The values of platelet counts among HIV positive patients in FMC Owerri

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ABSTRACT

The values of platelet counts were carried out. One hundred and sixty four subjects were sampled, comprising one hundred and fourteen HIV positive subjects and fifty HIV negative subjects which served as the control. Platelet count was analyzed using standard techniques. The results showed that HIV positive subjects had a significantly lower Platelet count (167 ± 78.40) when compared with the HIV negative subjects (238 ± 52.04).

INTRODUCTION

Human immunodeficiency virus (HIV)

Human Immunodeficiency Virus (HIV) is a lentivirus (a member of the retrovirus family) that causes acquired Immunodeficiency Syndrome (AIDS) (Weiss, 1993).

This is a condition in humans in which progressive failure of the immune system allows life threatening opportunistic infection and cancers to thrive.

Infection with HIV occurs by the transfer of blood, semen, vaginal fluid, pre-ejaculate or breast milk of the infected person to HIV free person. Within these body fluids, HIV is present as both free virus particles and virus within infected immune cells. The four major routes of transmission are unsafe sex, contaminated needles, breast milk, and transmission from an infected mother to her baby at birth (perinatal transmission) (Fox and Cottler-Fox, 1992).
Viruses such as HIV cannot grow or reproduce on their own, the need to infect the cells of a living organism in order to replicate. The human immune system usually detects and kills viruses fairly quickly, but HIV attacks the immune system itself, the very thing that would normally get rid of the virus (Ascher and Sheppard, 1990).

HIV is a causative organism of autoimmune deficiency syndrome which was recognized as a new disease syndrome in the early 1980’s in the USA with the unusual occurrence of pneumocystis carinii pneumonia and Kaposi’s sarcoma in previously healthy young men (Greene, 1991). This retrovirus was isolated from a young homosexual man with lymphadenopathy. The virus was identified and classified in the family Retroviridae genus lentviranae (Baker et al., 2007).

Under the electron microscope, the viruses were revealed as a cylindrical core with nucleic acid cloned and sequenced. The cylindrical core is 80-130nm in diameter, it has a unique three layered structure, and innermost is the genome nucleocapside complex. This complex is enclosed within a capsid which is surrounded by a host cell membrane derived envelope, from which viral envelope glycoprotein ‘spikes’ project. HIV infects a wide variety of tissues in humans including the marrow, lymph node, brain, skin and bowel (Baker et al., 2007). This retrovirus differs from other retroviruses such as human T lymphotrophic virus (HTLV) 1 and 2. The virus was eventually named Human Immunodeficiency Virus (Cohan et al., 1986).

It is transmitted mostly sexually in blood or blood products and pre-natally. The most at risk of acquiring HIV infection are homosexuals, injecting drug misuses and those with bisexual orientation. Others include individuals receiving unscreened blood or blood products, infants born of infected women.

There are various strains of HIV and are designated by a code with geographically informative letters and sequential numbers placed either in brackets, or as a number, or as a subscript. Example HIV-sf33 and HIV -2 (Pantaleo and Fauci, 1995).

If there is a laboratory evidence of HIV infection, certain indicator diseases that require presumptive and definitive diagnosis are diagnostic of AIDS, AIDS is an illness characterized by one or more indicator diseases (Safrit and Koup, 1995).

Acute HIV is usually characterized by fever, malaise, lymphadenopathy and rash. These conditions are subclinical. A chronic infection of AIDS that follows is asymptomatic in early stages. If an individual is infected with this virus, the virus acts so quick destroying the immune system making the individual prone to little infections. HIV is present all over the world and the long term consequences of this pandemic will affect every country one way or another over time. This is an evolving pandemic threatening global public health and health care provision, as well as political and economic stability (Kuby, 1997).

