Cardiac electrical system involvement in Alström syndrome: uncommon causes of dilated cardiomyopathies

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Abstract

Alström syndrome is a rare autosomal recessive disorder with dilated cardiomyopathy in 60% of patients. Despite the frequency of cardiac involvement in Alström syndrome, conduction system abnormalities or arrhythmias have not been characterized previously. We report two siblings with Alström syndrome with conduction system involvement with left bundle branch block on electrocardiogram (ECG). One patient had first degree atrioventricular block in addition to bundle branch block and underwent pacemaker implantation. This same patient developed intra-atrial re-entry tachycardia requiring anti-arrhythmic medication and eventual trans-catheter ablation. The second patient developed atrial and ventricular arrhythmias and underwent placement of a bi-ventricular defibrillator. These findings suggest that cardiac conduction system involvement and clinical arrhythmia may be significant yet under-recognized complications in patients with Alström syndrome. Patients should be routinely screened with ECG and Holter monitoring in addition to echocardiographic assessment and a cardiologist experienced with cardiomyopathy should be an integral part of the care team.

Case Reports

Case #1

The first patient is a 20-year-old male who initially presented to cardiology at the age of 17 years for tachycardia (Figure 1A). Prior to his evaluation by cardiology, he had been diagnosed with retinitis pigmentosa and had severe visual impairment. Upon presentation he had moderate ventricular dysfunction with a shortening fraction (SF) of 23% and mild left atrial enlargement (Figure 2A). During this admission, the patient was started on digoxin for treatment of his tachycardia. At his initial outpatient follow-up visit, he demonstrated improved ventricular function with a shortening fraction of 30%. He was initially diagnosed with tachycardia-mediated cardiomyopathy. His cardiac function was thought to likely improve in time with management of the arrhythmia.

Six months later he was noted to have concerning left ventricular (LV) dilation and a progressive widening of his QRS interval (148 ms combined; PR interval of 196 ms) with stable function (Figure 3A). These new electrocardiogram (ECG) findings in the context of his previously noted retinitis pigmentosa raised concern for Kears-Sayre syndrome. He was referred to a pediatric geneticist and a cardiologist with cardiomyopathy experience. On exam he was noted to have short stature (2%ile). Additional testing demonstrated high frequency sensorineural hearing loss and glucose intolerance. mtDNA testing in blood for common deletions/duplications including Kears-Sayre syndrome was normal. A cardiac biopsy showed displacement of myofibrils. By electron microscopy, the intermyofibril spaces were filled with increased numbers of mitochondria with striking proliferative features and, in some
areas, glycogen granules, consistent with a mitochondrial disorder. He developed progressive first degree AV block, and underwent placement of a dual chamber pacemaker. A skeletal muscle biopsy was performed while under anesthesia for his pacemaker placement. Testing of the muscle biopsy for deletions, the gold standard diagnostic test for Kearns-Sayre, was normal. Given this patient’s constellation of symptoms, Alström syndrome was considered. He was found to have two **ALMS1** mutations (c.7369_7370delGA (p.Asp2457Stop) and c.7374_7375delGA (p.Asp2459Stop)) which confirmed a diagnosis of Alström syndrome.

Over the next two years this patient developed progressive QRS widening with a left bundle branch block pattern and was admitted for atrial flutter which was successfully cardioverted. Five months following his device implantation he developed atrial flutter refractory to electrical cardioversion or amiodarone therapy, and required trans-venous ablation of several intra-atrial reentry circuits (Figure 3B). To date he remains clinically stable in sinus rhythm with mild LV dysfunction on a medical regimen of atenolol, lisinopril, metformin and lasix.

### Case #2

The second patient is a 25-year-old female (Figure 1B) who initially presented for cardiac evaluation at the age of 23 years following the diagnosis of dilated cardiomyopathy in her younger brother (Patient #1). Prior to evaluation in the cardiology clinic, she had been diagnosed with retinitis pigmentosa, systemic hypertension and Type II diabetes mellitus. Upon initial evaluation, her echocardiogram demonstrated borderline normal ventricular function with a SF of 27% and mild left atrial dilation (Figure 2B). Initial ECG demonstrated a left bundle branch block pattern with a QRS duration of 125 ms (Figure 3C). She was started on carvedilol in addition to her prior regimen of lisinopril and metformin.

**ALMS1** testing was performed on this patient for her known familial mutations which confirmed a diagnosis of Alström syndrome. Since her diagnosis she has developed progressive LV dysfunction, with a current SF of 19% and ejection fraction <35%. Associated with the decline in cardiac function, she developed frequent episodes of non-sustained ventricular tachycardia. Secondary to the combination of poor systolic function, left bundle branch block and ventricular arrhythmias she underwent placement of a bi-ventricular defibrillator system. Subsequently, she went on to develop recurrent atrial flutter and is currently well controlled on Amiodarone.

