Resolution of dilated cardiomyopathy in an adolescent with change of a failing highly active antiretroviral drug therapy

Twalib Olega Aliku1, Sulaiman Lubega2, Peter Lwabi1

1. Department of Paediatrics and Child Health, Gulu University.
2. Uganda Heart Institute, Mulago Hospital Complex.

Abstract

Background: Cardiovascular dysfunction is a recognized complication of HIV infection in children. Cardiac complications of HIV usually occur late in the course of the disease; they may be associated with drug therapy, and hence become more common as therapy and survival improve. Left ventricular (LV) dysfunction at baseline is a risk factor for death independent of the CD4 cell count, HIV viral load, and neurological disease.

Clinical case: We present the case of a 15 year old girl with HIV who developed left ventricular dysfunction while non-compliant on highly active antiretroviral therapy (HAART). She presented with features of heart failure over a course of two months. Her laboratory evaluation was significant for leukopenia with a low CD4 count, high viral load, elevated ESR and CRP. The ECG showed a sinus tachycardia with diffuse ST-T segment changes and LVH with strain. Initial echo revealed dilated left heart chambers with poor LV systolic function and a small pericardial effusion with the development of an LV thrombus on follow up echo evaluation. She was started on heart failure medicines and had anticoagulation for the LV thrombus. She received adherence counseling and her HAART regimen was changed. Six months after presentation she became asymptomatic with higher CD4 counts and a normal LV size and function on echo.

Conclusion: Immunoological recovery following a switch of a failing or potentially cardiotoxic HAART in addition to improved HAART adherence may result in resolution of left ventricular dysfunction. Early and regular cardiology evaluation may improve outcomes in these patients.

Key words: Dilated Cardiomyopathy, HAART

DOI: http://dx.doi.org/10.4314/ahs.v15i1.39

Background

Cardiovascular abnormalities in children infected with HIV are frequent and often persistent1. Cardiac complications of HIV usually occur late in the course of the disease with prolonged viraemia2-3. In HIV infected children not appropriately treated with highly active antiretroviral therapy (HAART), dilated cardiomyopathy is one of the commonest abnormalities seen. These abnormalities may range from subclinical ECG changes to overt heart failure from cardiomyopathy and sudden cardiac death1/4. Left ventricular function at baseline has been found to be a risk factor for death that is independent of the CD4 count, HIV viral load and neurological disease1. Long term HAART is cardioprotective for HIV infected children and adolescents1, and reduces the risk of cardiomyopathy by 50% in children compared to those not on HAART3.

Case report

We present the case of a 15 year old girl with HIV diagnosed at the age of ten years. For the past five years she received a HAART regimen containing zidovudine, lamivudine, and nevirapine. She was in boarding secondary school and occasionally missed some of her drugs (she had to take at one time ten different drugs). She presented with a two month history of cough, dyspnea, hemoptysis and right hypochondric pain. She underwent evaluation for pulmonary tuberculosis for her symptoms. Her sputum was negative for alcohol-acid fast bacilli. She was subsequently referred for cardiology consult on the basis of cardiomegaly on the chest radiograph. The physical exam revealed a heart rate of 118 beats per minute, respiratory rate of 24 breaths per minute, BP=106/79mmHg, and oxygen saturation of 96% in room air. She had orthopnea and mild pedal edema. Her weight was 42kg (between 5th and 10th percentile) and height=152cm (between 5th and 10th Percentile). Her cardiovascular exam was remarkable for tachycardia, small volume pulse, displaced apex beat to the 6th intercostal space in the anterior axillary line, a non-sustained apical heave, a third heart sound, basal chest crepitations and a tender hepatomegaly. She had elevated acute phase reactants. The serum electrolytes, urea and creatinine were normal. At initial cardiology evaluation her CD4 count was 51 cells per ml and the HIV viral load was 36,869 copies per ml. Initial lipid profile was not done. The chest radiography revealed cardiomegaly with cardiothoracic ratio of 0.7, interstitial edema and blunted left costophrenic angle. Her ECG showed a sinus tachycardia with heart rate of 122 beats per minute, diffuse ST-T segment changes and LVH with strain (see figure 1).

Corresponding author:
Twalib Olega Aliku
Department of Paediatrics and Child Health, Gulu University.
PO Box 166 Gulu
mobile:+256712551204.
Email: aliku90@yahoo.com

Figure1: The ECG of the patient at initial presentation. Note the diffuse inferolateral ST-T changes.

She had elevated acute phase reactants. The serum electrolytes, urea and creatinine were normal. At initial cardiology evaluation her CD4 count was 51 cells per ml and the HIV viral load was 36,869 copies per ml. Initial lipid profile was not done. The chest radiography revealed cardiomegaly with cardiothoracic ratio of 0.7, interstitial edema and blunted left costophrenic angle. Her ECG showed a sinus tachycardia with heart rate of 122 beats per minute, diffuse ST-T segment changes and LVH with strain (see figure 1).