**Coagulation**

This is a complex physiological process by which blood forms clot. It is important part of haemostasis (the cessation of blood loss from damaged vessel), wherein a damaged blood vessel wall is covered by a platelet and fibrin-containing clot to stop bleeding and begin repair of the damaged vessel (Shapiro, 2003).
Coagulation results from interactions among vessel wall, platelet and coagulation factors. When an injury occurs that results in bleeding, the coagulation system is activated and plugs the hole in the bleeding vessel while still keeping blood flowing through the vessels by preventing the clot from getting too large. The end result is the formation of insoluble fibrin threads that link together at the site of injury, along with aggregated cell fragments called platelets to form a stable blood clot. The clot prevents additional blood loss and remains in place until the injured areas have healed. The clot is eventually removed as the injured site is healed. In normal healthy individuals, this balance between clot formation and removal ensures that bleeding does not become excessive and that clots are removed once they are no longer needed (Shapiro, 2003).

A number of coagulation abnormalities have been described in human immunodeficiency virus (HIV) disease. High levels of plasma von Willebrand factor have been reported in HIV disease and might be indicative of activated endothelium. Endothelium is involved in important homeostatic mechanisms of non-thrombotic vascular surfaces, vascular tone regulation and immunomodulation (Karpatkin et al., 2002).

Injured endothelium leads to localized inflammatory response of which the direct consequence is the occurrence of occlusive thrombosis events mediated between leucocyte recruitment and platelet adhesion and aggregation, blood clotting activation and fibrinolysis derangement. HIV infection has been associated with endothelial dysfunction. Since HIV infection is associated with endothelial dysfunction it may therefore result in activation and consumption of coagulation factors and ultimately coagulation defect (Omoregie et al., 2009).

In HIV infection, the liver is affected. The liver is the major organ responsible for the synthesis of most coagulation factors and infection of the liver by HIV can lead to abnormal production of coagulation factors. The CD4+ count is used to measure immune status and HIV disease progression (Tolstrup et al., 2004).

Prothrombin time (PT) and an activated partial thromboplastin time (APTT) are screening tests for the extrinsic and intrinsic clotting systems respectively. They detect deficiency or inhibition of clotting factors in either system, and are the first tests in screening for coagulation disorders. As HIV infection progresses, endothelial dysfunction and liver damage will increase and this may result in severe clotting impairment.

In reference to the abnormality of coagulation in HIV positive individuals, the coagulation disorders will be investigated, by considering platelet count, prothrombin time, activated partial thromboplastin time, and blood fibrinogen concentration, as well as CD4 count and factor VII concentration.

Platelet count is a diagnostic test that determines the number of platelets in the patient’s blood. Platelet which are also called thrombocytes, are small disk-shaped blood cells produced in the bone marrow and involved in the process of blood clotting. There are normally between 150,000–450,000 platelets in each microlitre of blood. Low platelet count or abnormally shaped platelets are associated with bleeding disorders (Henry, 2001).

Prothrombin time (PT), is one of the coagulation factors produced by the liver. One of the final steps of the cascade is the conversion of Prothrombin (factor 11) to thrombin. The Prothrombin time test evaluates the integrated function of the
coagulation factors that comprises the extrinsic and common pathways.

The international Normalized Ratio (INR) is used to standardize PT result gotten (Horsti et al., 2005).

Activated partial thromboplastin time (APTT), is a screening test that is done to help evaluate a person’s ability to form blood clot. It assesses the amount as well as the function of coagulation factors XII, IX, VII, X, V, II and I which are part of haemostasis (Pagana and Pagana, 2006).

Fibrinogen (factor 1) is a soluble plasma glycoprotein synthesized by the liver and is converted by thrombin into fibrin during blood coagulation.

Fibrinogen deficiency (hypofibrinogenemia) or disturbed function of fibrinogen can lead to either bleeding or thromboembolic complications (Acharga and Dimichele, 2008).

CD4 count is the number of CD4 cells per microlitre of blood. It is used to stage the patient’s disease, determine the risks of opportunistic illness, assess prognosis and guide decisions about when to start antiretroviral treatment (CDC, 2009).

The main aim of this study to determine the value of platelet count among HIV positive subjects in FMC Owerri.

