Case Report

A Case of Sudden Cardiac Arrest in a Fabry Disease Patient with Kidney Transplantation - ə

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INTRODUCTION

Fabry Disease (FD) is an X-linked recessive lysosomal storage disorder associated with a deficiency α-galactosidase A resulting in accumulation of glycosphingolipids in the vascular endothelium which leads to potentially fatal renal, cardiac and cerebrovascular conditions [1]. Accumulation of its major substrate, Globotriaosylceramide (Gb3) which is mainly responsible for organ damage and its deacylated derivative, Globotriaosylsphingosine (Lyso-Gb3) in the fluids and cellular lysosomes throughout the body deteriorate organ functions, most frequently of the kidneys and heart [2].

The clinical features of Fabry disease include acroparaesthesia, angiokeratomas, hypohidrosis, corneal and lenticular opacities, and major end-organ disease (involvement of the kidneys, heart and brain) [3]. Renal pathology is one of the hallmarks of FD and is the most frequent cause of death, usually when patients are aged 30-50 years [4]. Because Fabry nephropathy does not recur in the allograft and transplanted Fabry patients appear to have better overall outcomes than those maintained on dialysis, kidney transplantation should be recommended as a first choice in Renal Replacement Therapy (RRT) for Fabry disease [5].

We present a male patient with FD who was transplanted with kidney from a living donor and had a sudden cardiac arrest on the 4th day after operation.

CASE REPORT

A 44-year-old male with FD had been on hemodialysis for 7 years. He had undergone enzyme replacement therapy with α-galactosidase A (0.2 mg/kg every two weeks) for nine months. His weight was 49 kg and height 166 cm (5.35 inches). According to the American Society of Anesthesiologists (ASA) physical status classification he was a ASA III patient. He had a history hyperlipidemia, coronary artery disease, of Anesthesiologists (ASA) physical status classification he was a ASA III patient. He had a history hyperlipidemia, coronary artery disease, arrhythmia (Ventricular Extrasystoles-VES), heart failure and ankylosing spondylitis. There was sinus rhythm on electrocardiogram (Figure 1). A screening pretransplant echocardiography showed left ventricular hypertrophy with ejection fraction of 60%, dilated left atrium, mitral valve regurgitation, and normal right ventricular and right atrial size with normal pulmonary artery pressure. Results of preoperative coronary angiography were left anterior descending artery with irregular borders and 50% stenosis at D2 branch, right coronary artery with irregular borders and plaque at ostium of circumflex artery. Medical therapy which were atorvastatin 20 mg/day, metoprolol 25 mg/day and acetylsalicylic acid 100 mg/day began for coronary artery disease. The patient had normal laboratory values except hemoglobin- hematocrit (10.8 g/dL-36.2 %), BUN (57 mg/dL), urea (123 mg/dL) and creatinine (5.01 mg/dL). He was on hemodialysis for 7 years and his estimated glomerular filtration rate was 13 mL/min (according to the Cockcroft gault formula).

On the physical examination for the patient whose mallampathy score was evaluated to be III, macroglossia and limited neck flexion were observed, and a video laryngoscope was prepared in case of a difficult intubation. The patient was taken into operation hall and anaesthesia induction was started after monitoring. A hemodialysis catheter which was placed before operation day was used as an intravenous (I.V.) route, and lidocaine 40 mg, propofol 100 mg, and fentanyl 100 μg was administered and the patient was ventilated with a mask. After that, muscular relaxation was obtained with 0.5 mg/kg rocuronium. The patient was being intubated and localization of tube was confirmed by auscultation and capnography and fixed. Maintenance of anaesthesia was continued with propofol 4mg/kg/hour and remifentanil 0,15 mcg/kg/hour infusions and 0.15 mg/kg rocuronium I.V. as required. End-tidal CO₂ and temperature monitorizations were made on the patient. Peripheral vascular route was opened with 16 G angiographic catheter, Central Venous Pressure (CVP) monitoring and left radial artery cannulation were performed and arterial blood pressure monitoring was made, and hourly urine output was monitored with a urinary catheter placed. The patient was hemodynamically stable maintaining a mean arterial pressure of 60-65 mmHg. After uncomplicated surgery, extubation was performed in operative room and awakening was uneventful. Analgesia was granted with intravenous tramadol. He was handed over to the clinic without problem after followed in postanaesthesia care unit for 1 hour during postoperative period. On the 4th day after operation, he had sudden cardiac arrest with asystole as the determined rhythm. Cardiopulmonary Resuscitation (CPR) was started immediately. Adrenaline was given 4 times at 1 mg doses

ABSTRACT

Fabry Disease (FD), also known as Anderson-Fabry disease, is an inherited X-linked disorder characterized by the absence (in men) or deficiency (in women) of α-galactosidase A, activity that causes a progressive accumulation of glycosphingolipids within lysosomes of cells in all the major organ systems and progressive organ damage that first manifests in childhood or early adulthood. End Stage Renal Disease (ESRD) is a major cause of morbidity and premature mortality in FD. We present a male patient with FD who was transplanted with kidney from a living donor and had a sudden cardiac arrest on the 4th day after operation. We suggest detailed preoperative examination including coronary angiography, echocardiography for these patients and also a multidisciplinary care is required for perioperative management of FD patients.

