The Impact of Genetic Diagnosis on Clinical Management of Patients with Congenital Heart Disease: Cilia Point the Way

Running title: Brueckner; Cilia defects in congenital heart disease

Martina Brueckner, MD
Yale University School of Medicine, New Haven, CT

Address for Correspondence:
Martina Brueckner, MD
Departments of Pediatrics and Genetics
Yale University School of Medicine
333 Cedar Street, Fitkin 426
New Haven, CT 06520
Tel: 203 785-2022
Fax: 203 737-2786
Email: martina.brueckner@yale.edu


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The first Blalock-Taussig shunt surgery in 1945 changed major congenital heart disease (CHD) from a universally lethal condition into a survivable one. In the 50 years that followed, rapid progress in anatomic diagnosis via echocardiography, and dramatic improvement in surgical and catheterization techniques, resulted in survival to adulthood and greatly improved quality of life for many individuals with CHD. Despite this, continuing progress in caring for individuals with the most complex forms of CHD has become incremental. One of the particularly frustrating aspects of managing patients with very complex CHD is that the outcomes are highly variable, even when accounting for surgical skill and anatomic complexity. In this issue of Circulation, Nakhleh et al show that a subset of patients with CHD and Heterotaxy has airway ciliary dysfunction that may account for increased morbidity and mortality independent of the complexity of the CHD. This finding highlights a critical point that will surely affect patient care in the years to come: improving our understanding of the genetic etiology of CHD is likely to impact response to therapy. Individualized therapy for CHD based on the underlying genetics will hopefully contribute to the next major improvement in outcome.

Heterotaxy is defined as any arrangement of organs across the body’s left-right axis that differs from complete situs solitus and complete situs inversus. CHD is one of the hallmarks of Htx, and is characterized by a wide range of highly complex cardiac anatomy, including but not limited to anomalous systemic and pulmonary venous return, complex atrioventricular canal, functionally single ventricle and a range of abnormalities of the outflow tract (Table 1). As such, the cardiac lesions associated with Htx represent a significant surgical challenge; however, the surgical outcome in Htx patients remains significantly worse than that in patients with comparably complex CHD as evaluated by the Risk Adjustment in Congenital Heart Surgery-1 (RACHS) score. This indicates that a subset of patients with Htx-associated CHD may have
abnormalities extending beyond their heart disease that affect their response to extensive cardiac surgery. In order to improve outcome, we will need to identify and treat the non-cardiac factors in Htx patients that contribute to their excessive morbidity and mortality.

Extensive evidence points to a predominantly genetic etiology for human Htx, and there are reports of dominant, X-linked, and recessive pedigrees affected by Htx. Further, the incidence of Htx is increased in populations with a high degree of consanguinity, and Htx is seen in association with aneuploidies and genetic syndromes including Primary Ciliary Dyskinesia (PCD) and Bardet-Biedl Syndrome (BBS)\textsuperscript{5}. Understanding the genetics of human Htx is complicated by high lethality leading to few large pedigrees and extensive phenotypic variability. At this time, mutations have been identified in 15 genes accounting for \(~10\%\) of human Htx. In contrast, the list of candidate genes for Htx is much larger \textsuperscript{6,7}. If one adds all genes associated with human syndromic and non-syndromic Htx, and then adds genes that have been implicated in left-right development in model organisms, the total list of Htx candidate genes currently stands at more than 125 genes encompassing a broad range of biological functions (Figure 1). This indicates that Htx is actually not a monogenic diagnosis, but a manifestation of a broad range of developmental disorders that affect the development of left-right asymmetry. The clinical manifestations observed in Htx patients are further complicated by prominent effects on the development and function of other organ systems; for example, many of the genes affecting LR development, through their role in cilia, also have critical effect(s) on the lung and kidney.

Almost 20\% of the genes known to control LR development have a role in the biogenesis and function of the cilium\textsuperscript{8}. Primary cilia are found on almost all cell types at some time, and they are central to the development of LR asymmetry. The vertebrate LR axis is initiated at the
mammalian left-right organizer (LRO) late in gastrulation when motile primary cilia found on LRO cells generate directional flow of extraembryonic fluid, termed “nodal flow”9. The direction of nodal flow is determined by the inherent chirality of the cilium itself. Sensory cilia transduce nodal flow into asymmetric gene expression that is then interpreted by the developing heart to direct asymmetric cardiac morphogenesis10. Cilia, however, have a host of other functions during embryogenesis: they are essential for development and function of the kidney, bones and brain11. They are found on the cells of the developing myocardium12, and after birth, motile cilia in the respiratory tract are required for effective clearance of secretions. Seminal work by Afzelius showed that patients with Primary Ciliary Dyskinesia (PCD) have structural ciliary defects as the cause of their sino-pulmonary disease13. Notably, 6.5% of patients with a primary diagnosis of PCD also have Htx14. Since patients with Htx have a higher incidence of perioperative respiratory complications than non-Htx patients undergoing equally complex cardiac surgery, the question addressed in the article by Nakhleh et al is whether a subset of Htx patients have previously undiagnosed ciliary defects predisposing them to respiratory complications? The authors use a combination of physiological measurements of ciliary function, including direct imaging of cilia obtained from a nasal biopsy, and nasal NO measurement to assess patients for ciliary dysfunction. These techniques are more sensitive than the standard transmission electron microscopy approach more commonly used to diagnose PCD15. Notably, 42% of Htx patients had ciliary dysfunction, compared to none of the healthy controls. Sequencing of 14 PCD genes demonstrated enrichment of novel coding variants in Htx patients with ciliary dysfunction as compared to unaffected controls. Thus during very early embryonic development, a subset of patients with Htx have abnormal left-right and cardiac development due to abnormal ciliary function. Because cilia are also required in the airway
throughout life, these patients also have chronic pulmonary disease that is a variant of PCD. The perioperative respiratory complications in at least some Htx patients are thus likely not caused exclusively by their cardiac surgery, but may be the product of intrinsic pulmonary abnormalities coupled with the stress of cardiopulmonary bypass and assisted ventilation.