Materials and Methods

Sample size calculation

The sample size for this study was calculated based on Ijeoma et al. (2010) 8.1% prevalence of HIV in Owerri in 2010.

\[ n = \frac{1.96^2 \times 0.08 (1.00-0.08)}{(0.05)^2} = 114 \text{ samples} \]

Informed consent

Participant information sheet (PIS) was given to the prospective participants. After reading and understanding the PIS, questions were asked and proper explanations given. They consented to participate in the study by signing the informed consent form.

Eligibility criteria

Informed consented subjects (both HIV/AIDS positive patients and HIV negative controls).

Subjects

One hundred and fourteen HIV positive subjects aged 18-65 years attending Heart to Heart clinic of Federal Medical Centre, Owerri were screened. Fifty HIV negative subjects were also screened and they served as controls.

Sample collection

Informed consented subjects were sampled. 2.5mls of blood was added into ethylene diamine tetra acetic acid (EDTA) bottle and mixed immediately by reverse uniform inversion for platelet counts. Platelet count was performed using sysmex automated haematology analyzer KY21N model manufactured by sysmex corporation Kobe, Japan.

Platelet (Direct Current Detection Method)

By automation using Sysmex automated haematology analyzer KY21N model manufactured by Sysmex Corporation Kobe, Japan.
The aspirated blood sample is measured to a predetermined volume, diluted at the specified ratio, and then fed into each transducer chamber, which has a minute hole - the aperture which also contains the electrodes in which direct current flows. Blood cells suspended in the diluents sample pass through the aperture, causing direct current resistance to change between the electrodes and the blood cell size is detected as electric pulses, and the histogram determined by the pulse sizes.

The sample in EDTA bottle was places in the spiral mixer and allowed to mix very well. Whole blood mode was activated in the LCD screen, the sample number (code) was inputed via the key board and then the enter key.

Then the sample was mixed very well again, the cap was removed and inserted into the probe, on that condition, the start switch was pressed. The LCD screen displays the sample analysis, and then sample was removed and recapped. The unit executes automatic analysis and displays the result on the screen.

Statistical analysis

The data obtained were subjected to some statistical analysis such as the mean (X), standard deviation (SD), standard error of mean (SEM), student’s t-test and Pearson moment of correlation using statistical package for social sciences (SPSS) version 17. The results were expressed in mean ± standard error of mean.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>HIV Negative subjects (n = 50)</th>
<th>HIV Positive subjects (n =114)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Platelet(×10^9)</td>
<td>238.00 ± 52.04</td>
<td>167 ± 78.40</td>
<td>P &lt; 0.05</td>
</tr>
</tbody>
</table>

NB: Figures are in mean ± standard deviation; n= number tested.

Results and Discussion

The mean value of platelet Count among the study subjects are shown in table below. Platelet counts were significantly lower in HIV patients (P< 0.05) when compared with the seronegative subjects (control).

HIV infection is associated with endothelial dysfunction and liver damage. Both endothelial dysfunction and liver damage can result in coagulation defects. It is therefore expected that as the HIV infection progresses, the coagulation abnormalities will increase (Linder et al., 1970).

The platelet count was significantly lower in HIV seropositive patients (p < 0.05) when compared to the seronegative controls which also agree with the works of Omoroge et al. (2009) and (van Gorp et al., 1999) who independently established a significant reduction in platelet count among HIV positive patients in their separate studies. Impaired thrombopoiesis and production of anti platelet antibodies have been suggested as possible mechanisms (Karpatkin et al., 2002). Impaired thrombopoiesis can result from infection of megakaryocytes by HIV, because megakaryocytes posses CD4 and CXCR4, which are known receptors for HIV, and various megakaryocytes lines are infectable with HIV (Najen and Rain, 1994).

It has been reported that patients with AIDS have decreased platelet production whereas patients with early onset HIV infection are more likely to have increased peripheral destruction of platelet by anti platelet antibodies.
Conclusion

Coagulation abnormalities have been described in HIV disease. HIV infection has been associated with endothelial dysfunction which may result in activation and consumption of coagulation factors and ultimately coagulation defect. The findings of this study indicate decrease in platelet count in HIV Positive.

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