Keywords: Fabry disease; Kidney transplantation; Cardiac arrest
intravenously every 3 minutes. The rhythm which was ventricular tachycardia returned at the 7th minute after CPR. Defibrillation was applied 4 times as 200 joules and amiodarone was given at 300 mg as a bolus dose in 10 minutes, 900 mg as a maintenance dose in 18 hours intravenously. The patient was stable hemodynamically at the 27th minute after cardiac arrest. He was admitted to the Intensive Care Unit (ICU) in an orotracheal intubated state. There was right bundle branch block and ST segment depression in the anterolateral leads on electrocardiogram (Figure 2). A transthoracic echocardiography was applied on post-cardiac arrest state. It showed left ventricular hypertrophy with ejection fraction of 50%, mitral valve regurgitation, hypokinetic interventricular septum, normal right ventricular and right atrial size with normal pulmonary artery pressure. His medical therapy was ordered as enoxaparin sodium 0.6 mL/day, acetylsalicylic acid 100 mg/day, metoprolol 50 mg/day, atorvastatin 20 mg/day and also immunosuppressive therapy. The patient was extubated on the 3th day. He was discharged to home without problem, and was followed up in the clinic of general surgery.

DISCUSSION

Fabry disease is a rare lysosomal storage disorder. The disease first described simultaneously but separately in 1898 by dermatologists from Germany, Johannes Fabry and the United Kingdom, William Anderson [6]. The incidence has been estimated to be 1:117,000 live births and 1:50,000 males [7]. Birth prevalence of the female Fabry carriers were only studied in some reports, and were therefore not known exactly. In 1947, the observation of abnormal vacuoles in blood vessels throughout the body at post mortem examination in 2 patients who had died from renal insufficiency led to the recognition of this disease being a generalized storage disorder [8].

Accumulation of the glycosphingolipid Globotriaosylceramide (Gb3) results and causes cellular dysfunction not only in several organ systems, mainly skin, kidney, heart, lung, and brain, but also in the gastrointestinal tract and cornea. The injured endothelial cells of small and large vessels and vascular smooth muscle cells cause dysfunction of the heart and brain leading to early onset of hypertension, concentric left ventricular hypertrophy without obstruction, and coronary heart disease, as well as vertebrobasilar artery signs and symptoms, excessive daytime sleepiness, and stroke [6]. The major causes of death include cardiac death, stroke, or the consequences of ESRD [9-10]. ESRD occurs in the majority of male patients and in a significant proportion of females who require dialysis or transplantation. Importantly, 12 % of ESRD patients with FD were female in both registries [5].

The first kidney transplant in a FD patient was conducted in 1967 in Basel, Switzerland. Sorbello et al. reported two patients with FD who underwent a deceased donor kidney transplantation, by focusing on the anaesthesiologic implications for this rare disease [4]. They demonstrated that a standard anaesthesiological protocol could be applied in these patients, but careful perioperative management is mandatory. Kruger et al. presented two female patients with FD who required general anesthesia twice for gynecological and trauma surgery, respectively, and discuss their perioperative management [6]. They reported that compared with equivalent patients without FD, they had to deal with early-onset arterial hypertension, hyperalgesia, cardiac myopathy, and renal dysfunction. They also concluded that patients with mild progression of the disease require only a few adjustments to keep them stable with a beneficial outcome.

In this report, we present a male patient with FD who had arrhythmia and ankylosing spondylitis. The patients with FD have rheumatologic findings such as joint pain, acroparäesthesia, Raynaud phenomenon. However, ankylosing spondylitis is not common finding of FD in the literature. In our case we planned difficult airway algorithm because of ankylosing spondylitis, mallampathy score III, macroglossia and limited neck flexion, but intubation did not force us as we expected. FD has serious cardiovascular manifestations like conduction system abnormalities, arterial hypertension, myocardial infarction, congestive heart failure, arrhythmia which can result in sudden death [11]. Our patient also had a cardiac arrest on the 4th day after operation which was managed successfully. He had a history of hyperlipidemia, coronary artery disease, arrhythmia and heart failure preoperatively. Although CAG was done and treatment was started according to the results of CAG before operation, surgical stress may have caused cardiac depression additionally. He didn’t have any electrolyte abnormalities. We didn’t observe any non-sustained ventricular tachycardia during monitoring. He was hemodynamically stable before cardiac arrest, so we have linked cardiac arrest to pre-existing cardiomyopathy. We informed the patient about his cardiac status and advised him to be examined by cardiologist after discharge.

CONCLUSION

Patient diagnosed with FD challenge anesthesiologist due to cardiac, renal and cerebral disfunctions, especially cardiac ones. Preoperative examination should make carefully including coronary angiography and echocardiography. According to our experience, we recommend advanced hemodynamic monitoring during surgery. Careful airway examination should be further performed and alternative strategies for airway management should be planned by considering a difficult intubation. A nephroprotective and cardioprotective strategies and a particular care to the associated end-stage organ disease may significantly improve the long-term outcome of patients with FD. A multidisciplinary care is required for perioperative management of these patients.

CONSENT

Written informed consent was obtained from the patient for publication of this case report and any accompanying images.

AUTHORS’ CONTRIBUTIONS

All of the authors contributed to the medical management of the
patient and preparation of the manuscript. All of the authors have read and approved the content of the manuscript.

REFERENCES