Figure1: The ECG of the patient at initial presentation. Note the diffuse inferolateral ST-T changes.
One month later her symptoms had improved. A repeat echo showed no thrombi with an improved LV function (FS=32%, EF=60%). Six months after diagnosis, she was asymptomatic. Her echo revealed normal LV size (LVEDD=5.1cm, Z-score=1.72), with normal systolic function (FS=32%, EF=60%). An upward trend in her LV function continued to 12 months of follow up (see figure 2). Warfarin and the other cardiac meds were discontinued except lisinopril. Her CD4 count after 6 months from switch of her HAART regimen was 360 copies per ml. The viral load was not done due to excessive cost. At 12 months of follow up her ECG showed diffuse T-wave inversions (see figure 3).

Discussion

In HIV infected children less than 10 years of age, 25% may die with chronic cardiac disease and 28% experience serious cardiovascular events after the onset of an AIDS defining illness, especially in those not appropriately treated on HAART. Our patient presented with longstanding cough and hemoptysis that prompted initial evaluation for pulmonary tuberculosis. In HIV infected patients, particularly those with advanced disease; concurrent respiratory infections, pulmonary hypertension, anemia, protein energy malnutrition, or HIV associated malignancies may alter or obscure the characteristic signs that often define congestive heart failure. In resource limited settings, this could potentially delay the cardiology referral of such patients and contribute to worse outcomes.

The presence of elevated acute phase reactants with tachycardia, diffuse ST-T changes on ECG and severe LV systolic dysfunction and pericardial effusion on echo could suggest myocarditis as the cause of the dilated cardiomyopathy in this patient. However, tachycardia is a nonspecific finding in patients with HIV and may be related to autonomic dysfunction. Dilated cardiomyopathy is strongly associated with CD4 counts lower than 100 cells per ml as was the case in this patient. Diffuse T-wave inversions are characteristic of myocarditis. Different possible mechanisms may explain the etiology of HIV associated cardiomyopathy. These include the HIV itself, autoimmune response to viral infection, cardiotoxicity or direct mitochondrial injury from therapeutic agents such as zidovudine, nutritional deficiencies and cytokine overexpression. HIV infected patients with dilated cardiomyopathy are more likely to have myocarditis and a wider spectrum of viral infections than HIV negative patients with idiopathic dilated cardiomyopathy. Immune deficiency may favor the selection of those viral variants with increased pathogenicity or enhance the cardiovirulence of viral strains. Our patient’s non adherence to her HAART regimen (in part due to high pill burden sometimes) most likely contributed to the treatment failure.

The combination of low CD4 counts, high viral loads and treatment with potentially cardiotoxic zidovudine in this setting of an infectious insult could have provided the perfect milieu for left ventricular dysfunction in this patient. Change in these risks through switch of HAART regimen to eliminate cardiotoxicity and promote immunological recovery; prompt treatment of concurrent infection and early management of the cardiac dysfunction could have contributed to the better outcome seen in this patient. Tudor et al described a similar case of an adolescent with HIV on HAART who had dilated cardiomyopathy. This patient had suboptimal immunological recovery and failure of virological suppression to undetectable levels. With change in the HAART regimen and immunological recovery, the left ventricular dysfunction improved.

Patients with LV systolic dysfunction are at an increased risk of LV thrombus formation due to associated stasis. However endothelial dysfunction that is common in patients with HIV contributes to a highly procoagulant state. Our patient had features suggestive of a transient ischemic attack and a further echocardiographic examination was able to demonstrate an intracardiac thrombus as the most likely cause.

Even though HAART substantially reduces the risk of cardiomyopathy, patients still remain at increased risk, particularly those with older age at HAART initiation, on zidovudine containing regimen or those with poor adherence to therapy and treatment failure. The patient is currently being followed regularly since she is on protease inhibitors that are associated with increased cardiovascular risk due to associated dyslipidaemia. Her ECG at 12 months of follow up had subclinical ECG abnormalities.

Echocardiography in children with HIV is recommended in patients at increased cardiovascular risk, those with clinical evidence of cardiovascular disease, unexplained or persistent pulmonary symptoms and every 1 to 2 years thereafter or as clinically indicated. Mondy et al found frequent echocardiographic abnormalities among generally healthy adults on HAART, including LV systolic dysfunction in 18%. Progressive LV dilatation is common in HIV infected children and may herald the onset of congestive cardiac failure. Rapid onset congestive heart failure has a grim prognosis in HIV infected adults and children, with more than half of patients dying from primary cardiac failure within 12 months of presentation.

Conclusion

With standard treatment of the heart failure, improving immunological recovery and optimal virological (suppression from improved adherence) or a change in HAART regimen, patients with HIV may have a resolution of their left ventricular dysfunction. However, they still need a regular cardiology follow up to detect any cardiovascular event, especially in the event of use of HAART with increased cardiovascular toxicity.

References
2. Lipshultz SE, Easley K, Orav EJ et al. Cardiac dys-


