plates and incubated aerobically for 18-24 hours at 37°C. The identities of bacteria growing on the culture plates were determined by colonial morphology, Gram and spore stains; as well as standard biochemical tests [22].

Antibiotic susceptibility testing

Selection of antibiotics is based on current treatment regimen for gram positive and negative. Susceptibility to antimicrobial agents was tested by the disc diffusion technique according to the guidelines by the Clinical and Laboratory Standards Institute (CLSI). The antibiotic discs used were ampicillin 10 μg; cotrimoxazole, 25 μg; erythromycin 15 μg; penicillin 10 units; tetracycline 30 μg; ciprofloxacin, 5 μg; gentamicin, 10 μg; ceftriaxone 30 μg and rifampicin 5 μg (Oxoid). The discs were placed on to the surface of inoculated Mueller- Hinton agar plates by an auto dispenser. After overnight incubation, the inhibition zone diameters were measured to the nearest millimeter, and isolates were classified as susceptible, intermediate, or resistant according to CLSI-specified interpretive criteria. E. coli ATCC 25922 was used as the control strain [23-27].

Statistical analysis

Data were entered, cleaned and analyzed using SPSS (Statistical Package for Social Science) version 16 by a trained data encoder. P values<0.05 were statistically significance.

Ethical considerations

Ethical clearance was obtained from Research and Publication Directorate Office of Debre Markos University, Ethiopia. Support letter was sent to Debre Markos referral hospital. Moreover, written consent was obtained from the hospital administration and laboratory head. All the data were recorded using codes, no name and other personnel identification. The result obtained in this research was confidential and only the result output disseminated for concerned bodies.

Result

Bacterial isolate and their susceptibility pattern

The length of storage of the blood ranged from 0 to 12 days (mean=4 days), and most of the contaminated samples (73.3%) had <1 week of storage (Figure 1). Of the 120 samples tested 15(12.5%) were found to be contaminated with various types of bacteria and making a 12.5% prevalence of bacterial contamination of whole blood at the Debre Markos and Fenote Selam hospital (Table 1).

Gram positive bacteria (S. pneumonia, S. aureus, coagulase-negative Staphylococcus, viridien streptococcus) and gram negative (S. typhi, E. coli and K. pneumonia) accounting 66.7 and 33.3%, respectively, were the common isolates identified (Table 1).

The two leading whole blood contaminant isolate were S. pneumonia and S. aureus. Some of the isolates were resistant to some antibiotics.

Microorganism Time of storage

<table>
<thead>
<tr>
<th>Microorganism</th>
<th>Time of Storage</th>
</tr>
</thead>
<tbody>
<tr>
<td>S. pneumonia</td>
<td>Day 0</td>
</tr>
<tr>
<td>Viridien streptococcus</td>
<td>Day 5</td>
</tr>
<tr>
<td>Viridien streptococcus</td>
<td>Day 2</td>
</tr>
<tr>
<td>Viridien streptococcus</td>
<td>Day 4</td>
</tr>
<tr>
<td>S. pneumonia</td>
<td>Day 0</td>
</tr>
<tr>
<td>S. aureus</td>
<td>Day 4</td>
</tr>
<tr>
<td>S. aureus</td>
<td>Day 3</td>
</tr>
<tr>
<td>S. aureus</td>
<td>Day 2</td>
</tr>
<tr>
<td>Coagulase negative staphylococcus</td>
<td>Day 5</td>
</tr>
<tr>
<td>Coagulase negative staphylococcus</td>
<td>Day 4</td>
</tr>
<tr>
<td>K. pneumonia</td>
<td>Day 8</td>
</tr>
<tr>
<td>S. typhi</td>
<td>Day 0</td>
</tr>
<tr>
<td>E. coli</td>
<td>Day 9</td>
</tr>
<tr>
<td>E. coli</td>
<td>Day 10</td>
</tr>
<tr>
<td>K. pneumonia</td>
<td>Day 12</td>
</tr>
</tbody>
</table>

Table 1: The isolated microorganisms and time of storage at Debre Markos and Fenote Selame hospital, North West Ethiopia, February 2013 to June 2013.
The organisms isolated in this study were both gram positive (S. aureus, coagulase negative staphylococci, viridian streptococcus, S. pneumonia) and gram negative (K. pneumonia, S. typhi and E. coli). Our finding was in agreement with study conducted in Uganda [15], Nigeria [16] and Kenya [23].

In this study, we depicted that the contamination of blood by gram positive organisms were documented within a week while contamination of blood by gram negative organisms were somewhat delayed. Our find was in agreement with different studies which claimed that Gram-positive commensals are isolated soon after donation, whereas Gram negative organisms not usually detectable until after a period of proliferation during storage [30-32].

Moreover, in contrast to other previous studies conducted elsewhere, in the present study both S. pneumonia and S. typhi were isolated at day=0 and this is an indication of bacterial contamination will be arising from donor bacterimia. As a result, efforts or measures to ensure blood transfusion safety such as adequate cleaning of phlebotomy sites, improved donor selection and screening should be strengthened.

In our finding, high rate of drug resistance for both gram negative and positive isolate were observed. These organisms might be cause septicemia and serious risk of fatality after post-transfusion takes place. Similarly, this high rate of drug resistance highlights the growing problem of antimicrobial resistance worldwide. Our finding was consistent with studies conducted in Ghana [14] and Uganda [15] and other parts of the world [31,33]. The possible explanation for the high resistance of donor blood isolates may be associated with the ease of procuring antibiotics, self medication and inefficient infection control procedure across the country.

**Conclusion and Recommendation**

Our result indicated that the prevalence of bacterial contamination of stored blood ready for transfusion at Debre Markos was high. In addition, high rate of drug resistance was observed for isolated bacterial strains. Therefore, preventive measures such as systemic and comprehensive donor selection and screening, scrubbing of the phlebotomy sites with improved disinfectants, and improved screening tests, as well as culturing of donor blood, particularly for immune compromised individuals should be carried out.
Limitation of the Study

The small sample size and our inability to follow up the recipients of the blood units to determine clinical outcome of infection may be some of the limitation of this study. In addition, we enrolled only whole blood since blood component separation does not undertaken in the hospital.

Acknowledgements

Our acknowledgements go to Ato Tariku Belachew, director of Debre Markos hospital and all members of RPO of Debre Markos University that facilitating all the bureaucratic procedures smoothly and swiftly. The authors also thank data collectors, Debre Markos hospital laboratory staffs for their unreserved support during the study period.

Competing Interests

This work was sponsored by Debre Markos University, Ethiopia. No financial aid was received from any organization for publication or other interest. There is no any competing of interest.

References