### Materials and Methods

Patient, family and medical history were evaluated in two siblings with Alström syndrome including genetic, echocardiographic and electrophysiological data. Patient information was collected as part of a clinical registry study approved by the Institutional Review Board of Cincinnati Children’s Hospital. Patient consent was obtained for photography and publication.
Discussion

Dilated cardiomyopathy is a frequently heritable and familial condition, however a pathogenic mutation is found in only approximately 30-40% of patients tested. In the pediatric age group there are both genetic and environmental causes of dilated cardiomyopathy. Genetic causes include mitochondrial disorders, fatty acid oxidation defects, genetic syndromic conditions, or mutations in sarcomeric or cytoskeletal genes. Although most genetic syndromes associated with dilated cardiomyopathy are rare, in aggregate they account for a substantial fraction of disease. Ciliopathies are an increasingly recognized group of heterogeneous genetic disorders. The molecular mechanism underlying dilated cardiomyopathy in Alström syndrome is not known. Furthermore, there is a paucity of information on cardiac involvement in other ciliopathies. Cardiomyopathy has been described in a subset of patients with Leber’s congenital amaurosis. A small case series described 2/8 patients with Bardet-Biedl syndrome with cardiomyopathy but provided no additional details. Finally, Elbedour et al. documented cardiac involvement in 50% of Bardet-Biedl patients including congenital heart defects, hypertrophy of the interventricular septum, and dilated cardiomyopathy, and recommended ongoing cardiac surveillance. Given that many of the ciliopathies share clinical and etiologic features, the possibility of cardiac dysfunction in other ciliopathies needs further evaluation.

Cardiac conduction system disease is often associated with various forms of cardiomyopathy, and clinical arrhythmias are frequently degenerative. In specific subtypes of cardiomyopathies electrocardiographic abnormalities may precede echocardiogram evidence of functional abnormality. These ECG findings have been associated with worse patient outcome. Thus, an abnormal ECG may be a prognostic indicator for development of future ventricular dysfunction and identify a high risk group of patients within the larger population of patients with cardiomyopathy. The mechanisms underlying conduction system abnormalities in patients with Alström syndrome are not known and potentially multi-factorial. Similar to central conduction system disease, arrhythmias may occur either as a result of the underlying muscle cell abnormality or as a result of poor systolic function and ventricular dilation in patients with cardiomyopathy. Although other ciliopathies syndromes including Bardet-Biedl and Kartagener’s syndrome have been associated with significant cardiac involvement, potential conduction system and arrhythmia involvement remains unknown. Accurate description of the frequency of ECG abnormalities and associated arrhythmias in patients with Alström syndrome compared to other cardiomyopathy cohorts is needed to delineate whether these abnormalities are secondary to the dilated cardiomyopathy phenotype or potentially related to the ALMS1 mutation itself.

Previously it was thought that ventricular dysfunction diagnosed during infancy in Alström syndrome was transient. More recent reports have shown that in up to 30% of patients normalization of ventricular function is merely a quiescent period, with cardiomyopathy recurring in adolescence or adulthood. The presence of cardiomyopathy in adolescent

![Image of electrocardiograms](image_url)
and adult patients with Alström syndrome is associated with a poor prognosis and significant mortality. In response to the significant incidence of cardiomyopathy in patients with Alström syndrome, serial echocardiographic assessment has been recommended as part of routine patient care. Although cardiac involvement is a common finding in patients with Alström syndrome, there has been no previous description of the central conduction system or cardiac rhythm disturbances in this patient population. This report represents the first description of both central conduction system disease and cardiac rhythm abnormalities in patients with Alström syndrome. Whether conduction system abnormalities are truly rare or whether there is under-reporting or under-recognition of these defects is unknown and more detailed analysis of ECG and arrhythmic involvement in larger Alström populations is needed.

Early detection of conduction defects in patients with Alström syndrome would allow for earlier and more aggressive treatment reducing morbidity and mortality. Prior studies in a variety of cardiomyopathy subgroups have demonstrated that preceding and predictive ECG changes can be used as a prognostic tool indicating the need for early treatment of cardiomyopathy. Hombach et al. reported that a QRS interval >110 ms is an independent predictor of both overall cardiac death as well as sudden cardiac death in a large cohort of adult cardiomyopathy patients. Whether ECG abnormalities precede echocardiographic evidence of ventricular dysfunction in a substantial number of patients with Alström syndrome will need to be evaluated.

In addition to the increased risk of cardiac death, significant conduction system delay also results in ventricular dyssynchrony and affects ventricular-ventricular interaction. Ventricular resynchronization has been found to improve functional outcomes and survival in adults with cardiomyopathy, and biventricular pacing has improved cardiac function in pediatric patients with cardiomyopathy. Given that both patients described in this report had early documentation of left bundle branch block, larger cohorts of patients with Alström syndrome should be evaluated with regard to conduction system involvement and possible response rates to cardiac resynchronization therapy. It has been our practice to follow published indications for both ICD and/or biventricular resynchronization therapy based on cardiac dysfunction and QRS duration.

Delay in diagnosis can delay early intervention and management of patients with ciliopathies such as Alström syndrome. Several of the ciliopathies that are associated with dilated cardiomyopathy have overlapping extra-cardiac findings with Kearns-Sayre syndrome. Although there is already a significant burden of medical care on these patients due to the multiple subspecialties involved, early detection of subtle cardiac findings may significantly impact patient management and eventual outcomes. Because of this potential impact, we recommend the involvement of a cardiologist experienced in treating patients with cardiomyopathies and their associated complications.

Conclusions

Patients with Alström syndrome are at risk of developing abnormalities of the central conduction system and clinical arrhythmias in addition to dilated cardiomyopathy. As with all forms of cardiomyopathy, we recommend routine ECG and ambulatory monitoring in all patients with Alström syndrome. In addition, a cardiologist with expertise in patients with cardiomyopathies should be an integral part of the treatment team in patients with Alström syndrome.

References

22. Hombach V, Merkle N, Torzewski J, et al. Electrocardiographic and cardiac magnetic resonance imaging parameters as predictors of a worse outcome in patients with...