With the advent of high throughput sequencing, we will be able to define the precise genetic abnormality(s) underlying a significant amount of congenital heart disease. The obvious question this raises is: We can diagnose it, can we treat it? In the case of the Htx patients who have ciliary abnormalities, there are promising approaches to improve their care. PCD without associated heart disease frequently presents with neonatal respiratory distress manifested by unexplained recurrent atelectasis, poor feeding and failure to thrive\textsuperscript{16}. These symptoms resemble those associated with many types of CHD, complicating and delaying the diagnosis of intrinsic pulmonary disease. Even without associated CHD, PCD leads to chronic bronchitis, recurrent pneumonia and eventually bronchiectasis in a subset of patients. The progression of PCD-associated respiratory disease can be significantly slowed by proper therapy, and currently the lifespan of PCD patients without CHD is approaching normal\textsuperscript{15,17}. In the cardiac patient population, the pulmonary manifestations of PCD are very likely exacerbated by CHD itself, and by some of the interventions required in the care of patients with complex CHD. For example, since PCD patients are entirely dependent on cough for airway clearance, intubation and mechanical ventilation that are a necessary evil in CHD surgery are particularly challenging. Careful perioperative management of secretions, and minimizing assisted ventilation may improve immediate perioperative care. It may be beneficial for CHD-PCD patients to have intensive regular chest physiotherapy, even while they appear clinically well. There is some evidence that beta-agonists increase ciliary beat frequency\textsuperscript{18}, and therefore may be beneficial in
those CHD-PCD patients with slowed ciliary beat. Finally, in contrast to care of CHD, aggressive antibiotic management of pulmonary symptoms is a cornerstone of PCD care. It is highly likely that patients with anatomically similar, but genetically distinct, cardiac lesions will benefit from individualized care tailored in no small part by their genetic diagnosis. An atrioventricular canal defect resulting from trisomy 21 is a different disease than that arising from a defect in cilia and the development of left-right asymmetry. We will need to evaluate whether therapies aimed at specific genetic diagnoses are beneficial to CHD patients. This will require large, multicenter studies combining CHD genomics with clinical research evaluating the benefit of focused clinical management strategies. The work presented in this issue of Circulation linking ciliary defects and the vexing congenital heart disease seen in heterotaxy is a first step in this direction.

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Conflict of Interest Disclosures: None

References:


Table 1. Congenital heart disease in Htx3,4:

<table>
<thead>
<tr>
<th>Venous anomalies</th>
<th>Interrupted IVC</th>
<th>13%-21%</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Bilateral SVC</td>
<td>44%</td>
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<tr>
<td>Anomalous Pulmonary Venous Return</td>
<td></td>
<td></td>
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<tr>
<td>Atria and Ventricles</td>
<td>Atioventricular septal defect</td>
<td>32%-79%</td>
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<td></td>
<td>Single ventricle</td>
<td>68%</td>
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<tr>
<td></td>
<td>Atrial Septal Defect</td>
<td>62%</td>
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<tr>
<td></td>
<td>Ventricular Septal Defect</td>
<td>71%</td>
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<tr>
<td></td>
<td>Ebstein’s Malformation</td>
<td>7%</td>
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<tr>
<td>Outflow and Great Vessels</td>
<td>VA discordance</td>
<td>70%</td>
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<tr>
<td></td>
<td>DORV</td>
<td>33%-45%</td>
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<tr>
<td></td>
<td>Pulmonary Stenosis/Atresia</td>
<td>63%-74%</td>
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<tr>
<td></td>
<td>Coarctation of the Aorta</td>
<td>5%-16%</td>
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<tr>
<td></td>
<td>Right Aortic Arch</td>
<td>28%</td>
</tr>
</tbody>
</table>

Figure Legend:

**Figure 1.** Predicted functional role of 162 genes with a possible role in Heterotaxy.
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